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1 **Gradients of disturbance and environmental conditions shape coral community**  
2 **structure for south-eastern Indian Ocean reefs**

3

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49 temperature, tropical cyclones

50

51 **Abstract**

52 **Aim**

53 To describe, model and assess the relative importance of environmental and climatic factors likely  
54 influencing the regional distribution of coral cover and assemblages with contrasting life histories  
55 and susceptibilities to bleaching.

56 **Location**

57 We compiled the first comprehensive empirical dataset for coral communities in the south-eastern  
58 Indian Ocean (SEIO), incorporating information from 392 sites along the west Australian coast and  
59 offshore atolls/islands across  $\sim 19^\circ$  of latitude.

60 **Methods**

61 We assessed hard coral cover and community composition to genus using point-intercept transects  
62 or point-count analysis of digital images taken along transects. We explored spatial variation in  
63 environmental conditions and in composition of corals with contrasting life histories. After de-  
64 trending the temporal patterns, we assessed the relative importance of environmental metrics to  
65 coral cover, life histories and bleaching susceptibility using a full subsets model-selection approach  
66 with generalised additive mixed models, accounting for both temporal and among site variation.

67 **Results**

68 The distribution of temperature, light, the frequency of temperature anomalies and tropical  
69 cyclones, appear to be drivers of coral community structure. Functional diversity of low to mid-  
70 latitude coral communities may convey some resilience to thermal stress, while higher-latitude  
71 communities dominated by Competitive and Bleaching-Susceptible taxa may lack this functional  
72 resilience. These patterns likely reflect varying historical exposure to cyclones and temperature  
73 anomalies.

74 **Main Conclusions**

75 As evident in recent years, changing background conditions and regimes of disturbance in coming  
76 decades will shift the distribution, functional diversity and resilience of coral reefs throughout the

77 SEIO. The rate and magnitude of environmental change will ultimately determine the future of the  
78 tropical reefs and whether the higher-latitude reefs provide some refuge from climate change. Our  
79 study highlights the need to quantify the distributional properties of key environmental metrics to  
80 better understand and predict reef condition through coming decades.

81

## 82 **1. Introduction**

83 Cycles of disturbance and recovery are a key feature of coral reef ecosystems (Connell,  
84 1978; Rogers, 1993), where multiple diversity-disturbance relationships exist depending on the  
85 interaction between the frequency and intensity of disturbances (Hall et al., 2012). Management  
86 actions aimed at maintaining the diversity, functional integrity and resilience of coral reef  
87 ecosystems are ideally based on understanding how inherent environmental conditions interact with  
88 disturbance regimes to shape coral community structure (Iwamura et al., 2010; Klein et al., 2013;  
89 Maynard et al., 2015a). This type of information is increasingly important, because many reefs face  
90 cumulative threats from a combination of natural and anthropogenic stressors operating at multiple  
91 scales (Hughes et al., 2003, 2010, 2017).

92 Spatial variation in environmental forces produce different ambient conditions, which  
93 influence the distribution of corals and create heterogeneity in the resilience of coral reefs (Richards  
94 & Hobbs, 2014; Graham et al., 2015). Coral communities are routinely structured by temperature  
95 regimes (McClanahan et al., 2007), light penetration (Anthony & Connolly, 2004; Muir et al.,  
96 2015; Sommer et al., 2017), wave energy (Madin & Connolly, 2006; Lowe & Falter, 2014), tidal  
97 amplitude (Richards et al., 2015), sediment delivery and re-suspension (Maina et al., 2013;  
98 Fabricius et al., 2014; Fisher et al., 2015), nutrient dynamics (Kroon et al., 2012), and ocean  
99 currents (Brinkmann et al., 2002; Lowe et al., 2012). For example, areas dominated by relatively  
100 benign conditions associated with shallow, clear waters as well as low wave action and nutrient  
101 loads are often characterised by the proliferation of corals with ‘Competitive’ life-history traits (e.g,  
102 branching *Acropora*), following the definition of Darling et al. (2012, 2013).

103 Acute disturbances, such as extreme temperature anomalies (Selig et al., 2010) and physical  
104 damage from waves associated with tropical cyclones can, however, disrupt normal environmental  
105 conditions and coral assemblages (Harmelin-Vivien, 1994; Fabricius et al., 2008, Beeden et al.,  
106 2015). Similarly, longer-term ecological stressors, such as outbreaks of coral-feeding crown-of-  
107 thorn starfish (Death & Fabricius 2010; Hock et al., 2014) and coral disease (Bruno et al., 2007;  
108 Ruiz-Moreno et al., 2012; Maynard et al., 2015b), can affect the distribution and composition of  
109 corals. If both the supply of propagules and time before the next disturbance are sufficient, recovery  
110 from these disturbances is possible (Sheppard et al., 2008; Lukoschek et al., 2013, Beeden et al.,  
111 2015; Gilmour et al., 2013; Graham et al., 2011, 2015). Major factors that mediate recovery rates  
112 and the impacts of disturbances are local environmental conditions. These conditions can, in some  
113 cases, alter communities by promoting some life-history traits over others (Carreiro-Silva &  
114 McClanahan 2012, Darling et al. 2013, McClanahan 2014a). For example, cooler water at greater  
115 depth (Tyler et al., 2014), or periodic upwelling of cool water (Riegl & Piller 2003), can mediate  
116 the effects of acute warm-temperature anomalies and may produce different coral responses  
117 (McClanahan & Maina 2003). Additionally, the effects of disturbances can be patchy, possibly due  
118 to fine-scale variation in exposure, bathymetry and reef structure interacting with each taxon's  
119 susceptibility to this stressor variation (Harmelin-Vivien 1994, Hoey et al. 2016). Finally, multiple  
120 stressors may combine to either enhance or reduce coral response to disturbance – such as reduced  
121 thermal stress from sea-surface cooling induced by cyclone wind (Manzello et al., 2007, Carrigan &  
122 Puotinen 2011, 2014; Hughes et al., 2017).

