

1 Quantifying clearance rates of restored shellfish reefs using 2 modular baskets

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18 **Keywords:** oyster reef, restoration, filtration, clearance rate, bivalve, climate change, marine
19 heatwave, flood

20 **Author Contribution Statement:** K.L.C., R.P., B.M., B.L.G. conceived the project. B.M.,
21 M.A., K.L.C., B.L.G. designed the experiments. B.M., R.P., K.L.C. provided resources.
22 M.A., B.M. built the seawater system and ran the experiments. M.A. collated and graphed the
23 data. B.M., M.A. analysed the data. M.A. and B.M. wrote the manuscript. B.M., K.L.C., R.P.,
24 and B.L.G. reviewed and edited the manuscript.

25 ***Abstract***

26 Shellfish reefs are among the most degraded ecosystems globally, prompting
27 substantial efforts to restore them. While biodiversity gains of restored reefs are well
28 documented, other ecosystem services such as water filtration remain poorly quantified. We
29 present a novel way of measuring water filtration by restored reefs using modular restoration
30 structures called Robust Oyster Baskets (ROB 400). Ten ROB 400s, colonised by mixed
31 invertebrate communities over 19 months in Moreton Bay, Australia, were retrieved and
32 placed individually in tanks within a recirculating seawater system. We first obtained
33 baseline clearance rates by measuring changes in the density of microalgae *Nannochloropsis*
34 *oceanica* (CS-246) over 1.5 h in tanks with and without ROB 400s. Mean clearance rates per
35 ROB 400 were $119.1 \text{ L h}^{-1} \pm 14.8 \text{ SE}$. Second, we tested how clearance rates in tanks with
36 and without ROB 400s were affected by (1) light/dark by covering tanks with black
37 polyethylene, (2) temperature by heating half of the total number of tanks $\sim 4 \text{ }^\circ\text{C}$ above
38 ambient for five days, and (3) reduced salinity by addition of freshwater from ambient (~ 36)
39 to ~ 25 or ~ 15 respectively. Clearance rates were reduced by $\sim 45\%$ when the salinity was ~ 15
40 compared to ~ 25 but were not affected by light vs dark, or temperature (ambient vs $+4 \text{ }^\circ\text{C}$).
41 Our results demonstrate modular restoration structures can be used to quantify ecosystem
42 services provided by restored reefs and to assess the vulnerability of natural and restored
43 shellfish communities to current and future threats such as light pollution, heatwaves, and
44 floods.

45 ***Introduction***

46 Shellfish reefs form when living bivalves aggregate on hard substrates in subtidal and
47 intertidal areas (Kennedy and Sanford 1999; Beck et al. 2011). These tightly bound molluscs
48 create distinct communities that engineer their surrounding environment (Kasoar et al. 2015).
49 The majority (~85%) of the world's shellfish reefs are gone, fallen to overharvesting, habitat
50 loss, pollution, and disease (reviewed by Beck et al. 2011; Gilby et al. 2018; Gillies et al.
51 2018). In North America, Europe, and Australia, several oyster reef habitats are functionally
52 extinct (Beck et al. 2011; Gillies et al. 2020). Numerous efforts are underway globally to
53 restore shellfish reefs using artificial constructions of recycled shell, crushed concrete, or
54 natural rock (Coen and Luckenbach 2000; Hernández et al. 2018; McAfee et al., 2022).

55 Most studies that have investigated the efficacy of artificial structures in restoring
56 shellfish reefs measured changes in biodiversity (e.g., Gilby et al. 2019; Xu et al. 2023). The
57 capacity of artificial reefs to restore other ecosystem services, such as water filtration,
58 remains less clear. One reason for this is the difficulty of measuring ecosystem services *in*
59 *situ*. Current methods to measure the amount of water filtered by organisms (hereafter
60 clearance rates) in the field can be large and heavy, are not easily replicated, are impractical
61 for manipulative experiments, vulnerable to tide and weather, confined to shallow water,
62 expensive, and/or limited to short deployment periods (Riisgård 2001; Grizzle et al. 2008;
63 Hansen et al. 2011). Some studies have attempted to overcome these limitations by
64 measuring clearance rates by bivalves in the laboratory (e.g., Castle and Waltham 2022;
65 Cottingham et al. 2023) or in the field using devices tailored to specific situations and species
66 (Riisgård 2001; Galimany et al. 2011). However, extrapolations from studies on single
67 species and individuals generally overestimate the clearance rates of invertebrate
68 communities by not accounting for inactive individuals (i.e., non-feeding) and interspecific or
69 intraspecific interactions (Hansen et al. 2011). Additional approaches that overcome
70 shortcomings in methods used to measure clearance rates by restored reefs are needed.

71 Some shellfish reef restoration projects are deploying modular restoration structures
72 to provide new habitat for wild animals or protection from predators for seeded spat,
73 juveniles, or adult shellfish (e.g., Walters et al. 2022; Grizzle et al. 2023). In Australia, the
74 not-for-profit organisation OzFish Unlimited designed a modular system called Robust
75 Oyster Baskets (hereafter ROB 400) and deployed ~5000 units in Moreton
76 Bay/Quandamooka, near Brisbane on Australia's east coast from 2018 (Fig. 1). Compared to

77 restoration methods that pile rock or shell to create shellfish ‘beds’, modular restoration
78 structures are easy to deploy and retrieve, can be deployed in complex configurations
79 including hanging designs, provide protection against predation, more easily facilitate public
80 participation, and can be tailored to suit the needs of each restoration project (pers. obs.; also
81 see Walters et al. 2022; Grizzle et al. 2023). In this study, through real-world application we
82 demonstrate that modular shellfish reef restoration structures can also be used to provide
83 baseline data on clearance rates of invertebrate communities colonising shellfish reef
84 restoration structures and for manipulative experiments measuring community responses to
85 stressors.

86 Shellfish reef restoration projects occur amid ongoing human impacts (Halpern et al.
87 2008; Babcock et al. 2019). To ensure effective and enduring restoration, understanding of
88 the ways in which restored shellfish reefs respond to stressors is crucial. While single species
89 studies show bivalve molluscs feed less when heat stressed, predicting the effects of
90 temperature changes on whole shellfish communities remains challenging due to interacting
91 direct and indirect effects (Turner et al. 2016; Cole et al. 2021). Freshwater influxes
92 negatively affect bivalve growth, survival, and recruitment (Pourmozaffar et al. 2020) and
93 reduces biodiversity on temperate restored shellfish reefs (Marshall et al. 2019). Floods are
94 expected to become more intense and frequent (Allen and Ingram 2002), and the
95 consequences of shifting freshet regimes for restored shellfish reefs is uncertain (Marshall et
96 al. 2019). The influence of recently recognised stressors on restored shellfish reefs, such as
97 light pollution (Davies et al. 2014), is poorly understood. Many marine shellfish can detect
98 and respond to light (Audino et al. 2020) and may depend on natural light cycles for
99 successful functioning (García-March et al. 2008; Gnyubkin 2010). Studies examining the
100 impacts of light pollution on shellfish reefs are lacking (but see Christoforou et al. 2023).

