

1 Fine-scale interplay between decline and  
2 growth determines the spatial recovery of  
3 coral communities within a reef  
4

5 **Abstract**

6 As coral reefs endure increasing levels of disturbance, understanding patterns of  
7 recovery following disturbance(s) is paramount to assessing the sustainability  
8 of these ecosystems. Given the slow dynamics of coral reefs and the increasing  
9 frequency of environmental pressures, management strategies focus on under-  
10 standing recovery patterns to drive efforts and actively promote the recovery of  
11 key coral populations. However, the fine spatial scale heterogeneity of coral dy-  
12 namics challenges our capacity to understand recovery patterns at large spatial  
13 scales and guide effective management actions. In this study, we developed a  
14 spatio-temporal statistical model to estimate the long-term trajectories of branch-  
15 ing, plate and massive corals at fine-spatial scales and predict their recovery pat-

16 terns at unobserved locations within a reef. We parameterized the model using  
17 repeated and georeferenced observations from 783 locations during 16 years at  
18 Heron Reef (Great Barrier Reef, Australia). We then developed indicators of  
19 recovery that capture the interplay between coral growth and relative decline  
20 from disturbance(s) across time, space and growth morphology. Our results re-  
21 veal that successful recoveries, expressed in terms of probability, are associated  
22 with minimum growth rate thresholds of 4.3% and 6.4% (absolute growth,  $y^{-2}$ )  
23 for branching and plate corals in reef locations that were impacted by distur-  
24 bance(s) at medium-high levels and historically abundant. As a product of the  
25 data revolution, predictive maps from statistical models support the development  
26 of new indicators that can support the identification of areas of concern to priori-  
27 tise management intervention. They should be integrated into reef management  
28 toolbox along with other approaches.

## 29 **Introduction**

30 Impaired recovery of hard coral communities has mainly been attributed to cu-  
31 mulative disturbances (Halpern et al., 2008; Darling et al., 2013; Osborne et al.,  
32 2017; Vercelloni et al., 2017; Hughes et al., 2018; Ortiz et al., 2018; Mellin  
33 et al., 2019; Vercelloni et al., 2020; Bozec et al., 2022). The poor recovery of  
34 critical communities, including branching and plate coral morphologies, dimin-  
35 ish their function as habitat providers and threatens the high marine diversity

36 associated with these species (Adjeroud et al., 2009; Fisher et al., 2015; Kayal  
37 et al., 2018; Ortiz et al., 2018; Darling et al., 2019). Traditionally, management  
38 strategies have focused on preserving ecosystem resilience (i.e., resistance and  
39 recovery) by reducing chronic pressures, such as nutrient pollution, overfish-  
40 ing, and predators, to enhance coral survival (Gilmour et al., 2013; Mcleod et al.,  
41 2019). More recently, active management interventions such as restoration are  
42 being explored to mitigate future effects of climate change by promoting faster  
43 rates of recovery, controlling chronic pressures and promoting the adaptive ca-  
44 pacity of corals to thermal stress (Anthony et al., 2017). Notwithstanding these  
45 efforts, the effectiveness of management interventions is challenged by the im-  
46 pacts of large-scale climate-driven disturbances that spatially isolate disturbed  
47 from undisturbed reefs across hundreds of kilometres (Dietzel et al., 2021), in-  
48 hibit connectivity and coral recruitment (Hughes et al., 2019) and may accentu-  
49 ate delays in coral reef recovery (Ortiz et al., 2018; Warne et al., 2022). Rapid  
50 assessment of the effectiveness of management interventions is one of the core  
51 challenges that need to be tackled to adapt management strategies in the light  
52 of new environmental regimes, and complex spatial dynamics (Anthony et al.,  
53 2020; Condie et al., 2021).

54 Modern coral reef management approaches propose using advanced technologies  
55 and analytical tools to model coral community coverage across space and time,

56 considering future changes in environmental conditions (Hickey et al., 2020).