123 Making reasonable predictions about the impacts of climate change on corals is thus  
124 expected to benefit from a better understanding of the interactions between local environmental  
125 conditions and large-scale disturbances. A first essential step towards this is to investigate these  
126 interactions where other human impacts, such as fishing and eutrophication, have not significantly  
127 altered reefs. Commercial, recreational and subsistence fishing occur on many reefs within South-  
128 East Indian Ocean (SEIO) and dredging threatens reefs in some areas (Hanley 2011; Fletcher et al

129 2017). However, impacts from these activities are localised, and anthropogenic stress at regional  
130 scales is low compared to many other reefs in the world (Burke et al., 2011). Moreover, the region  
131 is exposed to a wide range of background environmental conditions and large-scale natural  
132 disturbances, making it an ideal area for assessing how these processes influence coral assemblages.

133 Trait-based approaches to classifying organisms can reveal how coral communities, and the  
134 ecosystem services they provide, respond to disturbances (Darling et al., 2012, 2013). For example,  
135 large branching corals provide the structural complexity that supports reef-fish communities  
136 (Graham & Nash, 2013, Rogers et al., 2014). These Competitive corals grow rapidly and often  
137 dominate reefs, but decline rapidly following disturbances like heat stress, cyclones or outbreaks of  
138 predators (Hughes et al; 2017; Shedrawi et al 2017). Moreover, susceptibility of coral to  
139 disturbances varies considerably among taxa, with a meta-analysis of 68 studies revealing that  
140 *Acropora* and *Pocilopora* corals readily bleach following heat stress, whilst many of the faviid  
141 genera are less likely to bleach (Hoey et al., 2016). Thus, some corals have life-history traits that  
142 make them more resilient to disturbances, allowing them to persist over longer timeframes (Darling  
143 et al., 2013; McClanahan et al., 2014a-c). Consequently, understanding the spatial distribution of  
144 coral life-history traits may facilitate the prediction of future changes in community structure  
145 (Darling et al., 2013, Graham et al., 2014, Sommer et al., 2014, Done et al., 2015).

146 Here, we used coral life-history traits (LHTs) to examine regional responses of coral cover,  
147 community structure and bleaching susceptibility along a continuum of environmental conditions  
148 and disturbance regimes of various types. We compiled in situ coral reef survey data collected  
149 between 1998 and 2014 across 392 sites, spanning 19° of latitude, to build the first comprehensive  
150 empirical dataset for coral communities in the SEIO. Specifically, we asked: 1) how coral cover,  
151 coral life histories, and bleaching susceptibility are distributed over time and space; and 2) how key  
152 environmental factors likely shape the coral communities across space. Addressing these questions  
153 provides a basis for identifying resilient reefs and potential refugia from environmental change,  
154 which may be used to inform management of coral reefs.

155

## 156 **2. Methods**

### 157 **2.1 Study locations**

158           The western coastline of Australia forms the south-eastern margin of the Indian Ocean,  
159 covering nearly 19° of latitude in the southern hemisphere (Figure 1). Southward-flowing currents  
160 (Halloway and Leeuwin) push warm tropical water along the length of the coast (Condi &  
161 Andrewartha, 2008; Feng et al., 2008; Lowe et al., 2012), providing conditions favourable for  
162 extensive coral reef growth and development from the north Kimberley region as far south as the  
163 Abrolhos Islands (Veron & Marsh, 1988). Extensive coral reefs are also found on oceanic atolls and  
164 island territories adjacent to the north-west coast of Australia in the SEIO (Speed et al., 2013).

165           Data on percent coral cover and abundance at the level of individual genera were obtained  
166 from nine coastal and oceanic SEIO regions from the west coast of Australia (Figure 1). In each  
167 region, information was collated from 3-26 sites at 1 to 15 m depth that were typically sheltered  
168 from prevailing wind and wave exposure (Table S1). Surveys took place between 1998 and 2014,  
169 and include data about impacts from warm-water anomalies and cyclonic activity (Ceccarelli et al.,  
170 2011; Moore et al., 2012, Pearce & Feng, 2013). We define sites geographically as low- (North of  
171 17°S), mid- (~17-22°S) or high-latitude (~22-29°S; Tab. S1). Hard coral cover and community  
172 composition (identified to genus) were assessed using point-intercept transects, or point-count  
173 analysis of digital images taken along transects (Table S1). Comparative studies indicate that  
174 differences between these methods arise mainly for corals from the genera *Stylophora* and  
175 *Goniastrea*, and estimates of cover from other genera with contrasting growth forms are similar  
176 (Leujak & Ormond, 2007).

177

### 178 **2.2. Environmental data**

179           We examined nine environmental metrics representing potential drivers from 27 variables  
180 (Table 1). Seven metrics were derived from ocean satellite observations and/or modelled databases,

181 including: 1) sea surface temperature [SST]; 2) thermal stress metrics; 3) total suspended matter  
182 [TSM], 4) photosynthetically active radiation [PAR]; 5) tidal range; 6) nutrient concentrations  
183 (chlorophyll-a); and 7) frequency of exposure to extreme winds generated by tropical cyclones. The  
184 final two metrics, 8) depth and 9) physical location (latitude, longitude, isolation), were derived  
185 from in situ data. The nine metrics were specifically chosen for their relevance to physiological  
186 processes, productivity, and stress responses in Scleractinian reef corals (Maina et al., 2008; Maina  
187 et al., 2011). All environmental data, where appropriate, were aggregated to capture long-term (~30  
188 years; mean, median) averages, distribution (skewness and kurtosis), extremes (maximum) and  
189 variability (standard deviation (SD); Table 1). We accounted for potential bias in ocean-colour  
190 constituents by extracting estimates for our sites from a reanalysis database (Morel & Belanger,  
191 2006; Maina et al., 2011) that adjusts values for reflectance bias (Gove et al., 2015).

192 For each site, we obtained weekly SST data for the period 1982-2012 for our SEIO sites at a  
193 resolution of ~4x4 km from coral reefs thermal stress database (CoRTAD), which archives data  
194 from NOAA's Advanced Very High Resolution Radiometer (AVHRR;  
195 <http://www.nodc.noaa.gov/sog/Cortad/>; Selig et al., 2010). Site level SST time series were used to  
196 characterize the distribution (skewness and kurtosis) and variability (standard deviation) of SST at  
197 each site. From the same database, we extracted thermal stress anomalies (TSA) and weekly SST  
198 anomalies (SSTA) that define the spatial and temporal patterns of temperature anomalies associated  
199 with coral bleaching and disease (1982 to 2012; Selig et al., 2010).