101 Current approaches to measuring clearance rates by shellfish reefs are not readily
102 adapted to testing the impacts of stressors. *In situ* methods may not be practical for
103 manipulative experiments because of short deployment periods, the inherent difficulties of
104 working in the field, and high costs (Riisgård 2001; Grizzle et al. 2008; Hansen et al. 2011).
105 While small-scale laboratory approaches are useful for quantifying the impacts of stressors
106 (Benton et al. 2007), outcomes of manipulative experiments on single species may not
107 accurately represent the responses of assemblages to the same stressor/s (Wernberg et al.
108 2012). We propose modular shellfish structures can be used to reveal the vulnerability of

109 shellfish reefs to stressors. To demonstrate this, we examined how clearance rates of
110 shellfish-dominated invertebrate communities living on ROB 400s were affected by different
111 levels of light, temperature, and salinity in single-factor experiments.

112 ***Materials and Procedures***

113 **Robust Oyster Basket 400**

114 Robust Oyster Baskets (ROB 400s) have a triangular prism-shaped design (400 × 400
115 × 300 mm) made of mild steel welded wire mesh (2.5 mm wire, 25 × 25 mm aperture)
116 encasing recycled oyster shells (Fig. 1a) which, after deployment, are colonised naturally by
117 mixed invertebrate communities dominated by filter-feeding bivalves (particularly rock
118 oysters) (Fig. 1b). Ten ROB 400s that were positioned intertidally were collected (July 2023)
119 after 19 months of deployment in Moreton Bay/Quandamooka, Queensland, Australia (-
120 27.45684, 153.39528), transported out of water, and then housed individually in aerated
121 seawater in 113-L polyethylene tanks (white colour, 455 mm H, 637 mm diameter, Nally
122 Plastics IPO25) at The University of Queensland's Moreton Bay Research Station,
123 Minjerribah (North Stradbroke Island), Queensland, Australia from July to December 2023
124 (Fig. 1c). Seawater at ambient temperature (35–37 salinity, ~21 °C) was recirculated among
125 ten tanks containing ROB 400s, six tanks without ROB 400s, and a 400-L sump with
126 mechanical filtration (Fig. 1c). ROB 400s were fed a mixed diet of live *Nannochloropsis*
127 *oceanica* (CS-246, CSIRO Australian National Algae Culture Collection, Hobart, Tasmania,
128 F media [Cell-hi F2P, Varicon aqua], 25 °C, ~35 salinity, 22:2 h light/dark photoperiod) and
129 Shellfish Diet 1800 (Reed Mariculture) three times per week. Seawater (50–66%) was
130 exchanged monthly. When ROBs were added to the system, ammonia levels spiked to 0.5–
131 1.0 ppm (API Ammonia NH₃/NH₄⁺ test kit) but fell below 0.5 ppm within 3 days. Thereafter
132 ammonia was undetectable (<0.25 ppm). Temperature, salinity, and dissolved oxygen (DO)
133 were monitored with a Horiba U-52 Series MultiParameter Water Quality Meter. Values
134 recorded during experiments are reported in Supporting Information Table S1. ROB 400s
135 were left undisturbed for at least 14 days between experiments. The surface area colonised by
136 invertebrates, the number of living and dead filter feeding invertebrates (>10 mm), and the
137 height and length (*sensu* Galtsoff 1964) of 20 haphazardly selected rock oysters (the
138 dominant invertebrates present) were measured using a tape measure, by visual count, and
139 verniers, respectively, 14 days after the final (salinity) experiment (Supporting Information
140 Table S2).

141 **Baseline clearance rates**

142 To quantify baseline clearance rates, we measured changes in the density of live
143 microalgae (*N. oceanica*) in tanks with ROB 400s compared to tanks without ROB 400s. At
144 the beginning of the experiment, water flow was turned off, which created ten independent
145 tanks housing ROB 400s and six independent tanks without ROB 400s. Each tank was
146 continuously aerated to maintain DO and keep microalgae in suspension. Live *N. oceanica*
147 were added to each tank at an initial mean density of 1.3×10^6 cells mL⁻¹ \pm 5.2×10^4 SD.
148 Total absorbance (sum of $\lambda = 750, 664, 647, 630$ nm) was measured from one sample taken
149 from each replicate tank 40 min after microalgae were added and every 30 min thereafter
150 with a spectrophotometer (Hach DR 5000™ UV-Vis, Starn Pty Ltd glass cuvette, Type 1,
151 match code 7, path length 10 mm) until 2 h had elapsed or the density of the microalgae fell
152 below $\sim 9 \times 10^4$ cells mL⁻¹. Cuvettes were triple rinsed between samples, and samples were
153 directly pipetted from the respective tank (~ 100 – 150 mm from the edge of the tank and ~ 30 –
154 50 mm depth). Data on the density of *N. oceanica* in each tank at each time point were
155 derived from total absorbance data (Supporting Information Fig. S1). Clearance rates were
156 then calculated for each replicate per Eq. 1 modified from Riisgård (2001):

$$157 \quad Cl = (V/t) \ln(C_0/C_t) \quad (1)$$

158 where C_0 and C_t equal the concentration of microalgae (cells mL⁻¹) at the time points
159 zero and t respectively, and V equals the volume of water (Supporting Information Table S2).

160 **Effect of light on clearances rates**

161 Moreton Bay/Quandamooka is one of the most light polluted coastal regions in
162 Oceania, likely exposing restored shellfish communities to artificial light at night directly and
163 via ‘skyglow’ (Kamrowski et al. 2012; Davies et al. 2014). To test whether clearance rates by
164 invertebrate communities were different when exposed to extended periods of light or dark,
165 five ROB 400s tanks and three control tanks without ROB 400s were randomly assigned to a
166 ‘dark’ treatment and covered (top and sides) with black polythene (GRUNT GRGB0042,
167 mean 5.1 lux \pm 2.6 SD at the water surface, HOBO MX Temp/Light MX 2202 set to record
168 every 10 min), while the remaining five ROB 400s tanks and three control tanks were
169 assigned to a ‘light’ treatment and left uncovered exposed to constant light (LED ‘cool
170 white’, mean 455.7 lux \pm 5.6 SD at the water surface, HOBO MX Temp/Light MX 2202 set
171 to record every 10 min). After 18 h, water flow was turned off and live *N. oceanica* were
172 added to each tank at an initial mean density of 1.09×10^6 cells mL⁻¹ \pm 5.5×10^4 SD.

173 Absorbance was measured at the beginning of the experiment and every 30 min thereafter as
174 previously described and used to derive clearance rates for all replicates in the light and dark
175 treatments for the period 0–60 min after microalgae was added.

