The reef-building coral *Galaxea fascicularis*: a new model system for coral symbiosis

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Abstract

Reef-building corals owe their evolutionary success to their symbiosis with unicellular algae (Symbiodiniaceae). However, increasingly frequent heat waves lead to coral mass-bleaching events and pose a serious threat to the survival of reef ecosystems. Despite significant efforts, a mechanistic understanding of coral-algal symbiosis functioning, what leads to its breakdown and what can prevent it, remains incomplete. The main obstacles are low amenability of corals to experimental handling and, owing to its obligatory nature, the difficulties of manipulating the coral-algal association. Indeed, many studies on the symbiotic partnership are conducted on other cnidarian model organisms and their results may therefore not be fully transferable to tropical reef-building corals. Here, we identify the tropical stony coral species *Galaxea fascicularis* as a novel candidate coral model system. Individual polyps of this species can be separated, enabling highly replicated genotype studies, and are well suited to experimental investigation of the symbiosis as they can be easily and effectively rid of their algal symbionts (bleached). We show that bleached adult individuals can reestablish symbiosis with homologous and heterologous symbionts. We also report the completion of the gametogenic cycle *ex-situ* and the successful spawning in aquaria over multiple years. These achievements help overcome several of the major limitations to direct research on corals and highlight the potential of *G. fascicularis* as an important new model system for investigations of symbiosis functioning and manipulation.

Keywords: Model organisms; reef-building corals; menthol bleaching; symbiosis manipulation; *ex situ* sexual reproduction

Introduction

Reef-building corals form obligatory endosymbiotic association with unicellular algae of the family Symbiodiniaceae. This association is key to their evolutionary success, but it is also at the heart of corals' susceptibility to global climate change, which manifests in coral bleaching - the breakdown of the coral-algal symbiosis (Dietzel et al. 2020). Bleaching is mostly driven by marine heatwaves which are predicted to worsen, causing the loss of virtually all coral reefs by the end of this century (van Hooidonk et al. 2016).

The interaction between heat stress and bleaching has been studied for more than 30 years (Gates et al. 1992; McLachlan et al. 2020), but a detailed understanding of the mechanisms underlying coral bleaching is still missing. Progress in coral symbiosis research is hampered by two main aspects. First, corals are challenging to maintain in aquarium settings as they tolerate only a narrow range of environmental conditions. Second, the obligatory nature of the coral-algal association makes it particularly challenging to physically and functionally separate the partners, an approach often necessary to unravel the mechanisms underlying symbiosis breakdown and what could prevent it (Weis et al. 2008; Voolstra 2013).

Model organisms are integral for understanding fundamental biological principles and symbiotic cnidarians have successfully been used as "coral models", advancing our understanding of coral bleaching and holobiont functioning (Weis et al. 2008). These model organisms share important traits with corals, such as being cnidarians that associate with microalgae, yet they also lack other features that normally represent obstacles to research work such as the calcium carbonate skeleton. In addition, the ability to study the animal host and its algal symbiont in isolation is more easily achieved with facultatively symbiotic organisms such as *Hydra* spp., Aiptasia (*Exaiptasia diaphana*), and *Astrangia* spp. (Dimond and Carrington 2007; Weis et al. 2008; Galliot 2012).

60 Cnidarian model organisms allow the study of basal or shared traits or phenomena at a speed that
61 would not be possible with reef-building corals, but testing on the latter remains necessary for the
62 understanding of coral-specific or ecologically relevant aspects. The obligatory symbiosis with
63 Symbiodiniaceae in reef-building corals has important biological implications (Falkowski et al.
64 1984; Hoegh-Guldberg 1999), similar to the calcification process that is deeply intertwined with
65 coral physiology and ecology (Gattuso et al. 1999). It is therefore important to establish a "true"
66 tropical reef-building coral model species.
67 The richness of the scleractinian taxon offers a vast array of species to choose from in the quest
68 for a suitable candidate coral model species. These can be evaluated for their tractability (as
69 amenability to experimental work) considering several aspects which, as proposed by Puntin et al.
70 (2022) should include: 1) pre-existing knowledge: baseline information of the organism's biology
71 is necessary to interpret and contextualize results; 2) compatibility with aquarium rearing: the
72 possibility to maintain the organism and preferably to complete its life cycle in the lab is essential
73 to increase its availability, reduce confounding effects such as unknown life history, improve
74 reproducibility, and lower the pressure on threatened wild populations; 3) amenability to symbiosis
75 manipulations is necessary to help unravel complex coral-algal functional interactions, and can be
76 broken down into the organism's suitability to be rendered aposymbiotic (bleached), to be
77 maintained aposymbiotic for a certain time, and to then be re-infected with Symbiodiniaceae.
78 We have identified the coral species *Galaxea fascicularis* (Linnaeus, 1767) (clade: Complexa,
79 family: Euphyllidae) as a promising candidate coral model system. The species is relatively
80 common across the Indo-Pacific region (Veron et al. 2016), where the phylogeny of the genus
81 *Galaxea* is geographically well resolved (Wepfer et al. 2020b). It is also one of the more commonly
82 used species in heat-stress experiments (McLachlan et al. 2020) and in studies on coral

83 calcification (e.g., Marshall and Clode 2004; Al-Horani 2005). Importantly, other community
84 resources such as an annotated draft genome are available (Liew et al. 2016; Niu et al. 2016,
85 http://www.gfas.reefgenomics.org/). In addition, *G. fascicularis* is known for its tolerance to
86 stressors as it is "largely unaffected by bleaching" in the field (Marshall and Baird 2000) and for
87 being easy to grow and propagate in aquaria (Pavia and Estacion 2019). Further, its morphology
88 provides additional practical advantages with relatively big polyps, which can be easily isolated
89 (Al-Horani 2005). This conveniently allows reduction of the complexity of the organism from
90 colonial to individual, enables high clonal replication, and facilitates visualization (Al-Horani et
91 al. 2003; Marshall and Clode 2004). Its association with Symbiodiniaceae is relatively well
92 characterized (Huang et al. 2011; Wepfer et al. 2020a). Furthermore, the species' reproductive
93 mode and spawning patterns in the wild are well documented (Babcock et al. 1986; Harrison 1988;
94 Keshavmurthy et al. 2012). However, its amenability to symbiosis manipulation, to experimental
95 handling of bleached individuals, and the ability to complete gametogenic cycles and spawning in
96 closed aquaria has not yet been described.
97 The overarching aim of this study was to explore the potential of *Galaxea fascicularis* as a coral
98 model. Specifically, we investigated: 1) the feasibility to render polyps aposymbiotic by
99 eliminating Symbiodiniaceae (bleaching), 2) the amenability to experimental manipulation of
100 symbiotic and aposymbiotic polyps in a simplified system, 3) the re-establishment of the symbiosis
101 with various species of Symbiodiniaceae, and 4) the possibility to induce full gametogenic cycles
102 *ex situ*, with subsequent spawning under aquarium conditions. These aspects are central to
103 overcoming the two main obstacles to coral research work: low tractability of corals and their
104 complex nature.