200 The bleaching-related anomalies (TSA) occur in the warmest weeks of the year, whereas  
201 disease-related anomalies (SSTA) can occur at any time of year (Podesta & Glynn, 2001; Liu et al.,  
202 2003; Selig et al., 2006; Bruno et al., 2007). Following Selig et al. (2010), TSA is defined  
203 as observed weekly averaged temperature  $>1$  °C warmer than the warmest climatological week (52  
204 climatological weeks averaged over 30 years). Following Selig et al. (2010), SSTA are defined as  
205 observed weekly averaged temperature  $>1$  °C warmer than the weekly climatological value for each  
206 week of the year (over 30 years). Mean SST anomalies (mean SSTA) define the average number of

207 anomalies in any given year. We calculated both the frequency of TSAs (TSA frequency; Table 1)  
208 and SSTAs (SSTA frequency; Table 1) based on the number of anomalies in each calendar year and  
209 cumulatively over the 30-year study (as per Selig et al., 2010).

210 Time series data for total suspended matter (hereafter TSM,  $\text{g m}^{-3}$ ) and chlorophyll-a  
211 concentration monthly (2002-2010) were summarised to median values, distribution (skewness and  
212 kurtosis) and variability (standard deviation). Time series data (monthly; 2002-2010) of  
213 photosynthetically active radiation (PAR) were obtained from the Globcolour database  
214 (<http://hermes.acri.fr/GlobColour>) and summarised to median values, distribution (skewness and  
215 kurtosis) and variability (standard deviation) from the 8-year time series (Table 1).

216 Extreme winds generated during tropical cyclones can build large seas capable of damaging  
217 reefs. A particular coral colony's exposure and vulnerability to damage from such seas depends on  
218 a myriad of local-scale factors (Fabricius et al., 2008), most notably fine-scale bathymetry around  
219 the colony relative to the incoming wave direction during peak conditions. Such data are presently  
220 unavailable for most of our study area. Thus, we derived exposure to tropical cyclone winds as a  
221 proxy for the potential to cause damaging waves, accepting that damage within this zone will be  
222 patchy. We did this from 1985 to 2013 based on the International Best Track Archive for Climate  
223 Stewardship (IBTRACS – Knapp et al., 2010). Cyclone winds were defined as those of gale force ( $17$   
224  $\text{m.s}^{-1}$ ) or higher. These were mapped each day based on the reported or estimated radius of gale winds  
225 using methods detailed in Carrigan & Puotinen (2011). We extracted maximum cyclone days and their  
226 standard deviation per year from the 28-year database across the study area (Table 1).

227 We developed an Isolation Index to quantify each reef's relative potential for larval  
228 connectivity, given its location with respect to neighbouring reefs, assuming that more isolated  
229 coral communities may differ in structure and composition due to limited accessibility to coral  
230 larvae for recovery (Gilmour et al., 2009; Underwood et al., 2009). To measure isolation, we  
231 grouped reef habitat into 122 spatially distinct large-scale reef complexes, using remotely sensed  
232 reef data from the WCMC 2010 database (UNEP-WCMC et al., 2010) and West Australia habitat  
233 maps from the WA Department of Parks and Wildlife (Bancroft, 2003). We calculated the distance

234 in km between all pairs of reef complexes and calculated the Isolation Index as the normalised  
235 graph-theoretic closeness centrality (0 – isolated, 1 – maximum connected; Beger et al., 2010;  
236 Table 1).

237 As the data collected here quantified for the first time both the coral community and broad-  
238 scale environmental features in this region, the spatial variation in environmental conditions was  
239 first illustrated with Principal Components Analysis (PCA) of normalised environmental data  
240 (Clarke & Warwick 2001). Within the groups of environmental conditions (e.g. different measures  
241 of light, sediment, thermal stress; Table 1), a single combined metric was derived for cases when  
242 several metrics were highly correlated ( $>0.7$ ) with each other, resulting in 16 metrics out of the  
243 initial 27. The final metrics used for PCA corresponded to those identified as being the most  
244 important correlates to variation in coral community composition in the Generalised Additive  
245 Mixed Model (GAMM) analyses (Table 2).

246

### 247 **2.3. Coral community data**

248 To evaluate the distribution of coral assemblages across the SEIO, we standardised data to  
249 derive site-level estimates of total coral cover (%), coral life-history trait (LHT) groups (%), and  
250 bleaching susceptibility. Total coral cover was the average of live hard corals observed at each site  
251 for each sampling period. We classified corals into four coral LHT groups – Competitive, Stress-  
252 Tolerant, Weedy and Generalist - according to Darling et al. (2012), but adapted the categories for  
253 genera based on our expertise with Western Australia corals (co-authors ZR, JG, GS) (Table S2).  
254 For genera with species that grouped into different life histories, we assigned coral cover to each of  
255 the represented life histories in proportion to the number of species within each life history that  
256 occur in the Western Australian coral fauna (Veron & Marsh, 1988 sensu Darling et al., 2013).

257 Bleaching susceptibility (BS) of coral communities at each site was based on the relative  
258 abundance (RA) of genus  $i$  in the coral community weighted by a corresponding estimate of its

259 bleaching response ( $BR_i$ ) and summed across all genera in the community (Equation 1;  
260 McClanahan et al., 2007).

$$261 \quad \text{Site bleaching susceptibility} = \sum_i^n (RA_i - BR_i) \quad \text{Equation 1}$$

262 Bleaching responses were estimated by the observed bleaching intensity and mortality of  
263 genera during thermal stress events in the Western Indian Ocean (McClanahan et al., 2007, 2014a;  
264 McClanahan 2014b), which are comparable to bleaching events observed on the Great Barrier Reef  
265 (McClanahan et al., 2004).