57 The final products include predictive maps of reef indicators across management  
58 areas. On the Great Barrier Reef (GBR), predictive maps have been developed  
59 based on diverse information, including drivers of coral dynamics, environmen-  
60 tal gradients, exposure to disturbances from present and future regimes, green-  
61 house gas emissions and data integration from different monitoring programs,  
62 including citizen science and remote sensing (Mumby et al., 2014; De'ath et al.,  
63 2012; Wolff et al., 2018; Mellin et al., 2019; Peterson et al., 2020; Roelfsema  
64 et al., 2021; Bozec et al., 2022). A limitation of these maps is the dependence  
65 on coral reef monitoring data to predict changes at unobserved locations (Bozec  
66 et al., 2022). Uncertainty in predictions arises when monitoring observations  
67 are (1) too scarce in space to allow inferences about new locations, (2) not suffi-  
68 ciently representative across the combination of reef habitats and (3) too narrow  
69 with respect to types and exposures of disturbances and environmental gradients.  
70 This is particularly relevant to coral reef ecosystems when considering the high  
71 spatial heterogeneity of community composition resulting from complex space-  
72 time interactions throughout the time (Cumming et al., 2017). Consequently,  
73 robust estimation of coral cover trends at management scales (i.e., much larger  
74 than monitoring locations) continues to be a challenge (Vercelloni et al., 2017;  
75 Mellin et al., 2019). This motivates the development of spatially-explicit frame-

76 works that can better accommodate the fine spatial scale variability of commu-  
77 nity assemblages, recovery rates and susceptibility to a disturbance and provide  
78 management-ready products to inform effective decision-making.

79 Spatial patterns of coral reef recovery are driven by many variables acting at dif-  
80 ferent spatial scales, including aspects of recent and past disturbance(s) (Connell  
81 et al., 1997; Graham et al., 2011; Ortiz et al., 2018; Mellin et al., 2019), commu-  
82 nity structure and demographic processes (Gilmour et al., 2013; Adjeroud et al.,  
83 2017; Kayal et al., 2018; Holbrook et al., 2018; Darling et al., 2019), and the  
84 environmental climatology of the habitats (Connell et al., 1997; Gouezo et al.,  
85 2019; Castro-Sanguino et al., 2021; Tebbett et al., 2022). Combined, this knowl-  
86 edge enables more accurate prediction of recovery dynamics of different pop-  
87 ulations and communities within a reef. Importantly, it provides more targeted  
88 information to manage recovery progress towards pre-disturbed states and asso-  
89 ciated demographic drivers (Kayal et al., 2018; Mellin et al., 2019; Darling et al.,  
90 2019).

91 However, these spatial patterns are typically estimated at discrete hierarchical  
92 spatial scales, including sites, reefs, habitats and regions to accommodate a high  
93 variability of recovery patterns within a reef (Hughes et al., 2012; Tebbett et al.,  
94 2022). The systematic lack of studies accounting for fine-scale variability high-  
95 lights the need to explore the influence of space in the estimations of coral reef

96 recovery patterns to support model-based decision-support that better aligns with  
97 management goals (Zurell et al., 2022). By representing the space using continu-  
98 ous spatial processes, we allow information sharing between nearby locations  
99 and introduce a dependence (defined as spatial auto-correlation) into the ob-  
100 served data. Considering spatial dependency is key to estimating the spatial scale  
101 of coral recovery patterns and understanding the importance of drivers acting  
102 fine-scale, such as coral larval supply, coral recruitment and herbivory in shaping  
103 those patterns (Fletcher and Fortin, 2018).

104 Here, we developed a spatio-temporal model to (1) estimate long-term trajecto-  
105 ries of three groups of hard corals based on their growth morphology, (2) extract  
106 the spatial auto-correlation from the data, (3) use these information to predict  
107 coral cover at unobserved locations within a reef and (4) develop new indicators  
108 of coral recovery. Since 2002, benthic communities have been monitored annu-  
109 ally at Heron Reef, situated in the Southern Great Barrier Reef (Roelfsema et al.,  
110 2021). These surveys have captured 11 years of coral recovery (2008-2018)  
111 following the impacts of the white syndrome coral disease outbreak and storm  
112 damage. Spatial patterns are estimated using the complete time series (16 years,  
113 2002-2018) of fine-scale observational changes in branching, plate and massive  
114 corals across different habitats. The new indicators capture important aspects  
115 of coral recovery that can be used to assess the potential success of restoration

116 measures and explore alternative management options in the light of new envi-  
117 ronmental regimes.