Materials and methods

Coral collection and long-term aquarium rearing

Colonies of *Galaxea fascicularis* were collected from three geographical locations: the Red Sea, Hong Kong, and the Great Barrier Reef. Red Sea colonies were collected at 9-13 m depth at the North-Eastern protected end of the reef "Al Fahal" in the central Saudi Arabian Red Sea (22.3054°, 38.9655°) in March 2019, and transported to the Ocean2100 aquarium facility at Justus Liebig University Giessen (Germany) where they are part of the live collection (CITES permit 19-SA-000096-PD). In the aquarium system, light was provided by white and blue fluorescent lamps with a light:dark cycle of 12:12 h at 130 -160 µmol photons m^{-2} s^{-1}, which approximates light condition at the collection site (Ziegler et al. 2015). Salinity was maintained around 35 and temperature at 26 °C. Colonies were fed daily with a combination of frozen copepods, *Artemia*, krill, and *Mysis*. Hong Kong colonies were collected from 5 m (max. depth) from Crescent Island (22.5308°, 114.3150°) in November 2020 and transported to the University of Hong Kong, where they were fragmented and acclimated to aquarium conditions for one month. The aquarium system consisted of 6-L acrylic tanks placed within a Plant Growth Chamber (Panasonic MLR-352H-PA). Each tank held eight ~3 cm-nubbins and was fitted with a submersible pump (Atman AT301) and a small water filter (Shiruba PF120). The fragments used in symbiosis reestablishment were fed twice-weekly with a powdered blend of marine plankton (ReefRoids, Polyp Lab) followed by a partial water change. Light, salinity, and temperature conditions were consistent with those maintained in the Ocean2100 facility.

For *ex situ* spawning, 35 colonies were collected from Arlington Reef, Great Barrier Reef, Australia (-16.7000°, 146.0500°) in September 2019, and transported to the Horniman Museum and Gardens (United Kingdom; CITES permit 585319/01). Colonies were distributed into four

128 coral spawning systems that replicated the natural environmental parameters (seasonal
129 temperature, photoperiod, solar irradiance, and lunar cycle) required to stimulate reproduction.
130 Aquarium design and coral husbandry protocol was based on the mesocosms described by Craggs
131 et al. (2017), and the seasonal temperature profile were based on a non-sequential eight year
132 average (1998 - 2017) from Moore Reef (-16.8667°, 146.2334°')
133 (http://data.aims.gov.au/aimsrtds/datatool.xhtml?site=931&param=water%20temperature).

134 **Preparation of single polyps for experimentation**

135 For the menthol bleaching and thermal performance experiment, we used single polyps from three
136 Red Sea colonies each showing distinct coloration (Fig. S1). Replicate clonal polyps were isolated
137 from each colony using an electric rotary cutter, mounted on coral glue (Grotech, CoraFix
138 SuperFast), and allowed to heal for two weeks in the Ocean2100 aquarium facility (Schubert and
139 Wilke 2018).

140 **Menthol bleaching to remove algal symbionts**

141 An equal number of polyps from each colony were randomly assigned to the control group
142 (maintained in a symbiotic state), and the bleached group (chemically bleached with menthol).
143 Menthol bleaching followed a protocol modified from Wang et al. (2012) and consisted of three
144 days of treatment in 0.38 mM menthol solution in seawater, followed by a day of rest and a fourth
145 day of menthol treatment. Each day the polyps were incubated in menthol for 8 h under light and
146 stirring (Fig. S2). After, the polys were rinsed and kept in clean containers with air bubbling and
147 stirring. During menthol treatment, the treated polyps were only exposed to filtered (1.2 µm)
148 artificial seawater (FASW) to prevent exposure to Symbiodiniaceae. Following, all polyps
149 (including, for consistency, the symbiotic group) were maintained in FASW.

150 To confirm bleaching, half of the polyps from each group (bleached and symbiotic) and each
151 colony were visually assessed with a fluorescence stereomicroscope (Leica MZ16 F) 10 days after
152 the termination of the menthol treatment. Representative pictures were taken under natural light
153 (brightfield, light source: 2950 K) and under UV light with GF2 filters allowing visualization of
154 the green fluorescent proteins and chlorophyll (Fig. 1a). Following the same bleaching protocol,
155 menthol-bleached polyps were also documented using a compound epifluorescence microscope to
156 reach higher resolution (Leica DM 5500B, TX2 filter) (Fig. 1b).

157 **Thermal performance experimental design**

158 Symbiotic and bleached polyps were moved to the experimental tanks on the day after the last
159 menthol exposure and allowed to recover and acclimate for 10 days. The experimental system
160 consisted of eight 5-L glass tanks (20 cm × 30 cm) equipped with a small pump (Resun SP-500)
161 in a temperature-controlled water bath (26 ˚C). White fluorescent lamps were used to maintain the
162 light cycle and intensity consistent with long-term rearing conditions. Bleached and symbiotic
163 polyps were kept in separate tanks (four per group) filled with FASW, with three to six polyps
164 each (Fig. S3). Polyps were fed each day after the end of the dark incubation, followed by 10 %
165 water change after 2-3 h. Each polyp was fed one small frozen adult Artemia shrimp pipetted
166 directly on top of the oral opening. The polyps were cleaned every two to three days after the
167 incubations to remove fouling that would interfere with respirometry measurements.

168 To test the amenability to experimental handling of single polyps and to compare metabolic rates
169 of bleached and symbiotic polyps, we conducted a thermal performance curve experiment.
170 Photosynthesis and respiration of *G. fascicularis* polyps were measured across 12 ˚C temperature
171 over 10 days. For this, the polyps were acclimated to a different temperature each day, in the

172 following order: 20, 22, 23, 24, 25, 26, 27, 28, 30 to 32 °C. The temperature of the water bath was
173 changed overnight, and incubations took place the subsequent day at the respective temperature.