#### 266 267 **2.4. Environmental metrics and implications for coral communities**

268 To assess the relative contribution of spatial variation in environmental metrics in  
269 explaining the spatial variability of total coral cover, life histories and bleaching susceptibility  
270 whilst controlling for temporal trends, we adopted a full subsets model-selection approach, where  
271 models were compared using Akaike Information Criterion for small sample sizes (AICc) and AICc  
272 weight ( $\omega_i$ ) values (Burnham & Anderson, 2002). Prior to analyses, all environmental metrics were  
273 tested for collinearity, following Graham et al. (2003). To avoid issues with multicollinearity  
274 among metrics (predictors), we excluded any models where the absolute correlation between the  
275 metrics was greater than 0.28. To limit the maximum complexity of resulting models, we fitted only  
276 models that included up to three metrics (in addition to “null” model terms, see below). Individual  
277 metrics were carefully screened to ensure a relatively even distribution across sites. Three metrics  
278 (Chl a – SD, TSM – SD and TSM median) were transformed to a log scale because they were  
279 highly skewed. TSM – Kurtosis was excluded because it exhibited highly uneven spread across the  
280 study domain. These restrictions reduced the total model set to 360 unique models.

281 All models were fit using generalised additive mixed models, via the GAMM function from  
282 the mgcv package (Wood, 2006) in R (version 3.1.0, R Core Team 2014). GAMM was adopted  
283 rather than linear or non-linear parametric multiple regression to allow for possible non-linear  
284 effects of metrics on the response variable, without needing to define the functional form of each

285 model. Smooth terms were fit using cubic splines (Wood, 2006) and limiting the basis dimension  
286 'k', which controls the degree of flexibility in curve fitting, to a maximum value of 5 to avoid over-  
287 fitting and to ensure monotonic relationships. Percentage cover, rather than raw count data, were  
288 available for analyses, precluding a model using a binomial distribution. Accordingly, the mean  
289 proportional cover values were logit transformed and modelled using a Gaussian distribution. Site  
290 was included in all models as a random effect nested within Region. In addition, the year of  
291 sampling was included in all models as a continuous cubic regression spline to capture broad-scale  
292 temporal trends, with optimal basis dimension (k) identified via cross validation following Wood  
293 (2006). A null model consisting of a random site effect and year was also included in the model set.  
294 The random site effect was not nested within region, as region was collinear with many of the  
295 environmental metrics of interest. Analyses at the genus level were also carried out for genera  
296 occurring at more than 25% of locations (see Table S3).

297 The simplest model within 2 AICc values of the model with the lowest AICc value was  
298 assumed to be the optimal model. To determine the relative contribution of each predictor metric to  
299 the spatial variation in response metrics across the whole model set, we summed the  $\omega_i$  values for  
300 all models containing each predictor metric. The higher the combined weights for an explanatory  
301 predictor metric, the more important it was for the analysis (Burnham & Anderson, 2002).

302

### 303 **3. Results**

#### 304 **3.1. Environmental gradients**

305 The background environmental conditions at the SEIO reefs and their exposure to disturbances  
306 reflected their geographic setting, with variation being high among regions and comparably low  
307 among reefs within regions (Fig. 2; Table 1). Temperature distributions along the inshore reefs of  
308 northwest Australia had negative kurtosis, indicating flat distributions with frequent but modest  
309 deviations from the mean (Table 1). The distributions are less flat offshore, and were even slightly  
310 peaked or centralized in the further offshore reefs at Cocos Keeling Island, suggesting infrequent

311 extreme temperatures. Skewness of temperature data varied among regions. Positive skewness at  
312 mid- and high-latitude reefs indicate that unusually high temperatures occasionally occur, whilst  
313 positive skewness on high-latitude reefs suggest there are occasions when unusually low  
314 temperatures occur. Variation (SD) in SST was highest at Ningaloo Reef, Shark Bay and the  
315 Rowley Shoals, indicating that these sites are exposed to a wide range of temperatures (Table 1).  
316 Sediment (TSM) concentrations were high at the Montebellos and Ningaloo Reef, and  
317 comparatively low at the offshore reefs, particularly the Rowley Shoals, Christmas and Cocos  
318 Keeling islands (Table 1). The kurtosis and skewness of available light (PAR) were negative at all  
319 sites, with the exception of Ashmore Reef, suggesting most reefs are regularly exposed to the same  
320 levels of light, with few extremes. Chlorophyll concentrations were highest at the Montebello  
321 Islands, Ningaloo and Shark Bay, while chlorophyll skewness and kurtosis were positive at all  
322 reefs, indicating extremely high chlorophyll concentrations were sometimes experienced at these  
323 locations, except at the Abrolhos where kurtosis was negative (Table 1). Tidal range and mean  
324 maximum tides were highest at Ashmore Reef, Scott Reef, the Rowley Shoals and the Montebello  
325 Islands (Table 1).

326         There was a clear latitudinal pattern to cyclone activity, which was highest from Ningaloo  
327 Reef in the south to Scott Reef in the north, infrequent at the lowest-latitude reefs (Ashmore Reef,  
328 Christmas Island), and rare at the high-latitude reefs (Shark Bay, Abrolhos Islands – Table 1). In  
329 contrast, thermal stress varied according to both regional and local oceanography, with the highest  
330 frequency of temperature anomalies during the warmest months (TSA) at Scott Reef, Ashmore and  
331 Shark Bay, followed by Abrolhos and Rowley Shoals (Table 1).

332

### 333 **3.2. Coral community patterns**

334         Coral cover and community composition varied through time at all reefs (Fig. 3), influenced  
335 by their regional exposure to cyclones and particularly the impacts of temperature anomalies and  
336 coral bleaching across regions in 1998 and 2011 (Fig. 4). However, this temporal variation differed

337 among coral life-history groups, with large changes observed for the Bleaching-Susceptible and  
338 Competitive groups, and small changes for the Stress-Tolerant group (Fig 3b-d). The coral groups  
339 varied predictably according to their life-history traits (e.g. growth form) and susceptibility to  
340 disturbances, and the genera within groups generally displayed comparable temporal variation,  
341 although there were exceptions (Figs. S1-S3). For example, among the Stress-Tolerant genera,  
342 *Lobophyllia* changed little, but massive *Porites* displayed relatively large temporal change (Fig.  
343 S2). Furthermore, within the Generalist life-history group, the many contributing genera displayed a  
344 range of variation through time (Tab. S3).