176 **Effect of temperature on clearance rates**

177 Moreton Bay/Quandamooka is on Australia's east coast in a global climate change
178 hotspot, and forms part of the 45% of Australia's coastline which is experiencing
179 unprecedented numbers of extreme climate events (heatwaves, cyclones, floods, drought, low
180 sea level) which are expected to further intensify and become more frequent in future
181 (Hobday et al. 2016; Babcock et al. 2019). We simulated a marine heatwave lasting a period
182 of 5 days where mean temperatures were ~4 °C above ambient for that time of year (*sensu*
183 Hobday et al. 2016). The heatwave treatment was applied to five randomly allocated tanks
184 containing ROB 400s and three randomly allocated control tanks without ROB 400s that
185 were heated by 3–5 °C using 300 W titanium aquarium heaters (Aqua One TH300 or
186 Aqualogic). Five tanks with ROB 400s and three control tanks without ROB 400s were left at
187 ambient temperatures of ~20 °C. Water flow was turned off when heaters were added to the
188 tanks (time 0) and remained off for the duration of the experiment. After 24 and 120 h,
189 absorbance was measured when *N. oceanica* was added and every 30 min thereafter as
190 previously described and used to derive clearance rates for all replicates in the heatwave and
191 ambient treatments for the period 0–60 min after microalgae was added.

192 **Effect of salinity on clearance rates**

193 To test the effects of a simulated low salinity event, salinity was lowered with tap
194 water (21.5 °C, <0.01 salinity) in randomly allocated tanks to levels recorded in Moreton Bay
195 during the most recent major low salinity event, the 2010/2011 Brisbane River flood
196 (Oubelkheir et al. 2014; Clementson et al. 2021); ~15 salinity in three tanks containing ROB
197 400s and three control tanks without ROB 400s, or ~25 salinity in four tanks containing ROB
198 400s and three control tanks without ROB 400s. Salinity was not altered in three tanks
199 containing ROB 400s (salinity 36.2). AquaOne Water Conditioner© (Na₂S₂O₃, H₂O) was
200 added to all tanks (10 mL per tank). Water flow to all tanks was turned off before the
201 freshwater was added (time 0) and remained off for the duration of the experiment. After 24,
202 72, and 120 h, absorbance was measured when *N. oceanica* was added and every 30 min
203 thereafter as previously described and used to derive clearance rates for the reduced salinity
204 and ambient treatments for the period 0–120 min after microalgae was added. After 120 h,

205 salinity levels had increased by ~1 in all tanks due to evaporation (Supporting Information
206 Table S1).

207 **Statistical analysis**

208 For the baseline study, data on algal density in tanks with and without ROB 400s were
209 analysed separately using repeated measures ANOVA design with ‘time/interval’ as a fixed
210 factor. Replicate (tank) was included in the model to account for non-independence of
211 measurements taken from the same replicate over time. A type I sum of squares was used.
212 Data on clearance rates in the light experiment were analysed by two-way ANOVA using
213 ‘presence/absence of ROB 400’ and ‘treatment’ as fixed factors, and tank as the level of
214 replication. A type III sum of squares was used. Data on clearance rates in the temperature
215 experiment were analysed using repeated measures ANOVA design with ‘day’,
216 ‘presence/absence of ROB 400’, and ‘temperature’ as fixed factors. Replicate (tank) was
217 included in the model to account for non-independence of measurements taken from the same
218 replicate over time. A type I sum of squares was used. For the salinity experiment, replicates
219 with ROB 400s and replicates without ROB 400s were analysed separately. Data on
220 clearance rates were analysed using repeated measures ANOVA design with ‘day’ and
221 ‘treatment’ as fixed factors. Replicate (tank) was included in the model to account for non-
222 independence of measurements taken from the same replicate over time. A type I sum of
223 squares was used.

224 All ANOVAs were done using the permutational analysis of variance
225 (PERMANOVA) routine in Primer v7.0.23 software package. Pair-wise comparisons of
226 untransformed data were generated using Euclidean distance and 9999 permutations of the
227 raw data. Significant outcomes ($p < .05$) with more than two levels were interrogated by pair-
228 wise tests. Monte-Carlo values were used when there were fewer than 100 permutations.
229 Normality and heterogeneity of variance were examined using Q-Q residual plots, values for
230 skewness and kurtosis, and Kolmogorov-Smirnov and Shapiro-Wilk tests in IBM SPSS v29.0
231 (Quinn and Keough 2002; Field 2018). Data for the salinity experiment with ROBS present
232 were not normally distributed, but as normality is not an assumption for PERMANOVA
233 (Anderson 2017), untransformed data were analysed. All other data met the assumptions of
234 traditional ANOVA. Any outliers were included in analyses as exclusion had no effect on the
235 outcomes.

236 ***Assessment***

237 **Baseline clearance rates**

238 Density of *N. oceanica* fell in tanks containing ROB 400s (Fig. 2a), decreasing by
239 almost 85% after 90 min (Fig. 2b). In contrast, the density of *N. oceanica* in control tanks
240 without ROB 400s remained stable (Fig. 2a) and was not significantly different at the final
241 measurement (130 min) compared to the initial measurement (repeated measures ANOVA,
242 $F_{1,9} = 0.03, p = .204$, Fig. 2b). The mean clearance rate of tanks with ROB 400s present was
243 $119.1 \text{ L h}^{-1} \pm 14.8 \text{ SE}$, though clearance rates were not consistent over time (Fig. 2c). For
244 instance, mean clearance rates were initially $70.8 \text{ L h}^{-1} \pm 4.8 \text{ SE}$ but more than doubled to
245 $175.4 \text{ L h}^{-1} \pm 33.4 \text{ SE}$ in the period from 100–130 min after *N. oceanica* were added (repeated
246 measures ANOVA, $F_{2,24} = 6.63, p = .004$, followed by pairwise test: $40-70 < 70-100 < 100-$
247 130 , Fig. 2c).

248 **Effect of light on clearance rates**

249 There was no effect of light on the clearance rates of tanks containing ROB 400s (Fig.
250 3, Table 1). Tanks with ROB 400s present had significantly higher clearance rates than
251 control tanks without ROB 400s (Table 1, present > absent).

252 **Effect of temperature on clearance rates**

253 An increase in temperature ($\sim 4 \text{ }^\circ\text{C}$) had no effect on clearance rates of the invertebrate
254 communities living on ROB 400s over five days (Fig. 4, Table 1). Tanks with ROB 400s
255 present had significantly higher clearance rates than control tanks without ROB 400s (Table
256 1; present > absent). Clearance rates did not significantly vary between day 1 and day 5 (Fig.
257 4, Table 1), and there were no significant interactions among any factors (Table 1).