174 **Photosynthesis and respiration of symbiotic and bleached polyps**

175 We measured oxygen evolution of individual *G. fascicularis* polyps in light and dark incubations
176 to calculate net photosynthesis (PN) and dark respiration (R) respectively. For these measurements
177 four bleached and four symbiotic polyps from three colonies were used (n = 12). Every day, each
178 polyp was placed in individual 160 mL glass incubation jars (Weck, Germany), equipped with a
179 magnetic stirrer (~5 cm s^{-1}). Three additional jars containing only seawater and stirring bars were
180 used to control background biological activity. Light intensity of 230 - 280 µmol photons m^{-2} s^{-1}
181 during light incubations (ATI 80W Aquablue Special) and temperatures were maintained through
182 thermostat-controlled water baths placed on a custom-made multiplate stirring system (Rades et
183 al. 2022).

184 Dissolved oxygen (DO) was measured at the start and at the end of each incubation using a
185 handheld multiparameter probe (WTW Multi 3620 IDS set). Light incubations lasted 2 h around
186 mid-day followed by 1.5 h of dark incubation. Net photosynthesis and respiration were calculated
187 using the following formula (Schneider and Erez 2006):

188 $$PN\ or\ R[mgO_2 \times cm^{-2} \times h^{-1}] = \frac{\Delta O_2[mg \times L^{-1}] \times V_{incubation}[L]}{T[h] \times SA[cm^2]}$$

189 Where ΔO_2 is the difference in dissolved oxygen between the end and the start of the incubation
190 (DO_{end} - DO_{start}) corrected for the controls' ΔO_2. This is multiplied by the volume of the incubation
191 chambers in L ($V_{incubation}$), and divided by incubation time (T) in hours and polyp surface area (SA)
192 in cm^2.

193 **Surface area measurements**

194 Polyp surface area was measured through photogrammetry. Between 40 to 50 pictures of each
195 polyp were taken with a phone camera two days before the beginning of the incubations. These
196 were used to create 3D models with 3DF Zephyr Free (v.4.523), cleaned on Artec3D (Studio 11
197 Professional v.11.2.2.16), and loaded on MeshLab (v.2016.12, Cignoni et al. 2008) for size scaling
198 and calculation of live coral surface area.

199 **Thermal performance data analysis**

200 All analyses were conducted in the R statistical environment (v.4.1.0, R Core Team 2021) and
201 plotted with ggplot2 (v.3.3.5, Wickham 2016). The effects of symbiotic state ('symbiotic' vs.
202 'bleached') or colony identity on net photosynthesis (PN) and respiration (R) were analyzed using
203 linear mixed-effect models. The effect of symbiotic state on R was tested by setting 'symbiotic
204 state' as a fixed factor and 'polyp identity', 'colony', and 'temperature' as random factors. The
205 effect of colony identity on PN and R was tested considering light and dark incubations separately
206 and setting 'colony identity' as fixed factor and 'polyp identity', 'temperature', and 'symbiotic
207 state' as random factors. The package 'lmerTest' (v.3.1-3, Kuznetsova et al. 2017) was used to
208 construct the models and calculate p-values. The package 'performance' was used to check the
209 residuals of fixed and random factors and to compare alternative models (v.0.7.3, Lüdecke et al.
210 2021). Differences between colonies were tested as pairwise comparisons of estimated marginal
211 means (Searle et al. 1980) with Bonferroni correction with the 'emmeans' package (v.1.6.2-1).

212 **Symbiosis reestablishment after bleaching**

213 We explored the possibility of returning polyps to the symbiotic state after menthol bleaching
214 using three approaches. The first and simplest approach consisted of testing natural symbiont
215 acquisition. One month after the menthol treatment, bleached polyps were placed next to symbiotic

216 colonies of *G. fascicularis* to expose them to Symbiodiniaceae naturally released from conspecifics
217 into the water column. The second approach consisted of inoculating bleached polyps with freshly
218 isolated Symbiodiniaceae from symbiotic *G. fascicularis*. For this, a donor polyp from each colony
219 was placed upside-down inside a 1.5 mL tube and centrifuged (Heraeus Biofuge Pico, England) at
220 8,300 *g* for 1 min to detach the tissue from the skeleton (Wilkinson et al. 2016). The slurry was
221 homogenized with 600 µl of FASW, centrifuged at 3,400 *g* for 1 min, and the resulting pellet
222 resuspended twice to clean the algal fraction. This Symbiodiniaceae suspension was used to
223 inoculate two to three polyps each with Symbiodiniaceae from the same colony or from one
224 respective other colony. For the inoculation, bleached polyps were placed in individual beakers,
225 the Symbiodiniaceae inoculum was pipetted on the oral opening together with one Artemia shrimp,
226 and allowed to feed for 3 h. Subsequently, inoculated polyps were transferred to the long-term
227 rearing aquarium facility. Symbiosis reestablishment was visually assessed after one month
228 (Fig. 3).
229 The third approach consisted of inoculating menthol-bleached coral nubbins with cultured
230 Symbiodiniaceae. Two species of symbionts, *Cladocopium goreaui* and *Durusdinium trenchii*
231 were chosen for targeted reinfection of the bleached fragments, as species from both these genera
232 are known to form stable symbiosis with *G. fascicularis* across the South China Sea (Tong et al.
233 2017). Batch cultures of *C. goreaui* (AIMS-SCF-055, isolated from *Acropora tenuis*) and *D.*
234 *trenchii* (AIMS-SCF-088, isolated from *A. muricata*) were obtained from the Symbiont Culture
235 Facility at the Australian Institute of Marine Science and maintained in f/2 media (Guillard 1975)
236 under the same conditions as the corals. A total of 16 fragments were inoculated with a
237 concentration of 200 symbiont cells ml^{-1} in each 6-L aquarium. The concentration of symbionts
238 was divided in a 90:10 ratio (180 cells ml^{-1} *D. trenchii* : 20 cells ml^{-1} *C. goreaui*). This ratio was

239 chosen because *G. fascicularis* is more commonly associated with symbionts from the genus
240 *Cladocopium* under the temperature conditions of this experiment (Dong et al. 2009; Zhou et al.
241 2017b). The symbiont aliquots were centrifuged to remove the culture media, washed once with
242 FASW before final resuspension in 10 mL FASW. The bleached fragments were first fed with
243 ReefRoids before being exposed to the chosen symbionts. For this, a suspension of ReefRoids in
244 ASW was pipetted directly into the oral opening of each polyp. The same process was then
245 immediately repeated with the suspended symbionts. Aquarium filters were turned off for 1 h to
246 allow the corals to feed and to prevent removal of symbiont cells from the water column.
247 Fragments were inoculated twice a week for 6 weeks, until symbiosis reestablishment was visually
248 confirmed. Symbiodiniaceae populations *in hospite* were then characterized with fluorescent in-
249 situ hybridization (FISH) and flow cytometry (McIlroy et al. 2020).