345 After accounting for temporal trends, coral cover and the bleaching susceptibility of  
346 communities were similar among the broad regions, with few notable trends (Fig 3a, b; Figs. S4-  
347 S8). Of the life-history groups, only the cover of Weedy corals showed a slight trend, with highest  
348 cover at low to mid-latitudes, and higher-latitude reefs showing very low cover (Fig. 3e; Fig, S7)

349

### 350 **3.3. Environmental metrics and coral communities**

351 After temporal trends were accounted for, spatial variation in total coral cover was best  
352 explained by temperature variation (SST kurtosis) and anomalies (mean SSTA), and exposure to  
353 cyclones (maximum days) (Table 1, Figs. 4 and 5). Total coral cover declined at most sites with  
354 increasing temperature anomalies (mean SSTA) and with increasing exposure to cyclones, and was  
355 highest when SST kurtosis was negative (even spread of temperatures lacking extremes) (Figure 5).  
356 Temperature variation (SST SD) alone explained the small spatial variation in the Bleaching  
357 Susceptibility of the coral communities (Table 2; Figs. 4 and 5), with cover remaining constant  
358 before declining when SST SD was greater than 2.0 (Figs. 4 and 5).

359 Among the life-history groups, the variation in cover after the temporal trends were  
360 removed was often explained by measures of temperature distribution, in addition to the  
361 environmental metrics that reflected the group's susceptibility to disturbances and their life-history  
362 traits (Figs. 4 and 5). For the Competitive corals, there were six models of similar explanatory

363 power (Table 2), but most included exposure to temperature anomalies (SSTA) and cyclones, and  
364 the distribution of available light (PAR skewness) (Figs. 4 and 5). Competitive corals declined in  
365 cover with increasing PAR skewness (more extreme outliers) and SSTA, but there were some sites  
366 with high cover and SSTA (Figure 5). Within the assemblage of Competitive corals, *Acropora* was  
367 the dominant and most typical genus, and a similar pattern of change was explained by their  
368 exposure to temperature anomalies (SSTA) and cyclones (Table S3, Figs. S1-S3).

369 For the Stress-Tolerant corals, three competing models explained their spatial variation in  
370 cover, but all models included measures of their temperature distribution (Figs. 4 and 5; Table 2).  
371 The cover of Stress-Tolerant corals decreased as both SST skewness and kurtosis became more  
372 positive, but there was little change when kurtosis was above -1.0. Within the assemblage of Stress-  
373 Tolerant corals, massive *Porites* was the most abundant genus, and its variation was best explained  
374 by water depth and exposure to cyclones (Figs. S1-S3). The cover of massive *Porites* increased to a  
375 depth of approximately 8 m, and was low in both the absence of cyclones and at intermediate levels  
376 of exposure. In response to cyclone exposure, the variation in cover of massive *Porites*, the  
377 dominant Stress-Tolerant genus, was inverse to that displayed by *Acropora*, the dominant  
378 Competitive genus (Figs. S1-S8; Table S3).

379 Of all the life-history groups, the Weedy corals showed the only latitudinal variation in  
380 cover once the temporal trends were removed. Five competing models in which measures of water  
381 quality were consistently represented best explained this variation, especially the distribution of  
382 available light (PAR) (Table 2; Figs. 4 and 5). Weedy coral cover was lowest at high-latitude reefs,  
383 when PAR distribution was flat or had negative kurtosis ( $< -1.2$ ), or when the distribution of  
384 suspended solids (TSM) was positively skewed (Table 1). The most widespread and typical of the  
385 Weedy corals was *Seriatopora*, whose variation in cover was also best explained by the distribution  
386 of available light (PAR kurtosis, skewness) (Figs. S1-S3; Table S3).

387 The maximum number of cyclone days and depth best explained the variation in cover of  
388 Generalist corals (Table 2; Table S3; Figs. 4 and 5). Cover of Generalist corals was highest at

389 intermediate depths (4-10 m), but varied unpredictably with exposure to cyclones, other than being  
390 highest when maximum cyclones days were low to moderate (4-8 days) (Figures 5). For the many  
391 (12) diverse genera (Table S2) within the Generalist life-history group, there was a corresponding  
392 range in the number of competing models and physical metrics explaining their variation once the  
393 temporal trends were removed (Table S3; Figs. S1-S3). For example, the cover of *Turbinaria*,  
394 *Pocillopora* and *Isopora* predominantly varied in response with their background physical  
395 conditions (Figs. S1-S3). Tides (mean maximum, range) were the dominant metric affecting the  
396 *Turbinaria* cover. *Pocillopora* cover varied with available light (PAR skewness) and water quality  
397 (chlorophyll kurtosis). *Isopora* cover varied with latitude and depths (Figs. S1-S3; Table S3).

398

#### 399 **4. Discussion**

400 Coral communities of the SEIO have varied considerably through time due to episodic  
401 disturbances. Pervasive changes in coral cover were likely a consequence of extreme temperature  
402 anomalies in 1998 and 2011 (Gilmour et al., 2013, Moore et al., 2012; Hughes et al., 2017), whilst  
403 localised impacts within some regions are attributable to cyclones (Speed et al., 2013). Declines  
404 were evident across all coral life-history groups following broad-scale temperature anomalies,  
405 although the magnitude of this impact varied. Declines were most noticeable among bleaching-  
406 susceptible taxa with competitive life histories.

407 Once these broad-scale disturbances were accounted for, our models indicated that coral  
408 community composition across the SEIO was associated with gradients in background  
409 environmental conditions. Importantly, the variance and frequency distribution of environmental  
410 metrics were generally better predictors of coral cover and community composition than the mean  
411 values. Indeed, kurtosis, skewness, and standard deviation of temperature (SST) or water quality  
412 (PAR, TSM) commonly explained observed variation in cover and some life-history traits. This  
413 indicates that models based on mean values that do not consider a full set of environmental  
414 predictors and the distribution of these predictors may not accurately predict coral niches or their

415 responses to disturbances (McClanahan & Maina, 2003, McClanahan et al., 2015, van Hooidonk et  
416 al., 2013; Cacciapaglia & van Woesik, 2015).