## 118 **Methods**

### 119 **Heron Reef Benthic Surveys and Geomorphic Zonation**

120 The Heron Island field survey was originally designed to develop annual benthic  
121 habitat maps of coral reef by integrating field data and satellite imagery (Roelf-  
122 sema and Phinn, 2010). The benthic compositions are semi-automatically de-  
123 rived from georeferenced photoquadrat collected at 2-3m interval along a tran-  
124 sect in different zones. Each photoquadrat represents a 1x1 m<sup>2</sup> footprint of the  
125 benthos (see Roelfsema et al. (2021) for additional details on the methodology).  
126 The geomorphic zones define different habitats across a reef that are formed by  
127 physical attributes including depth and wave exposure (Kennedy et al., 2020;  
128 Roelfsema et al., 2021). Using this method, Heron Reef is divided into four geo-  
129 morphic zones (Figure 1a). The northern and southern reef slope areas are char-  
130 acterized by high and low wave intensities, respectively, and depth of 4-7m. The  
131 inner and outer reef flats are shallower areas (0-2m) without influences of wave  
132 exposure.

133 The composition of coral community is estimated using a convolutional neu-

134 ral network and point-sampling methodology (González-Rivero et al., 2020;  
135 Roelfsema et al., 2021). Coral communities are subsequently aggregated into  
136 branching, plate and massive coral types to produce relative abundances of each  
137 of the three types for each photoquadrat. Sub-sites are defined as areas of 100 m<sup>2</sup>  
138 in size in each surveyed year and generated using hierarchical clustering based  
139 on Euclidean distance between geo-located photoquadrats (Roelfsema et al.,  
140 2021). This method of data aggregation has been used previously to forecast the  
141 effects of multiple and intensifying disturbances in the northern GBR (Vercelloni  
142 et al., 2020), the efficiency of management zoning in Indonesia (Kennedy et al.,  
143 2020) and the identification of hotspots of coral cover across Heron Reef using  
144 the same datasets (Roelfsema et al., 2021).

145 The generated result is a total of 783 sub-sites based on an average of 8.7 ( $\pm$  4.6  
146 standard deviation) photoquadrats per sub-site for each of the 16 years (2002-  
147 2018). The abundances of branching, plate and massive corals are then averaged  
148 within each sub-site and year. Observations of coral cover at the sub-site scale  
149 were used to model 16 years of coral changes across the habitats and three forms  
150 of corals within Heron Reef.



## 151 **Spatio-temporal model for coral cover**

152 Recently published studies have shown the importance of the spatial structure of  
153 benthic data to estimate long-term trajectories and highlight the strength of spa-  
154 tial clustering of coral communities (Aston et al., 2019; Ford et al., 2021; Levy  
155 et al., 2018). The typical methods employed in these papers characterize spa-  
156 tial auto-correlation across different forms of corals and reef locations and gain  
157 knowledge about the spatial scales on which ecological drivers are acting. We  
158 developed a spatio-temporal model to go beyond these approaches, with the goal  
159 of reusing the estimated spatial structure to interpolate coral cover at unobserved  
160 locations and thus increase the volume of information to interpret. This aim is  
161 the essence of modern spatial statistical modelling: to estimate spatial patterns  
162 while considering the effects of drivers in species responses, and interpolate over  
163 a continuous spatial field to predict responses at unobserved locations (Lindgren  
164 et al., 2011).