250 **Fluorescent In-Situ Hybridisation (FISH)**

251 After 6 weeks of inoculation, a single polyp was removed from each fragment with a chisel. Tissue
252 was removed from the skeleton with an airbrush containing deionized water. The resulting slurry
253 was homogenized using a 20-µl pipette tip. 1.5 mL of homogenate was transferred to a 2-mL
254 Eppendorf tube and centrifuged at 500 rpm for seven minutes, after which the supernatant was
255 discarded. The remaining pellet was washed once before being preserved in 5X SET buffer and
256 stored at -80 °C for further analysis.

257 Samples were prepared following an adapted protocol from McIlroy et al. (2020). Briefly, after
258 washing with 5X SET in IGEPAL at 0.4 and 0.1 % vol/vol samples were split equally between
259 three 1.5-mL black Eppendorf tubes. Each sample had one tube treated with a SymC probe, one
260 tube with a SymD probe and one without any probe as a control. Both SymC and SymD probes
261 were designed to have genus-level specificity for *Cladocopium* and *Durusdinium*, respectively.

262 After overnight hybridization at 45 °C (5X SET, 0.1 % vol/vol IGE- PAL, 10 % vol/vol formamide
263 and 100 pmol of the relevant probe), samples were washed once in warm 1X SET, vortexed and
264 resuspended in 500 µl of 1X SET for same-day cytometric analysis on a BD FACSAria Fusion
265 flow cytometer (BD Biosciences, CA). FlowJo™ Software (v.10.8.1, BD Life Sciences) was used
266 to determine the percentage of probe-positive cells (either *Cladocopium* or *Durusdinium*), to
267 identify a) the presence and b) the relative abundance of each algal genus within the sample.

268 **Completing gametogenic cycles and *ex-situ* spawning**

269 *Galaxea fascicularis* in the GBR spawns between October and December, 0.05-200 minutes after
270 sunset (MAS) and 1 – 8 nights after full moon (NAFM) (Babcock et al. 1986; Baird et al. 2021).
271 To determine if the *ex situ* reproductive pattern remained in synchrony with the wild we followed
272 our colonies over three consecutive reproductive seasons (2019 – 2021).
273 To ascertain gamete development in each colony, two months prior to predicted spawning, and
274 two to four days prior to the full moon, whole polyps were removed from each colony and
275 longitudinal sections imaged (Canon 5d MKIII, Fig. 4a,d). Based on the stage of gamete
276 development in these sections *ex situ* spawning activity was predicted.
277 To record spawning observations each year, colonies were observed with red head torches during
278 the predicted spawning months. Observations commenced two NAFM and 30 mins prior to the
279 predicted spawning time, during which time the broodstock aquariums were isolated from the
280 filtration system. In addition, all internal water pumps were also turned off leaving the aquariums'
281 water static. Observations continued over multiple consecutive nights until all colonies that had
282 developed gametes, during that reproductive season, spawned. Spawning date and time of each
283 colony was recorded, with onset of spawning being denoted as the time of first gamete release.

Following release, gametes were pooled to allow fertilization to occur, and subsequent embryos were reared and larvae settled following the method described by Craggs et al. (2019).

Results

Production of single polyps and long-term rearing of bleached polyps

We produced clonal replicates of *G. fascicularis* colonies from the Red Sea and Hong Kong and bleached them with menthol. All polyps (n = 24, including additional polyps not used in this experiment) and all coral fragments (n = 16) survived the menthol treatment and appeared visually completely white. Visual inspection under a fluorescent stereomicroscope revealed no detectable algal cells in the bleached polyps (Red Sea colonies, Fig. 1a), but inspection at higher magnification (compound fluorescent microscope, ×400) revealed the presence of few scattered algal cells, limited to the base of the tentacles (Fig. 1b).

All polyps that were employed in the thermal performance experiment (n = 12) remained visually completely bleached and viable throughout the 10 days of temperature treatment, with the exception of two polyps that appeared dead on the 7th and 8th day of incubation, respectively. These polyps showed atrophic and unresponsive tentacles, and overall thinning of the soft tissue until it was not recognizable. Overall, we successfully maintained the bleached polyps for three weeks from the termination of the menthol treatment.