417         Our results also indicate that exposure to cyclones may exert a strong effect on coral  
418 distribution in the SEIO, as was found for coral cover across the Great Barrier Reef over the period  
419 1985-2012 (De'ath et al 2012). Tropical cyclones typically damage large branching or plating  
420 colonies, which can lead to a predominance of smaller encrusting or massive Generalist and Stress-  
421 Tolerant corals, as demonstrated for the Great Barrier Reef (Cheal et al., 2017; Massel & Done,  
422 1993, Madin et al., 2012, 2014). Intermittent cyclones maintain community diversity by preventing  
423 fast-growing, competitively-dominant, species from monopolising space (Connell et al., 1997).  
424 Recovery from cyclones may, however, be rapid as asexual fragmentation can facilitate  
425 proliferation of Weedy corals. Such corals were common at mid-latitude reefs. Levels of cyclone  
426 exposure at mid to low-latitude SEIO reefs may rise in future, with an increase in total wind energy  
427 from cyclones predicted for Australia's NW shelf (Emanuel 2006) as the most intense cyclones  
428 become more frequent worldwide (Kossin et al., 2016). If, as predicted, cyclones track further  
429 poleward when at their most intense ( $67\pm 55$  km per decade for the South Indian basin – Kossing et  
430 al 2014), this may also increase exposure of high-latitude SEIO reefs to cyclones. How this  
431 increased exposure will effect SEIO reefs depends on interactions with other stressors. For example,  
432 repeated cyclone exposure combined with overfishing led to severe degradation in the Caribbean  
433 (Gardner et al., 2005).

434         Despite the recent bleaching events along the SEIO, the Competitive corals and their  
435 dominant taxa (Acropora) were common at many sites, suggesting the historic disturbance regime  
436 has not been so severe as to cause their total replacement by Stress-Tolerant and Generalist corals.  
437 Additionally, consistent exposure to a range of water temperatures may confer some resistance to  
438 coral bleaching (McClanahan & Maina, 2003; McClanahan et al., 2007; Ateweberhan et al., 2011).  
439 This may change, however, if the frequency and intensity of warm water anomalies increases  
440 (Hughes et al 2017).

441 Across all study regions, the coral community was composed of taxa susceptible to  
442 bleaching, but the dominance of susceptible taxa declined when temperature variation exceeded 2.5  
443 SD. A study of coral mortality after the 1998 El Niño found that mortality declined as sea surface  
444 temperature variation increased up to ~2.5 SD but increased for variations >2.5 SD, thus producing  
445 a U-shaped mortality curve (Ateweberhan and McClanahan 2010). Consequently, while  
446 background temperature variation and distributions may infer some ability to acclimate to acute  
447 temperatures, there are limits. Extreme temperature anomalies are increasingly likely to reduce  
448 Bleaching-Susceptible taxa and change the structure of SEIO reefs possibly at both the low and  
449 high ends of background SST SD (Halpern et al., 2015; Ainsworth et al., 2016). Indeed, in the last  
450 two decades, abnormally intense warm-water events have affected both high and low-latitude reefs  
451 in the SEIO, of which the 2011 heatwave was the most severe (Abdo et al., 2012; Depczynski et al.,  
452 2013; Feng et al., 2013; Hobbs & McDonald, 2010; Moore et al., 2012; Wernberg et al., 2012;  
453 Zinke et al., 2015; Zhang et al., 2017). Moreover, since 2011, anomalously warm SSTs have caused  
454 persistent summer heat stress and severe coral bleaching at many SEIO reefs (Caputi et al., 2014;  
455 Feng et al., 2015; Lafratta et al., 2016). These stresses are likely to interact with aspects of  
456 background temperature variation to produce changes in coral communities that may not be linearly  
457 related to historical temperature variation.

458 Coral life histories provided a useful approach to understanding how gradients of  
459 environmental conditions and disturbances across the SEIO reefs likely shape reef communities.  
460 Reefs with a relatively higher frequency of environmental disturbances (e.g. cyclones, bleaching)  
461 were characterized by communities with more diverse life histories, whilst at less-frequently  
462 disturbed reefs, Competitive corals were more common. This is in accord with studies showing  
463 Stress-Tolerant, Generalist and fast-growing Weedy corals as more common in disturbed  
464 communities in Kenya (Darling et al., 2013), the Maldives (McClanahan & Muthiga, 2014), the  
465 Red Sea (Riegl & Piller, 2003), the Great Barrier Reef (Graham et al., 2014), and subtropical  
466 Australian reefs (Sommer et al., 2014). Here, we provide the first analysis of how different life-

467 history coral groups respond to putative environmental drivers on reefs with limited exposure to  
468 local human impacts. The patterns of change in the different life-history groups and their key  
469 environmental correlates are also often similar for dominant and most-representative taxa, such as  
470 *Acropora* within the Competitive corals, and the massive *Porites* within the Stress-Tolerant corals.  
471 Nonetheless, taxa within the life-history groups can show substantial variation in their responses to  
472 environmental drivers, suggesting that approaches encompassing life-history traits may benefit from  
473 further refinement. This was especially evident among corals in the Generalist life-history category,  
474 where environmental variables best predicting coral distribution varied among genera.

475

## 476 **5. Conclusion**

477 In summary, we find that a diverse, and possibly more resilient, community prevails at reefs  
478 exposed to regular disturbances. However, extreme, extensive warming events have had a major  
479 impact on the current distribution, cover and community composition of corals in the SEIO, raising  
480 questions about the long-term stability of these patterns. Where and how frequently intense  
481 warming occurs will have a major impact on corals across the region. Reefs at high latitudes that  
482 have historically had little exposure to disturbances could be among the most susceptible to future  
483 climate change because climate impacts, including more intense cyclones (Kossin et al 2014),  
484 penetrate further into subtropical reefs (Hobday & Lough, 2011; van Woesik et al., 2011; van  
485 Hooidonk et al., 2013; Cacciapaglia & van Woesik, 2015). Conversely, functionally diverse corals  
486 at lower latitudes that are regularly exposed to some level of disturbance may be more resilient to  
487 ocean warming and environmental disturbances even as exposure to cyclones increases (Emanuel  
488 2006). Overall, regional compilations of community patterns are important to disentangle the  
489 effects of natural environmental variability. Our results provide a unique perspective on how natural  
490 environmental drivers likely shape coral community structure in the SEIO, providing a reference  
491 point to evaluate ongoing impacts of global change on coral reef ecosystems (Hughes et al., 2017).