258 **Effect of salinity on clearance rates**

259 In tanks containing ROB 400s, salinity had a significant effect on clearance rates (Fig.
260 5, Table 1). Post hoc pair-wise tests (Table 1) indicated there was an overlapping hierarchy of
261 significance, with clearance rates higher in the 25 salinity treatment than in the 15 salinity
262 treatment, but clearance rates in the ambient treatment were not significantly different than
263 either the 25 and 15 salinity treatments (Table 1; $25 = \text{ambient} > \text{ambient} = 15$). In tanks
264 without ROB 400s, there was no effect of salinity, but the density of the algae behaved in
265 different ways depending on the day, with no significant interactions among factors (Fig. 5,
266 Table 1, $1 = 5 > 5 = 3$ days). On day 1, mean clearance rates in tanks without ROB 400s were
267 positive (i.e. > 0 , Fig. 5a) as the density of *N. oceanica* decreased, perhaps due to cells
268 clumping together or falling out of suspension. However, on day 3, mean clearance rates in

269 tanks without ROB 400s were negative (i.e., <0 , Fig. 5a) as the density of the *N. oceania*
270 increased, perhaps due to reproduction or disaggregation of cells. On day 5, mean clearance
271 rates in tanks without ROB 400s were also positive (i.e. >0 , Fig. 5c), but were not
272 significantly different than clearance rates at either day 1 or day 3.

273 **Discussion**

274 In ambient conditions, mean clearance rates by invertebrate communities on ROB
275 400s varied between 251.0 and 522.6 L h⁻¹ m⁻² among experiments. These values are similar
276 to clearance rates reported for invertebrate communities dominated by oysters and mussels in
277 field studies (e.g., Hansen et al. 2011; Vismann et al. 2016; Rullens et al. 2022). For instance,
278 bivalve beds dominated by *Crassostrea gigas* and *Mytilus edulis* had clearance rates of 138.6
279 ± 32.7 L h⁻¹ m⁻² ($n = 18$) and 447.2 ± 97.8 L h⁻¹ m⁻² ($n = 16$) respectively (Vismann et al.
280 2016). In a mussel-dominated shellfish bed, clearance rates increased from 193.5 to 806.1 L
281 h⁻¹ m⁻² after microalgae was added (Hansen et al. 2011). We also found rock oyster-
282 dominated invertebrate communities living on ROB 400s increased their clearance rates
283 through time, a trend also observed for freshwater rainbow mussels, *Villosa iris*, fed a high
284 food ration (Gatenby et al. 2013). One explanation for this could be compensatory food
285 intake where filter feeders increase the amount of water they pass through their bodies as
286 algae concentrations decrease (Bayne et al. 1987; Barillé et al. 1993; Bayne et al. 1993). Our
287 results demonstrate modular restoration structures can be used to obtain data on clearance
288 rates of whole communities *ex situ* without the inherent difficulties of sampling in the field
289 and over-estimation of clearance rates extrapolated from measurements on individuals
290 (Hansen et al. 2011). With further refinement and validation of our approach, modular
291 restoration structures could enable restoration programs to monitor clearance rates easily and
292 cheaply, facilitating access to national and international programs that fund verifiable
293 ecosystem services.

294 We found no effect of light on clearance rates of invertebrate communities occupying
295 ROB 400s. Our results are similar to those of Christoforou et al. (2023) who reported no
296 effect of ALAN (artificial light at night) on the total proportion of phytoplankton consumed
297 by the mussel *Mytilus edulis*. We are not aware of any other study that has tested the effects
298 of light on clearance rates by a community of marine filter feeders, but studies done on
299 freshwater mussels indicate that the effects of light on clearance rates is likely species
300 specific (Hills et al. 2020; Pouil et al. 2021). For instance, exposure to darkness led to

301 increases in clearance rates of the Asian clam, *Corbicula fluminea*, but not the paper
302 pondshell, *Utterbackia imbecillis* (Hills et al. 2020). The negligible impact of light in this
303 study might be because communities were dominated by intertidal rock oysters. Intertidal
304 species are generally less affected by light because they must feed while they are submerged
305 to gain adequate nutrition, regardless of the time of day (Loosanoff and Nomejko 1946).

306 We found no effect of an increase in temperature on clearance rates of invertebrate
307 communities living on ROB 400s. This contrasts with studies performed on single bivalve
308 species which found that clearance rates generally increase with temperature until a thermal
309 limit is reached beyond which feeding is depressed and clearance rates rapidly decline (e.g.,
310 Ren et al. 2000; Yukihiro et al. 2000; Parker et al. 2024). One explanation for why our results
311 differ from those of previous studies could be that the invertebrate communities we tested
312 were adapted to an intertidal environment where they are regularly exposed to a broad range
313 of temperatures (Potter and Hill 1982; Helmuth et al. 2006). We also simulated a marine
314 heatwave occurring during winter or early spring. An increase in temperature of 4 °C is more
315 likely to influence clearance rates during summer when the rock oysters that dominated the
316 ROB 400s would be closer to their upper thermal limit, especially in scenarios of aerial
317 heatwaves occurring during low tide (Dove and O'Connor 2007; Scanes et al. 2020).

318 Clearance rates were on average ~45% lower when salinity was ~25 compared to ~15,
319 perhaps because the invertebrates living in the ROB 400s approached their tolerance limit at
320 the lower extremes of salinity these invertebrate communities experience in nature
321 (Oubelkheir et al. 2014; Clementson et al. 2021). Reductions in clearance rates due to
322 exposure to low salinity have been reported for a wide range of invertebrate communities and
323 species (e.g., Navarro and Gonzalez 1998; McFarland et al. 2013; Casas et al. 2018). Our
324 results indicate the invertebrate communities living on the ROB 400s may have been stressed
325 when exposed to very low salinity but seemingly continue to provide substantial filtration
326 services regardless. As clearance rates were greatest at a salinity of ~25, shellfish reef
327 restoration may be as effective in boosting water filtration ecosystem services in estuarine
328 and coastal waterways, where the salinity is usually lower than in the ocean.

329 ***Comments and recommendations***

330 The density of microalgae in the control tanks without ROB 400s was not always
331 stable, sometimes declining presumably as microalgae fell to the bottom or slightly

332 increasing as microalgae reproduced or clumps of microalgae broke up. Filtering algae
333 cultures prior to use or using a microalga that is neutrally buoyant and/or swims may help to
334 prevent this occurring. The propensity of the density of living microalgae to vary over time
335 highlights the importance of using controls for accurate evaluation of clearance rates. To
336 date, most studies investigating bivalve clearance rates included controls only when
337 performed in a laboratory setting. Filtration studies on invertebrate communities performed
338 by adding microalgae or silt *in situ* often lack controls (e.g., Hansen et al. 2011). We suggest
339 future studies should include controls to avoid overestimating the clearance rates of filter-
340 feeding invertebrate communities.

341 The ROB 400s used in this study were initially deployed in the intertidal zone and
342 subsequently colonised by organisms adapted to intermittent emersion. This likely explains
343 why these communities appeared to cope well with transport to the laboratory and subsequent
344 experiments. The proportions of deceased shellfish within communities on the ROB 400s at
345 the end of this study are similar to those found on ROB 400s deployed in Moreton Bay
346 (Porter, unpublished data). Likewise, there was only a small spike in ammonia levels in the
347 days after transport, suggesting there was little mortality and decomposition (Canfield et al.
348 2010). Marine invertebrates living on subtidal modular restoration structures may be less
349 robust when exposed to air compared to the same species living in the intertidal (e.g.,
350 Widdows and Shick 1985; Giomi et al. 2016). Future studies should consider the need to
351 transport communities in water to reduce lethal and sublethal impacts that could influence
352 clearance rates in subsequent experiments.