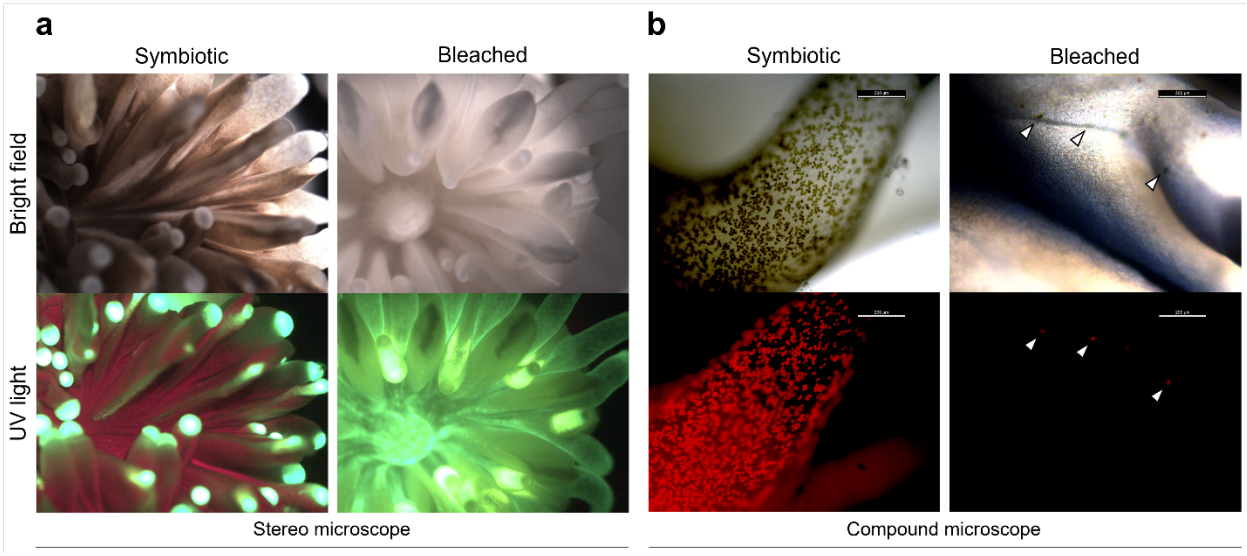
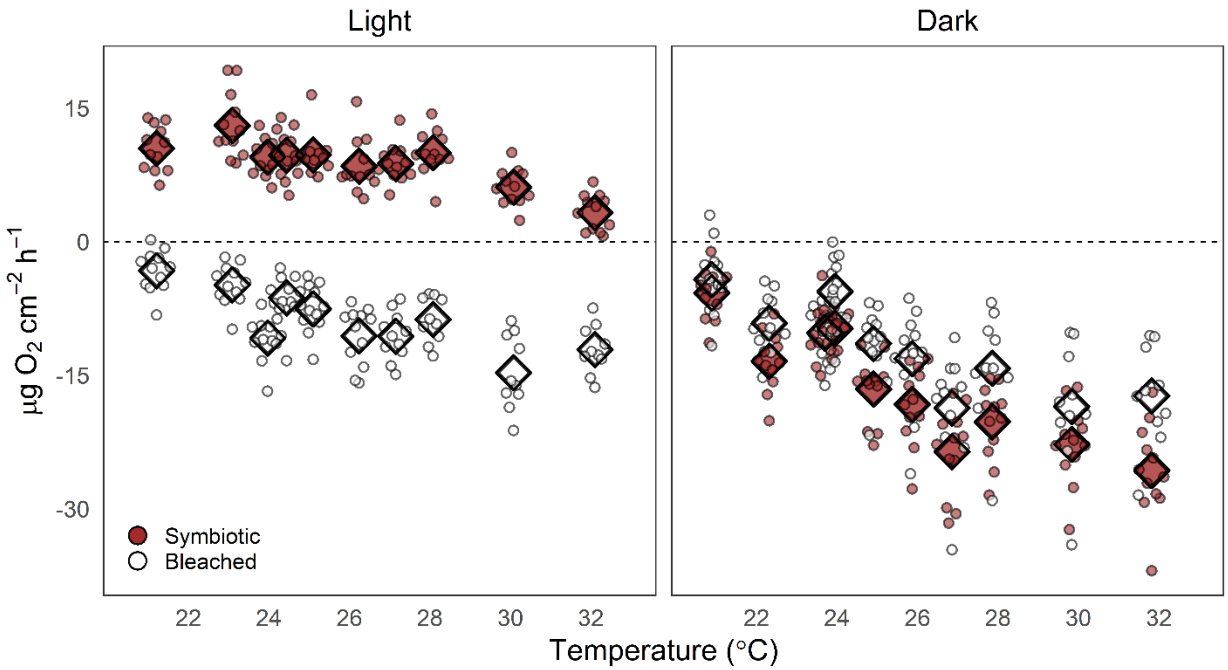


Fig. 1 Microscopic comparison between symbiotic and menthol-bleached polyps. Representative stereo and compound micrographs in bright field and UV light, with filters for chlorophyll (red) and coral tissue (green) autofluorescence. **a** Symbiodiniaceae cells are abundant in symbiotic polyps while not detectable in bleached polyps. **b** At higher magnifications, few algal cells are still detectable in bleached polyps only at the base of the tentacles (arrows)

Amenability to experimental manipulation and broad thermal tolerance

Symbiotic (untreated) polyps had positive net photosynthesis across all temperatures from 21 to 32 °C (Fig. 2). Net photosynthesis peaked at 23 °C (mean 13.05 µg O_2 cm^{-2} h^{-1}) and reached the minimum at 32 °C (mean 3.26 µg O_2 cm^{-2} h^{-1}). In contrast, respiration overall increased with temperature with minimum values at 21 °C (mean 5.77 µg O_2 cm^{-2} h^{-1}) and maximum at 32 °C (mean 25.70 µg O_2 cm^{-2} h^{-1}; Fig. 2). Bleached polyps, as expected by their lack of photosymbionts, exhibited a net oxygen depletion. Similar to symbiotic polyps, respiration in bleached polyps increased with temperature, both in light and dark incubations (Fig. 2). Overall, respiration in symbiotic polyps was higher than in bleached polyps ($P_{lmer} < 0.001$).

316 While net photosynthesis was similar between colonies ($P_{lmer} > 0.05$), respiration was significantly
317 higher in one colony ($P_{lmer} < 0.001$) compared to the other two (RS2-RS1: $P_{Bonf} < 0.01$; RS2-RS3:
318 $P_{Bonf} < 0.001$; RS1-RS3: $P_{Bonf} > 0.05$; Fig. S4).



319

320 **Fig. 2 Net photosynthesis and respiration rate of symbiotic states in *Galaxea fascicularis.***
321 Symbiotic (brown symbols) and Bleached (white symbols) polyps in light and dark incubations.
322 Noise added (jittering) for ease of visualization, means per group depicted as diamonds. n = 12

323 **Symbiosis re-establishment after bleaching**

324 Natural symbiont uptake by placing menthol-bleached polyps next to symbiotic colonies resulted
325 in the re-establishment of the coral algal symbiosis. In the long-term, these polyps grew into small
326 colonies that were visually indistinguishable from untreated corals (Fig. S5a). The degree and
327 speed of repopulation differed between polyps from different colonies. One month after natural
328 uptake, polyps from one colony appeared fully symbiotic while polyps from another colony were
329 only intermediately populated (Fig. 3a).

330 Inoculation with freshly isolated Symbiodiniaceae was followed by symbiont repopulation in three
331 out of seven bleached polyps. These belonged to the same colony and repopulation was
332 independent of the source inoculum (i.e. symbionts isolated from the same or from a different
333 colony). Similar to the successfully repopulated polyps from natural uptake, successfully
334 inoculated polyps grew into small colonies visually indistinguishable from untreated corals one
335 year later (Fig. S5b).
336 Inoculation with cultured symbionts resulted in successful symbiosis re-establishment for 100 %
337 of the samples (n = 16), which were macroscopically visibly symbiotic. Of these, nine samples
338 were successfully FISH hybridized, with the genus-specific probes identifying the presence of both
339 target genera in each sample (Fig. 3b). Individual variation in the relative ratio of each genus was
340 evident among the fragments, with *Durusdinium* contributing to 4 – 63 % of cells and
341 *Cladocopium* to 37 – 82 % (Fig. 3b). A small, but significant proportion of symbionts remained
342 unlabeled in some samples. These may represent additional symbiont genera or be the result of
343 variable probe efficiency.