492

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507

508 **Biosketch**

509 Between May 13-15<sup>th</sup> 2014, a group of international scientists (Wildlife Conservation Society; The  
510 University of North Carolina) and experts from federal and state government agencies in Western  
511 Australia (AIMS, CSIRO, DPaW, DoF, WA Museum) with colleagues from Australian Universities  
512 (The University of Western Australia, Curtin University, University of Queensland, James Cook  
513 University) met to discuss and develop a new framework that maps the susceptibility of Western  
514 Australian coral communities to chronic and acute thermal and cyclone stress events. The multi-  
515 disciplinary team includes experts in coral reef ecology, management, oceanography and spatial  
516 modeling. Author contributions: J.Z., S.K.W., J.P.G., M.S., M.P., E.D., T.R.M., Z.T.R. and R.F.  
517 conceived the ideas; All authors helped in collection of the data; R.F., M.P., E.D., J.M., M.B.,  
518 J.P.G., Z.T.R. and S.K.W. analysed the data; and J.Z., R.F., J.P.G., M.P. and S.K.W led the writing.

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949 **Supporting Information**

950 Additional supporting information can be found in the online version of this article.

951 **Appendix S1** {Coral genera levels patterns }

952 **Figure S1** { Pooled environmental metric importance (heatmap) }

953 **Figure S2** {GAMM coral genera vs. environmental predictors }

954 **Figure S3** { Community patterns of total hard coral cover and dominant coral genera }

955 **Figure S4** { Bleaching susceptibility spatial pattern }

956 **Figure S5** {Competitive corals spatial pattern }

957 **Figure S6** { Stress-Tolerant corals spatial pattern }

958 **Figure S7** {Weedy corals spatial pattern }

959 **Figure S8** {Generalist corals spatial pattern }

960 **Table S1** {Study locations }

961 **Table S2** { Genera level life-history assignments }

962 **Table S3** {GAMM fits for top models }

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970 **Figure captions**

971

972 **Figure 1** Study sites and regions in the southeast Indian Ocean Reefs (SEIO): A) Cocos Keeling,  
973 B) Christmas Island, C) Ashmore and Scott Reefs, D) Rowley Shoals, E) Montebello Islands, F)  
974 Ningaloo Reef, G) Shark Bay, and H) Houtman Abrolhos Islands.

975

976 **Figure 2** Spatial variation in physical conditions across the southeast Indian Ocean Reefs (SEIO).  
977 Principal Components Analysis of environmental predictor metrics at replicate reefs at each of the 9  
978 coral reef regions. The vectors and environmental predictor metrics (Table 1) responsible for the  
979 spatial separation among reefs are in grey; predictor metric abbreviations are PAR  
980 (Photosynthetically Active Radiation), TSM (Total Suspended Materials), Chl (Chlorophyll a), SST  
981 (Sea Surface Temperatures), SSTA (Sea Surface Temperature Anomalies), TSA (Thermal Stress  
982 Anomalies) and skew (skewness), kurt (kurtosis), med (median), max (maximum), freq (frequency)  
983 and av (average).

984

985 **Figure 3** Temporal and spatial trends in community patterns of total hard coral cover (a), bleaching  
986 susceptibility index (b) and the four coral life-history groups (c-f) following Darling et al. (2012,  
987 2013) across the southeast Indian Ocean Reefs (SEIO). Left hand panels show temporal trends  
988 fitted via GAMM smoothers (see methods) and right hand panels show boxplots of residuals for  
989 each region, once this temporal trend is accounted for (these are effectively de-trended regional  
990 patterns). The box highlights the interquartile range with the mean for each indicated by a solid line.  
991 The whiskers show the maximum range and the open circles are outliers.

992

993 **Figure 4** Variable importance (summed AICc weights) of environmental spatial predictor metrics  
994 in driving summed coral cover, coral groups with contrasting life-history traits (LHT) following

995 Darling et al. (2012, 2013) and the estimate of bleaching susceptibility for the southeast Indian  
996 Ocean (SEIO) Reefs. Environmental metrics are defined in Table 1.

997

998 **Figure 5** Generalised Additive Mixed Model (GAMM) fits for the best models for coral cover (a),  
999 bleaching susceptibility (b) and the four coral life-history groups following Darling et al. (2012,  
1000 2013) (c-f; Table 2). Partial residuals for each smooth term are the residuals that would be obtained  
1001 by dropping the predictor metric concerned from the model, while leaving all other estimates fixed  
1002 (Wood, 2006). Note that all models were fit with year included to ensure temporal trends were  
1003 accounted for. Where multiple models scored within 2 AICc of one another, the most  
1004 “parsimonious” model (least summed estimated degrees of freedom) was plotted. All model plots  
1005 are available in the supplementary material.

1006 **Table captions**

1007 **Table 1** Regional mean of environmental metrics used in this study grouped into nine overarching  
 1008 categories. All environmental metrics included in this study and their mean values are summarised  
 1009 in Supplementary Table 1. The environmental factors analysed in this study include changes in  
 1010 background conditions (e.g. sediment, chlorophyll, temperature, tidal amplitude, light and depth)  
 1011 and periodic disturbances (e.g. extreme temperature anomalies, cyclones).