353 This study demonstrates modular restoration structures can be used to test the effects
354 of physiochemical parameters (e.g., light, temperature, salinity) on clearance rates of
355 invertebrate communities. Modular baskets bypass common challenges that occur during
356 filtration studies on bivalves, both *ex situ* (e.g., high level of disturbance, small volumes,
357 often restricted to few animals and/or single species) and *in situ* (e.g., dependence on weather
358 and tide, inability to manipulate physiochemical parameters, lack of controls, poor
359 replication). Additional studies are recommended to expand our baseline study by, for
360 instance, evaluating the clearance rates of the invertebrate communities in a wider range of
361 experimental conditions predicted over the next century (e.g., reduced pH, reduced oxygen
362 levels, increased turbidity), and to test identical and more extreme conditions over extended
363 periods as extreme events often last beyond the 5 days tested here (e.g., reduced salinity over
364 weeks).

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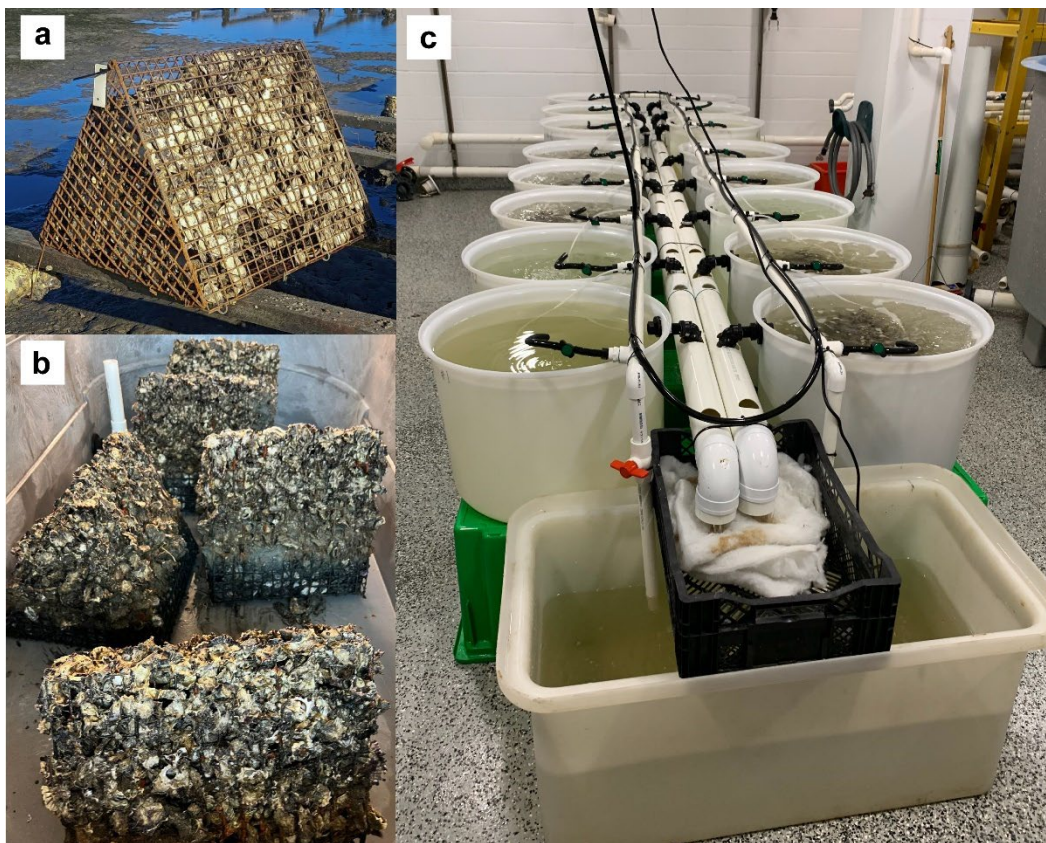
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541 **Acknowledgments**

542 We extend our deepest respect and recognition to all First Nations Peoples of the
543 Quandamooka Country, where this study was conducted, who continue cultural and spiritual
544 connections to Country. We recognise their valuable contributions to Australian and global
545 society. We thank the volunteers and staff at OzFish Unlimited Moreton Bay chapter for their
546 work in creating, deploying, and retrieving the ROBs, with special thanks to Dave Smith, Bill
547 Milligan, Gavin Gray, and Randall Kirkwood for assistance with counting invertebrates. We
548 also thank UQ students Myles Munro and Raileh Linton for assistance with sampling and the
549 kind professional staff at the Moreton Bay Research Station for help with logistics and
550 administration. We thank Associate Professor Ian Tibbetts UQ for reviewing the manuscript.
551 BM was supported by an Australian Research Council-funded DAATSIA (Discovery
552 Aboriginal and Torres Strait Islander Award, IN2000100026).

553 **Figures**



554

555 **Fig. 1.** Modular shellfish reef restoration structures and experimental setup used in this study.

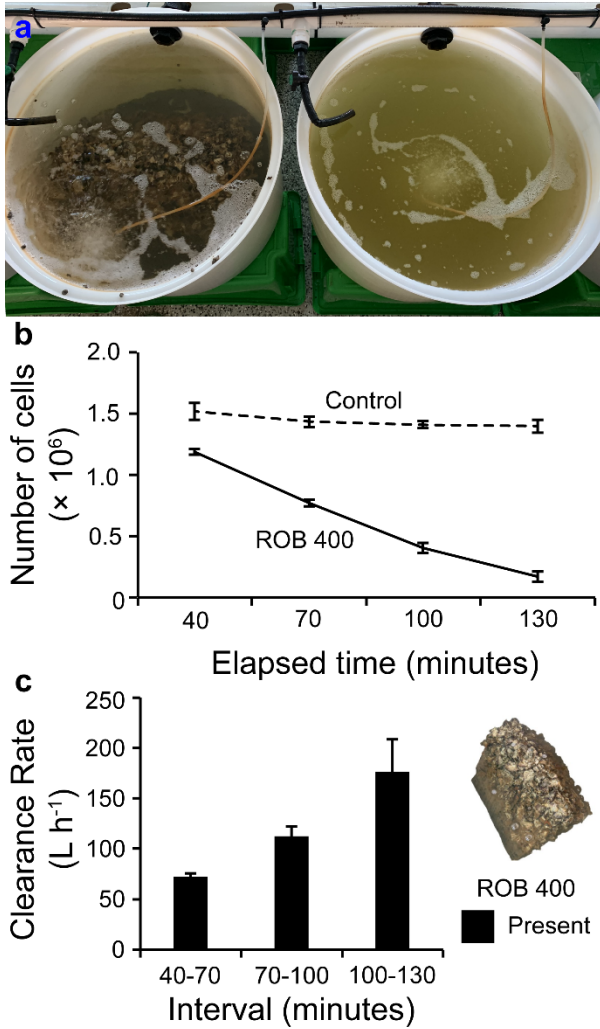
556 **a.** The Robust Oyster Basket (ROB 400) prior to deployment. OzFish Unlimited (n.d.). **b.**