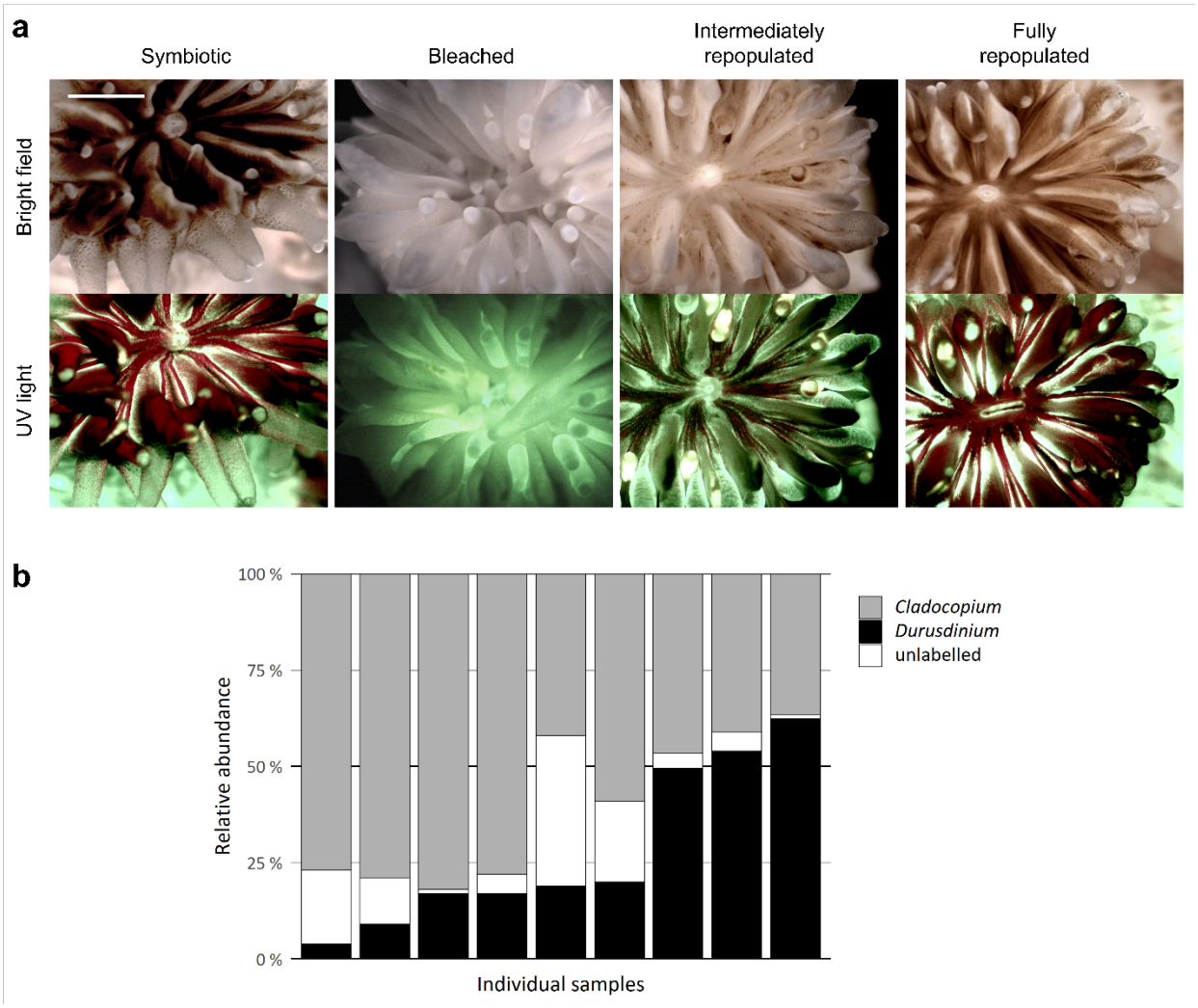


Fig. 3 Symbiosis reestablishment of menthol-bleached polyps following inoculation with Symbiodiniaceae. **a** Representative stereomicrographs of untreated ('Symbiotic') polyp and fully aposymbiotic ('Bleached') polyp 10 days after menthol treatment. One month after inoculation with freshly isolated Symbiodiniaceae, polyps originating from different colonies show different degrees of repopulation. UV light visualizes chlorophyll (red) and host tissue (green) fluorescence. Bright field: natural light (2950 K); UV light + GF2 filter; (Leica, MZ16 F). Scale bar = 2 mm. **b** Relative abundance (%) of *Cladocopium* (gray), *Durusdinium* (black), and unlabeled cells (white) following menthol bleaching and inoculation with cultured algae of these genera. The presence of both genera is confirmed, with individual variation in relative ratios across fragments (n = 9).

354 **Ex-situ spawning**

355 Spawning was observed each year over the three-year observation period for all 35 colonies.
356 Annual fecundity, as a percentage of colonies completing a full gametogenic cycle, was 43.97 %
357 (3.02 sd) (2019: 40.48 %, 2020 and 2021: 45.71 %) during these three spawning cycles.
358 Spawning occurred in synchrony with wild predictions, with gametes being released in November
359 and December each year, between 0.26 - 259 MAS and 5 - 9 NAFM (Fig. 4b-c & e-f). Following
360 embryo rearing, Symbiodiniaceae were observed within primary polyps 14 days post settlement
361 (Fig. 4g). Polyps were fully developed 40 days post settlement (Fig. 4h) and grew into small
362 colonies within the first six months (Fig. 4i).