Latitude		Low				Mid			High	
Environmental metric	units	Cocos Keeling	Christ-mas	Ash-more	Scott Reef	Rowley Shoals	Monte-bello	Ninga-loo	Shark Bay	H. Abrolhos
<b>Location</b>										
Depth	m	5.17	8.71	7.00	6.73	7.26	4.87	1.34	1.75	10.33
Latitude (South)	°	12.11	10.45	12.24	14.02	17.32	20.67	22.57	25.50	28.70
<b>Thermal stress</b>										
SSTA (mean)	No. yr <sup>-1</sup> 30yrs	16.87	15.88	15.89	16.91	18.21	20.98	19.99	20.88	17.78
SSTA (frequency)	No. yr <sup>-1</sup> 30yrs	4.99	4.83	4.64	5.55	6.15	7.67	6.93	7.60	5.83
TSA (frequency)	No. yr <sup>-1</sup> 30yrs	1.35	1.42	2.18	2.51	1.72	1.26	1.57	2.18	1.82
<b>Temperature variability</b>										
SST (kurtosis)	°C	0.19	-0.42	-0.66	-0.82	-1.05	-1.13	-0.81	-0.71	-0.67
SST (skewness)	°C	0.03	-0.23	-0.14	-0.13	-0.07	0.10	0.32	0.25	0.19
SST (SD)	°C	1.05	1.36	1.44	1.52	1.89	2.58	2.09	2.02	1.57
<b>Cyclones</b>										
Cyclone days (mean)	counts/ yr	1.63	0.55	1.30	2.75	3.68	2.39	1.34	0.72	0.03
Cyclone days (SD)	counts/ yr	2.67	1.31	2.31	3.71	3.14	3.28	2.66	1.91	0.18
Cyclone days (max)	counts/yr	11.00	5.00	8.00	14.00	10.52	12.00	10.41	7.90	1.00
<b>Sediment</b>										
TSM (kurtosis)	g cm <sup>3</sup>	1.99	0.64	15.00	16.22	3.03	5.43	7.54	7.62	2.65
TSM (median)	g cm <sup>3</sup>	0.35	0.35	0.45	0.50	0.30	0.74	0.62	0.49	0.42
TSM (skewness)	g cm <sup>3</sup>	-1.01	0.06	0.67	2.61	0.45	1.52	0.96	-0.74	0.20
TSM (SD)	g cm <sup>3</sup>	0.08	0.10	0.09	0.53	0.10	0.36	0.58	0.09	0.12
<b>Light</b>										
PAR skewness	E.m <sup>2</sup> .d	-0.24	-0.25	0.05	-0.10	-0.31	-0.21	-0.21	-0.15	-0.09
PAR kurtosis	E.m <sup>2</sup> .d	-0.94	-0.72	-1.06	-1.11	-1.08	-1.34	-1.36	-1.41	-1.43
<b>Tide</b>										
Average tidal range	m/day	0.68	0.77	1.57	1.59	1.68	1.37	0.78	0.56	0.51
Tide mean maximum	m/day	0.28	0.34	0.92	0.95	1.02	0.79	0.34	0.18	0.14
<b>Nutrients</b>										
Chl (kurtosis)	mg.cm <sup>3</sup>	2.36	1.53	1.01	3.91	2.14	4.69	10.95	0.91	-0.14
Chl (median)	mg.cm <sup>3</sup>	0.11	0.12	0.38	0.36	0.13	0.73	0.62	0.48	0.38
Chl (skewness)	mg.cm <sup>3</sup>	1.48	1.34	0.63	1.09	1.26	1.67	2.59	1.05	0.49
Chl (SD)	mg.cm <sup>3</sup>	0.04	0.07	0.12	0.28	0.05	0.38	0.46	0.18	0.12
<b>Isolation</b>										
normalized centrally		0.01	0.36	0.43	0.56	0.78	0.98	0.86	0.53	0.28
distance (median)	km	2379	1688	1273	1035	621	355	394	660	1012

distance (skewed)	km	-2.10	-0.83	0.21	0.64	1.71	0.85	0.75	0.69	0.46
distance (kurtosis)	km	11.05	6.37	2.78	3.57	6.81	2.82	2.22	2.09	2.34

1012

1013 **Table 2** Generalised Additive Mixed Model (GAMM) fits for best models (the simplest model  
1014 within 2 AICc of the lowest AICc) for environmental predictor metrics influencing changes in  
1015 cover of all corals, and those with contrasting life-history traits (LHT; following Darling et al.,  
1016 2012, 2013) and bleaching susceptibility. Shown are the predictor metrics included in the best  
1017 models, Akaike Information Criterion (AICc), Delta AICc, AICc weight ( $\omega_i$ ) values,  $R^2$ , and the  
1018 number of other competing models within 2 AICc. Best models illustrated in Figure 6 are shown in  
1019 bold. TSM = total suspended matter, PAR = Photosynthetically Active Radiation, TSA = Total  
1020 stress anomaly, Dist = Isolation metric (distance).

1021

LHT	All best models (<2 AICc of min AICc)	AICc	$\Delta$ AICc	$\omega_i$	R2
Coral cover	SSTA (mean)+SST (kurtosis)+Cyclone days (max)	1139.3	0.0	0.492	0.41
Competitive	SSTA (frequency)+Cyclone days (max)	960.7	0.0	0.117	0.39
	PAR (skewness)+SSTA (frequency)	961.1	0.4	0.094	0.35
	SSTA (mean)+SST (kurtosis)+Cyclone days (max)	961.3	0.6	0.086	0.41
	SSTA (mean)+Cyclone days (max)	961.7	1.1	0.069	0.38
	SSTA (frequency)+Cyclone days (max)+log[TSM (median)]	961.9	1.2	0.065	0.42
	PAR (skewness)+SSTA (frequency)+log[TSM (SD)]	962.5	1.8	0.048	0.36
Stress-Tolerant	SST (kurtosis)+SST (skewness)+log[TSM (SD)]	593.1	0.0	0.319	0.42
	SST (kurtosis)+SST (skewness)	593.9	0.8	0.214	0.38
	PAR (skewness)+SST (kurtosis)+SST (skewness)	594.1	1.0	0.193	0.39
Weedy	PAR (skewness)+PAR (kurtosis)	374.7	0.0	0.14	0.37
	Isolation+Tide (mean maximum)	374.9	0.2	0.125	0.35
	Isolation+average tidal range	375.6	0.9	0.09	0.35
	PAR (skewness)+PAR (kurtosis)+TSM (skewness)	375.9	1.3	0.074	0.38
	PAR (kurtosis)+TSM (skewness)	376.4	1.7	0.058	0.36
Generalist	Depth+Cyclone days (max)	701.2	0.0	0.692	0.31
Bleaching-Susceptibility	SST (SD)	-195.2	0.0	0.463	0.30

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