557 ROB 400s colonised by invertebrate communities dominated by rock oysters following 19

558 months deployment in the intertidal zone in Moreton Bay, Australia. Andersson (2023). **c.**

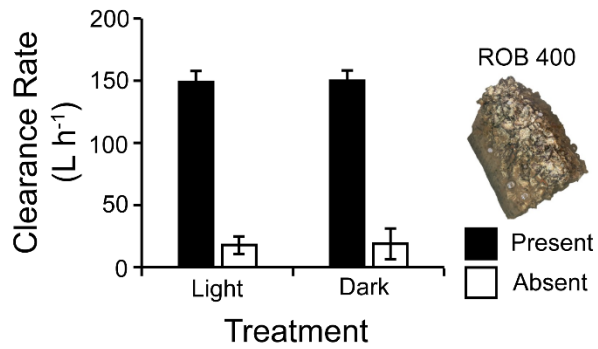
559 ROB 400s in the experimental setup used to test the effects of light, temperature, and salinity

560 on clearance rates. Mos (2023).



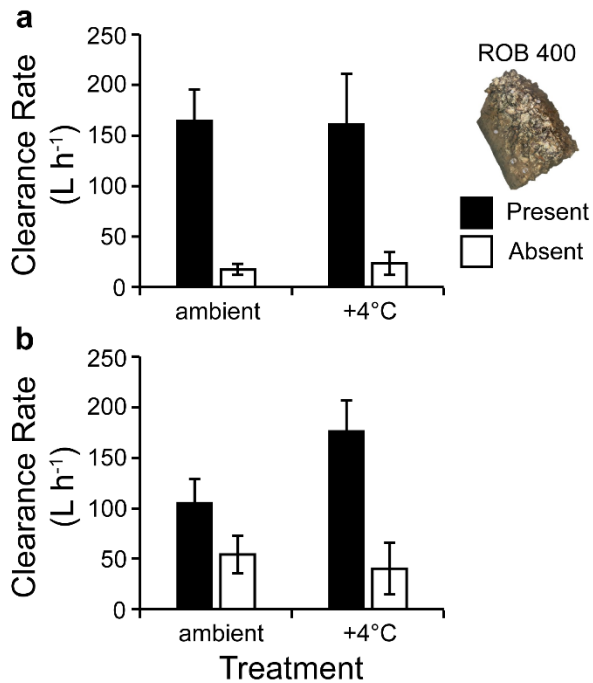
561

562 **Fig. 2.** Clearance of microalgae, *Nannochloropsis oceanica* (CS-246) by marine invertebrate
 563 communities living on modular shellfish reef restoration structures (ROB 400). **a.** Illustrative
 564 density of *N. oceanica* in tanks with ROB 400s present (left) and absent (right) 2 h after
 565 microalgae were added. Mos 2023. **b.** Density of *N. oceanica* over 90 min in static, aerated
 566 tanks without modular shellfish reef restoration structures (Control, dashed line) and with
 567 modular shellfish reef restoration structures (ROB 400, solid line). The first measurements
 568 were taken 40 min after *N. oceanica* were added to tanks at an initial density of $\sim 1.3 \times 10^6$
 569 cells mL^{-1} , with subsequent measurements taken every 30 min thereafter. Data are means \pm
 570 SE, $n = 9$ for ROB 400, $n = 5$ for control. **c.** Clearance rates of invertebrate communities
 571 living on ROB 400s in three 30-minute intervals. Data are means \pm SE, $n = 9$.



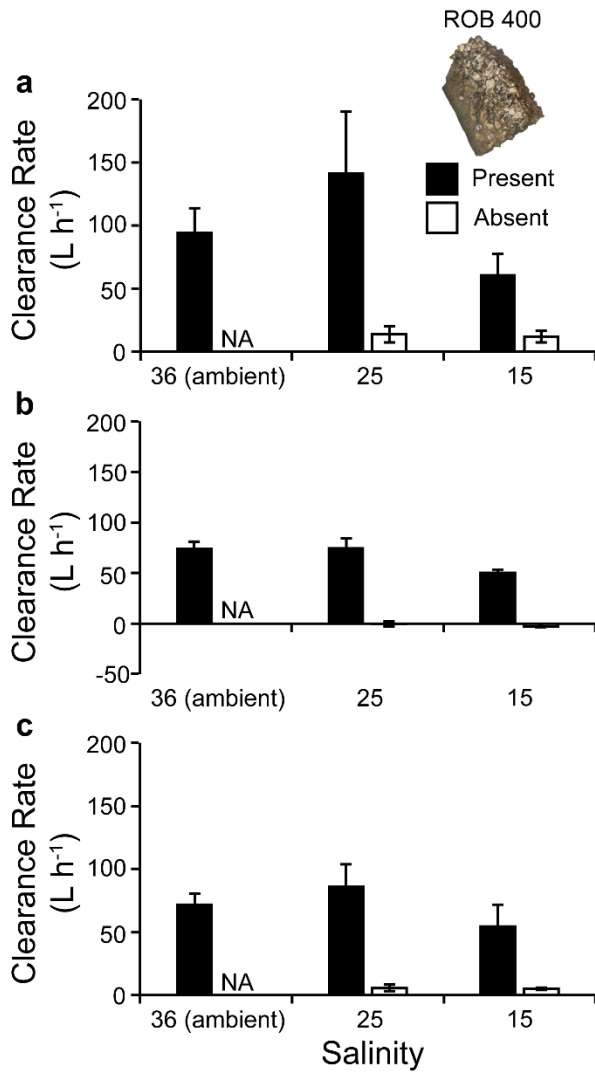
572

573 **Fig. 3.** Effects of light and the presence/absence of modular shellfish reef restoration
 574 structures housing rock oyster-dominated invertebrate communities (ROB 400) on clearance
 575 rates in static, aerated tanks. Tanks were held in darkness (mean 5.1 lux \pm 2.6 SD) or light
 576 (mean 455.70 lux \pm 5.60 SD) for 18 h before *Nannochloropsis oceanica* CS-246 was added at
 577 a density of $\sim 1.1 \times 10^6$ cells mL⁻¹. Clearance rates were derived from data on changes in the
 578 density of *N. oceanica* after 60 min. Clearance rates were affected by the presence of ROBs
 579 (present > absent) but were not influenced by treatment nor the interaction between these
 580 factors (Table 1). Data are means \pm SE; $n = 5$ for present, $n = 3$ for absent.