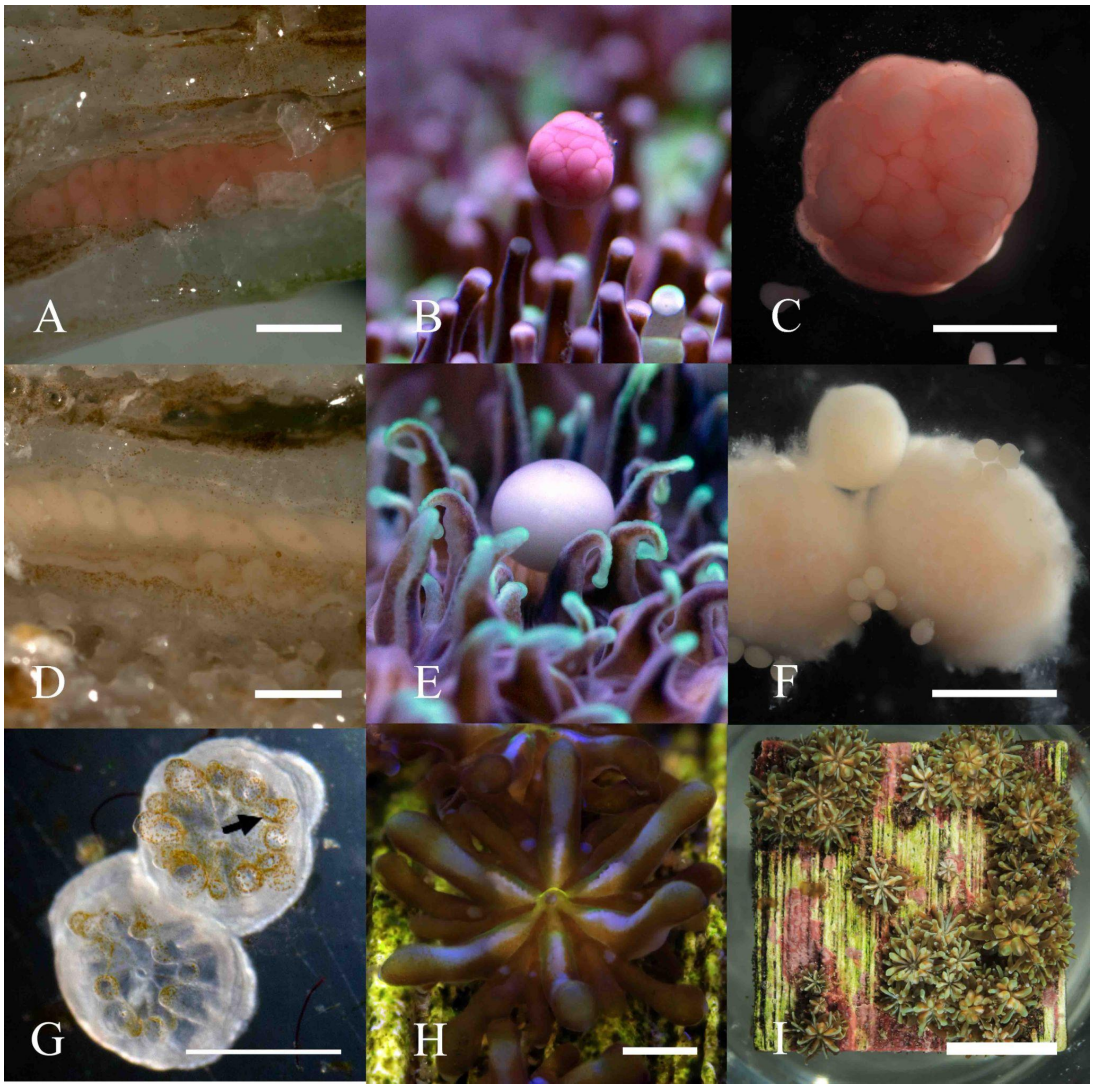


Fig. 4 ***Galaxea fascicularis*** **spawning** ***ex situ***. Representative images of gamete development and spawning in female and male colonies. **a** Longitudinal section of female polyp with pinkish-red pigmented oocytes. **b** Female colony releasing oocyte bundle *ex situ*. **c** Close-up of female bundle post release. **d** Longitudinal section of male polyp with white oocytes. **e** Male colony releasing oocytes/sperm bundle *ex situ*. **f** Close-up of male gamete bundles post release undergoing disassociation. **g** Newly settled *G. fascicularis* primary polyps showing initial Symbiodiniaceae (arrow) uptake 14 days post settlement. **h** Primary polyp 40 days post settlement. **i** Multiple colonies six months post settlement. Scale bar: a-h = 1 mm, i = 1 cm

372 **Discussion**

373 Here, we explored the potential of *G. fascicularis* as a novel coral model by assessing its
374 amenability to aquarium rearing, experimental handling, and symbiosis manipulation.

375 **Successful bleaching with menthol**

376 We showed that *G. fascicularis* can be readily and reliably rendered aposymbiotic through menthol
377 bleaching, similar to other cnidarian model systems (e.g., Matthews et al. 2015; Röthig et al. 2021).
378 We lowered the menthol concentration in the protocol developed by Wang et al. (2012) to further
379 limit stress on the host, and indeed all coral polyps survived this bleaching procedure on three
380 independently replicated occasions. Overall, menthol treatment was very effective, with only a
381 few algal cells remaining at the base of the tentacles. From a photophysiological perspective, these
382 scant remnants can be considered negligible, as illustrated by the light and dark oxygen production
383 data. Furthermore, bleached polyps remained aposymbiotic for weeks and we therefore considered
384 this bleaching protocol suitable for experimental investigation of aposymbiotic *G. fascicularis* in
385 physiological studies and symbiosis manipulation. However, as menthol has photo-inhibitory
386 (Wang et al. 2012; Clowez et al. 2021) and antimicrobial properties (İşcan et al. 2002), further
387 studies should investigate the effects of menthol bleaching on the remaining microbiome.

388 **Tractability of symbiotic and aposymbiotic *G. fascicularis***

389 Tractability, as amenability to experimental work and laboratory conditions, is one of the most
390 important requirements for model organisms. The *G. fascicularis* polyps and fragments could be
391 maintained for three and ten weeks respectively in simple systems consisting of small independent
392 tanks (5-6 L) with regular feeding and basic water quality care. These time frames match the
393 requirements for most experimental designs and we believe could be further extended. Indeed,
394 symbiotic individuals looked healthy and responsive throughout the whole period, testifying to a

395 comparatively hardy coral with only low demands on rearing conditions. Additionally, the positive
396 net photosynthetic rate across a large temperature range confirmed the broad thermal tolerance of
397 this species (Al-Horani 2005; McIlroy et al. 2019), which was retained in the simplified single
398 polyp application.
399 Moreover, bleached polyps could be maintained in the simple experimental system for three weeks
400 even with the additional stress of the thermal experiment. However, they showed signs of
401 deteriorating condition over time indicating that not all nutritional needs were met. Thus one
402 priority is to develop bespoke feed for long-term rearing of bleached coral polyps (Wang et al.
403 2012).