581

582 **Fig. 4.** Effects of temperature and presence/absence of modular reef restoration structures
 583 housing rock oyster-dominated invertebrate communities (ROB 400) on clearance rates in
 584 static, aerated tanks. Tanks were held at ambient (~21 °C) and warmed (+4 °C, ~25 °C)
 585 temperatures for **a.** 24 h and **b.** 120 h before *Nannochloropsis oceanica* CS-246 was added at
 586 an initial density of ~1–1.5 × 10⁶ cells mL⁻¹. Clearance rates were derived from data on
 587 changes in the density of *N. oceanica* after 60 min. Clearance rates were affected by the
 588 presence of ROB 400s (present > absent) but were not influenced by temperature, time, nor
 589 any interaction among these factors (Table 1). Data are means ± SE; *n* = 3–5 for present, *n* =
 590 3 for absent.



591

592 **Fig. 5.** Effects of salinity and the presence/absence of modular reef restoration structures
593 housing rock oyster-dominated invertebrate communities (ROB 400) on clearance rates in
594 static, aerated tanks. Tanks were held at ambient salinity (~36) or reduced salinity (~25 or
595 ~15) for **a.** 1 day, **b.** 3 days, and **c.** 5 days before *Nannochloropsis oceanica* CS-246 was
596 added at a density of $\sim 1.5 \times 10^6$ cells mL⁻¹. Clearance rates were derived from data on
597 changes in the density of *N. oceanica* after 120 min. No data for ambient salinity-absent
598 treatment was available (NA). Note different scale for the y-axis in panel **b.** Clearance rates
599 for tanks with (present) and without (absent) ROB 400s were statistically analysed separately
600 (Table 1). When ROB 400s were present, clearance rates were affected by salinity (25 = 35 >
601 35 = 15) but were not influenced by day nor any interaction between these factors (Table 1).
602 When ROB 400s were absent, clearance rates varied across day (day 1 = day 5 > day 5 = day
603 3) but were not influenced by salinity nor any interaction between these factors (Table 1).
604 Data are means \pm SE; $n = 3$ for present except for present-25 where $n = 4$; $n = 3$ for absent.

605 **Tables**

606 **Table 1.** Outcomes of ANOVA analyses examining the effects of light, temperature, and
 607 salinity on the clearance rates of tanks with and without invertebrate communities living on
 608 modular shellfish reef restoration structures (ROB 400) in laboratory experiments. df, degrees
 609 of freedom; MS, mean square; p/a, presence/absence; temp, temperature. Significant factors
 610 are in bold ($p < .05$).

611

Parameters	Source	df	MS	F	p	Post hoc tests
Light	presence/absence	1	6.45E4	185.80	.0002	present > absent
	treatment	1	3.63	0.01	.981	
	p/a × treatment	1	0.03	7.51E-5	.994	
	residual	12	346.99			
Temperature	day	1	2.43E3	1.17	.308	present > absent
	presence/absence	1	1.16E5	18.81	<.0005	
	temperature	1	3.21E3	0.59	.468	
	p/a × temp	1	4.52E3	0.54	.754	
	p/a × day	1	351.91	0.17	.700	
	temp × day	1	2.89E3	1.38	.277	
	temp × day × p/a	1	6.26E3	3.00	.117	
	replicate (temp × p/a)	12	6.10E3	2.92	.067	
	residual	9	2.09E3			
Salinity (Present)	treatment	2	5.34E3	2.33	.040	25 = 35 > 35 = 15
	day	2	3.82E3	2.09	.118	
	treatment × day	4	938.08	0.51	.797	
	replicate (treatment)	7	2.13E3	1.17	.370	
	residual	14	1.82E3			
Salinity (Absent)	treatment	1	11.19	1.34	.348	day 1 = day 5 > day 5 = day 3
	day	2	325.66	8.05	<.001	
	treatment × day	2	0.99	0.25	.976	
	replicate (treatment)	4	37.50	0.93	.491	
	residual	8	40.44			

612

Supporting Information for

Quantifying clearance rates of restored shellfish reefs using modular baskets

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This file includes:

Tables S1 to S2

Figure S1

Table S1. Mean water parameters (temperature (°C), salinity, and dissolved oxygen (mg L⁻¹)) for **a.** temperature and **b.** salinity treatments during laboratory experiments testing clearance rates in tanks containing mixed invertebrate communities living on oyster baskets (ROB 400) and tanks without ROB 400s (Control). Values in parentheses are standard deviation. – data not available

Experiment	Treatment	Elapsed time (h)	Temperature	Salinity	Dissolved oxygen		
a	ROB 400	+4°C	24	24.1 (0.2)	–	–	
	Control	+4°C	24	24.7 (0.7)	–	–	
	Rob 400	Ambient	24	21.9 (0.1)	–	–	
	Control	Ambient	24	21.7 (0.2)	–	–	
	ROB 400	+4°C	72	24.1 (0.2)	–	–	
	Control	+4°C	72	24.1 (0.8)	–	–	
	ROB 400	Ambient	72	19.5 (0.2)	–	–	
	Control	Ambient	72	19.4 (0.1)	–	–	
	ROB 400	+4°C	120	24.0 (0.2)	39.5 (0.1)	4.8 (0.3)	
	Control	+4°C	120	24.3 (0.8)	38.9 (0.2)	4.7 (0.2)	
	ROB 400	Ambient	120	19.3 (0.1)	38.3 (0.2)	6.9 (0.5)	
	Control	Ambient	120	19.1 (0.3)	37.8 (0.3)	6.9 (0.8)	
	b	ROB 400	15 salinity	24	21.0 (0.1)	15.5 (0.5)	6.9 (0.2)
		Control	15 salinity	24	21.0 (0.1)	15.1 (0.1)	7.2 (0.3)
ROB 400		25 salinity	24	20.9 (0.1)	25.6 (0.2)	6.0 (0.2)	
Control		25 salinity	24	21.0 (0.1)	24.7 (0.2)	7.1 (0.1)	
ROB 400		36 salinity	24	21.0 (0.1)	35.7 (0.3)	6.0 (0.1)	
ROB 400		15 salinity	72	21.1 (0.1)	15.8 (0.1)	7.1 (0.4)	
Control		15 salinity	72	20.8 (0.1)	15.7 (0.1)	8.0 (0.7)	
ROB 400		25 salinity	72	21.0 (0.1)	26.0 (0.2)	6.2 (0.3)	
Control		25 salinity	72	21.0 (0.1)	24.8 (0.1)	6.6 (0.4)	
ROB 400		36 salinity	72	21.0 (0.1)	36.4 (0.1)	6.3 (0.3)	
ROB 400		15 salinity	120	21.5 (0.1)	16.7 (0.1)	7.3 (0.2)	
Control		15 salinity	120	21.2 (0.1)	16.1 (0.2)	7.8 (0.2)	
ROB 400		25 salinity	120	21.4 (0.1)	26.1 (0.3)	7.1 (0.3)	
Control		25 salinity	120	21.4 (0.1)	25.3 (0.2)	6.5 (1.1)	
ROB 400		36 salinity	120	21.3 (0.1)	36.8 (0.1)	5.9 (0.2)	

Table S2. Information on the tank setup and robust oyster baskets (ROB 400) used in experiments; the volume of seawater in each tank (L), the surface area of each ROB colonised by filter-feeding invertebrates (m²), the number of living and non-living (in parentheses) individuals (>1 cm) of the dominate filter-feeders found on the ROB400s, the total number of filter feeders (>1 cm) on each ROB, the height and length (mean ± SD) of 20 rock oysters haphazardly measured from each ROB400, and the treatment randomly allocated to each tank for the three experiments (light/dark, heatwave, and flood).