404 **Thermal performance of symbiotic and aposymbiotic *G. fascicularis***

405 Studies that compare symbiotic and aposymbiotic coral holobionts are particularly informative for
406 understanding coral response to climate change, and therefore we investigated the comparability
407 between symbiotic and aposymbiotic *G. fascicularis* polyps. As expected, these differed in their
408 photosynthetic rates, where the latter were net oxygen consumers even under illumination. Overall,
409 symbiotic polyps had higher respiration rates than aposymbiotic polyps, which could be ascribed
410 to the additional metabolic burden of the associated Symbiodiniaceae, as well as a greater
411 metabolic capacity derived from the availability of photosynthates (Muscatine et al. 1981; Al-
412 Horani et al. 2003). For all polyps, respiration rates overall increased with temperature in line with
413 optimum thermal performance of corals and macroalgae from the central Red Sea (27-33 ˚C)
414 (Anton et al. 2020).
415 Interestingly, we observed intraspecific variability in physiological performance. One colony had
416 significantly higher respiration rates than the others (Fig. S4). This manifested exclusively under
417 dark conditions and was therefore more likely linked to host rather than symbiont identity. This

418 variability warrants further investigations to better characterize host genotypes, as in the long term
419 it will be necessary to establish representative host clonal lines.

420 **Symbiosis reestablishment in adult corals**

421 We showed that bleached *G. fascicularis* polyps successfully reestablished symbiosis with
422 Symbiodiniaceae using qualitative and quantitative approaches. The simpler approaches included
423 exposure to symbionts through the water medium (by proximity with symbiotic colonies), and a
424 one-time raw inoculum of Symbiodiniaceae. In the short-term, bleached hosts retained the ability
425 to feed, which facilitates symbiont uptake (Fitt and Trench 1983) and likely contributes to
426 symbiosis re-establishment. In the long-term, these polyps continued to grow into small colonies
427 and became visually indistinguishable from the untreated controls. Therefore, these trials allowed
428 us to qualitatively demonstrate that symbiosis reestablishment of aposymbiotic adults after
429 menthol treatment is possible.

430 Further, the quantitative approach also demonstrated that it is possible to reestablish symbiosis in
431 adult *G. fascicularis* with cultured Symbiodiniaceae. Remarkably, these were likely heterologous
432 strains (different from the natively associated ones) as they were initially isolated from *Acropora*
433 spp. from the GBR (Australia) and inoculated to *G. fascicularis* from Hong Kong. The
434 experimental repopulation of adult corals with cultured heterologous symbionts has only recently
435 been demonstrated (Scharfenstein et al. 2022), and it is particularly relevant in the context of
436 symbiosis manipulations to enhance heat tolerance in reef-building corals (van Oppen et al. 2015).
437 Further assessments of Symbiodiniaceae identity from algal cultures, symbiotic corals, and
438 inoculated corals are however necessary to confirm that this heterologous symbiosis
439 reestablishment occurred in our experiment.

440 Nevertheless, our targeted inoculation highlighted other interesting prospects for symbiosis
441 manipulations. The high thermal tolerance of *G. fascicularis* (Marshall and Baird 2000) has been
442 hypothesized to be linked to its flexible host-algal association, where *Cladocopium*, *Durusdinium*,
443 and their simultaneous co-occurrence have been documented, and with an equatorial-ward shift
444 from *Cladocopium* dominance to *Durusdinium* dominance (Huang et al. 2011, and references
445 therein). At Hong Kong's latitude, and at our rearing temperature, *G. fascicularis* is expected to
446 be exclusively associated with *Cladocopium* (Huang et al. 2011; Zhou et al. 2017a), yet the
447 targeted inoculation produced mixed-genus associations. This shows that targeted inoculation can
448 successfully shift Symbiodiniaceae community composition towards a higher proportion of heat
449 tolerant strains (Silverstein et al. 2015) without thermal treatment. Although the longevity of these
450 partnerships is still unclear, these are promising results for future development of *in hospite* studies
451 on the role of temperature on inter-partner symbiosis dynamics.

452 ***Ex situ* spawning**

453 *Galaxea fascicularis* is a good model species candidate for sexual reproductive work due its high
454 survivorship in aquarium conditions. As demonstrated herein, a combination of seasonal
455 programming and husbandry enabled the species to complete a full gametogenic cycle *ex situ* over
456 multiple years, which closely mimics that observed in the wild (Babcock et al. 1986; Baird et al.
457 2021). Furthermore, we showed *in vitro* fertilization, embryological development, and *ex situ*
458 larval settlement resulting in spat of known ages. As the age at which the species reaches sexual
459 maturity is currently unknown, future studies should focus on this aspect, which is also important
460 for producing an F2 generation and for closing the species life cycle *ex situ* (Craggs et al. 2020).
461 Access to a F2 generation of known parental lineage will provide a platform to conduct
462 experiments in areas such as mapping quantitative trait loci (Zhang 2012) or phenotypic traits such

463 as growth, disease resistance, or thermal tolerance (van Oppen et al. 2015). It also opens
464 possibilities for studies on heritability in symbiont selection, and for rearing of axenic or
465 gnotobiotic coral lineages, paralleling procedures established for *Hydra* (Fraune et al. 2015).

466 **Conclusions**

467 The study of coral-algal symbiosis has been constrained by the difficulties of maintaining corals
468 *ex situ* and of manipulating the coral-algal association to unravel symbiosis functioning. Here, we
469 showed that adopting *G. fascicularis* as a model system can alleviate these limitations. Its
470 demonstrated compatibility to rearing in simplified systems and experimental applications,
471 together with the repeated spawning in aquaria, confirm the high tractability of this coral species.
472 Further, the possibility to readily obtain aposymbiotic individuals suitable for detailed study and
473 symbiosis re-establishment set the stage for exciting future developments in coral symbiosis
474 research which is necessary to understand corals' potential to cope with changing oceans.
475 Besides these strengths, our study also highlights areas worth future efforts on the path to
476 developing the *G. fascicularis* model system further. These include: i) better characterization of
477 the effect of menthol on the rest of the coral microbiome (e.g., bacteria); ii) development of custom
478 feed for long-term maintenance of bleached individuals; iii) assessment of long-term persistence
479 of heterologous Symbiodiniaceae, and testing of a variety of symbiont strains to understand the
480 breadth of symbiont diversity that can be accommodated by the *G. fascicularis* host; iv) further
481 characterization of host colonies and genotypes with the aim of establishing clonal lines, mirroring
482 for example the development of the Aiptasia model; and v) development of collaborative platforms
483 and open-access community resources around this emerging model system to accelerate research
484 and discovery.

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489 the flow cytometry equipment.

490 **Data and code availability**

491 Data and R scripts used for this study are available at:
492 https://github.com/sPuntinG/Galaxea_Coral_Model/

493 **Author contributions**

494 GP and MZ conceived and designed the study. GP, JC, RH, KEE, SM produced and analyzed data.
495 MS, DMB, MZ contributed reagents/materials/analysis tools. GP and MZ wrote the first draft with
496 contributions from all authors.

497 **Conflict of Interest**

498 The authors declare no conflict of interest for this submission.

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