Tank #		Volume of seawater in tank (L)	Surface Area m ²	rock oysters <i>Saccostrea</i> spp.	hairy mussel <i>Trichomya cf. hirsuta</i>	pearl oyster <i>Pinctada cf. albina</i>	total number of filter feeders	alive (%)	height mm	length mm	Light/dark	Heatwave	Flood
1	Control	107									Light	Ambient	15
2	Control	107									Dark	Ambient	25
3	ROB 400	81.5	0.246	93 (30)	1 (0)	0 (5)	94 (35)	72.9	26.8 (6.9)	32.7 (8.7)	Dark	Ambient	15
4	ROB 400	80	0.259	102 (55)	3 (0)	8 (5)	113 (60)	65.3	26.5 (5.4)	21.8 (6.0)	Light	Heated	36
5	ROB 400	80.5	0.285	121 (43)	0 (0)	5 (8)	126 (51)	71.2	23.8 (9.2)	32.7 (6.9)	Dark	Heated	36
6	ROB 400	80	0.289	170 (144)	7 (1)	3 (3)	180 (148)	54.9	26.0 (9.0)	21.4 (5.4)	Dark	Ambient	25
7	Control	107									Light	Heated	25
8	ROB 400	80	0.299	122 (65)	65 (0)	18 (4)	205 (69)	74.8	21.8 (6.9)	36.0 (8.60)	Light	Heated	25
9	ROB 400	82	0.264	104 (80)	36 (0)	7 (0)	147 (80)	64.8	29.9 (9.6)	21.7 (7.6)	Dark	Ambient	25
10	Control	107									Dark	Heated	15
11	Control	107									Light	Ambient	15
12	ROB 400	83	0.315	157 (138)	25 (1)	8 (7)	190 (146)	56.5	24.5 (6.1)	32.9 (3.2)	Light	Heated	36
13	ROB 400	85.5	0.310	84 (53)	2 (0)	3 (3)	89 (56)	61.4	29.2 (8.4)	25.6 (9.5)	Light	Ambient	25
14	Control	107									Dark	Heated	25
15	ROB 400	81	0.309	169 (173)	22 (0)	2 (10)	193 (183)	51.5	26.6 (7.4)	32.0 (6.6)	Dark	Heated	15
16	ROB 400	81	0.298	172 (103)	7 (1)	0 (4)	179 (108)	62.4	31.5 (9.5)	25.4 (8.3)	Light	Ambient	15
		MEAN	0.287	129.4	16.8	5.4	151.6	63.6					
		±SE	0.024	10.9	6.6	1.7	43.5	2.5					

The mean total surface area colonised by invertebrates on each ROB did not differ among treatments for the light/dark (ANOVA, $F_{1,8} = 1.44$, $p = .242$), heatwave ($F_{1,8} = 0.61$, $p = .458$), or flood ($F_{2,7} = 0.05$, $p = .925$) experiments.

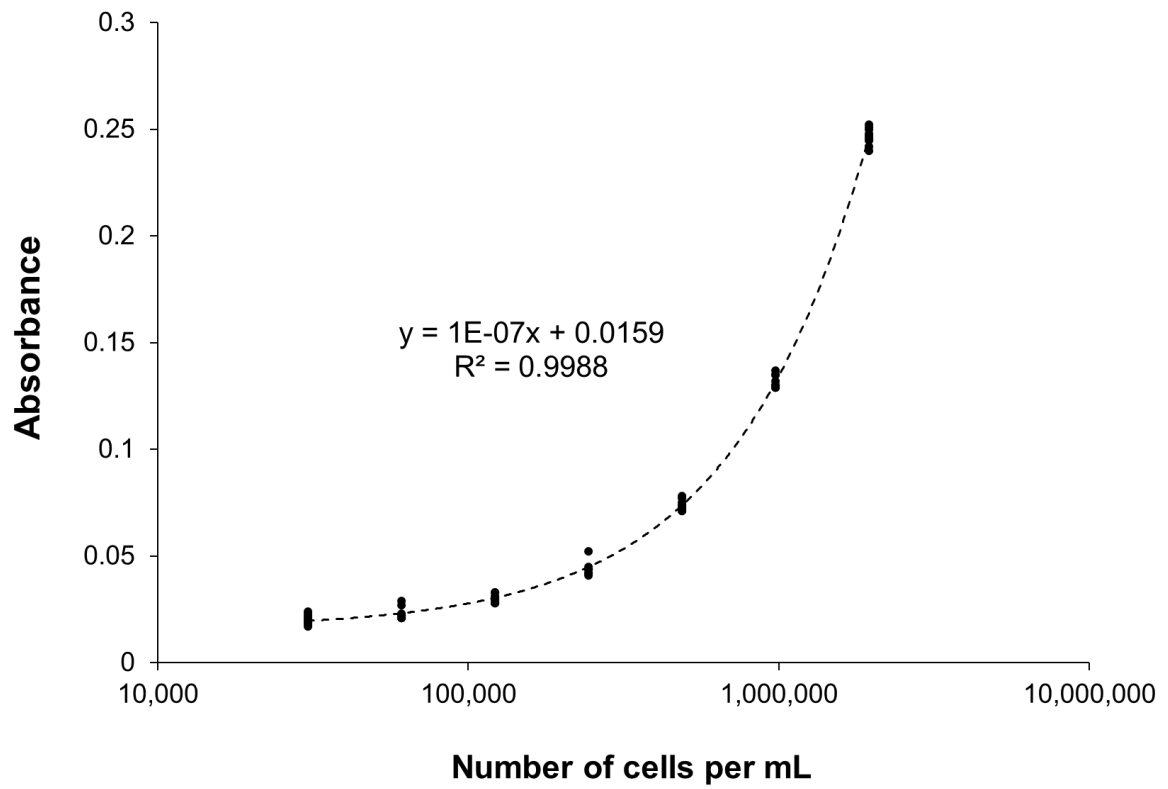


Fig. S1. Relationship between the number of cells per mL of *Nannochloropsis oceanica* (log scale) and total absorbance (sum of $\lambda = 750, 664, 647, 630$ nm) measured using a spectrophotometer (Hach DR 5000™ UV-Vis, Starn Pty Ltd glass cuvette, Type 1, match code 7, path length 10 mm).