165 A spatio-temporal Bayesian model is developed to estimate the trajectories of  
166 three groups of corals from 2002 to 2018 (Eqs. 1). The cover of branching, plate  
167 and massive corals ( $y_{it}$ ), for observations  $i$  sampled at location  $s_i$  and time  $t$  was  
168 modelled independently. For each model, a Beta distribution is used because  
169 observed values of coral cover are proportions bounded between 0 and 1. The  
170 Beta likelihood is parameterized in terms of a variance  $\phi$  and linked to the linear

171 model components via a logit transformation (Ferrari and Cribari-Neto, 2004).

$$\begin{aligned} y_{it} &\sim \text{Beta}\left(\phi, \text{logit}^{-1}(x_i^T \beta_j + r(s_i, t) + V_i)\right) \\ r(s_i, t) &= \omega \cdot r(s_i, t - 1) + Z(s_i, t), \\ Z(s, t) &\stackrel{\text{ind}}{\sim} \mathcal{GP}(0, K), \quad t = 2002, \dots, 2018 \end{aligned} \tag{1}$$

172 where,  $x_i^T \beta_j$  is a function of an intercept  $\beta_0$  and the four habitats  $\beta_j$  and  $V_i$  inde-  
173 pendent random effects at the sub-site level. The spatio-temporal random effects,  
174  $r(s_i, t)$ , is composed of a first-order autoregressive process,  $\omega \cdot r(s_i, t - 1)$ , in time  
175 and a Gaussian field,  $Z(s, t)$ , that is approximated using a Gaussian Markov ran-  
176 dom field (GMRF) and a covariance kernel. The GMRF is approximated using a  
177 stochastic partial differential equation (Lindgren and Rue, 2015). See Appendix  
178 S1 for more information about the spatio-temporal modelling.

## 179 **Indicators of coral recovery**

180 The years of recovery were set from 2008 to 2018 because no coral loss attributable  
181 to documented disturbances that was reported during this period. Coral recovery  
182 followed the impacts of four years of white syndrome disease outbreak between

183 2004-2008 and storm damage in 2008 (Haapkylä et al., 2010; Roff et al., 2011).  
184 Despite the proximity of cyclone Hamish in 2009, there was no recorded impact  
185 on Heron Reef (Haapkylä et al., 2010).  
186 Preliminary analyses presented in Appendix S2 showed that the use of partial  
187 recovery (80% of pre-disturbance values) instead of the full recovery (100% of  
188 pre-disturbance values) allowed almost double the number of reef locations to be  
189 considered as recovered. This is mostly due to the high pre-disturbance cover of  
190 branching corals in some locations of Heron Reef.

#### 191 **Probability of recovery**

192 The probability of recovery was used to determine the outcome of recovery  
193 while considering uncertainty in the capability of branching and plate corals to  
194 recover from the disturbance(s). For every unobserved location  $\hat{s}_i$ , and coral mor-  
195 phology, the probability of recovery is estimated from the difference between  
196 predicted coral cover at time  $t$  of recovery (2008-2018) and their pre-disturbance  
197 values ( $\hat{y}_{\text{baseline}}$ , Eq. 2).

$$\hat{p}_{\text{rec},t} = \frac{1}{2000} \sum_{k=1}^{2000} I \left\{ \left( \hat{y}_t^{(k)} - 0.8 \hat{y}_{\text{baseline}}^{(k)} \right) > 0 \right\}, \quad t = 2008, \dots, 2018$$

$$\hat{y}_{\text{baseline}} = \max_{t_{\text{baseline}} \in \{2002, \dots, 2007\}} \hat{y}_{t_{\text{baseline}}} \quad (2)$$

198 with 2000 corresponding to the number of draws from posterior distributions of  
 199 estimated coral cover  $\hat{y}_{\hat{s}_i t}$ .  
 200 Recovery was asserted when  $\hat{p}_{\text{rec}}$  was greater than 0.75 at any time during the  
 201 years of recovery and locations  $\hat{s}_i$ . When this condition was satisfied, recovery  
 202 was defined as successful. As such, successful recovery is interpreted as "75%  
 203 chance of recovering to at least 80% of pre-2008 coral cover".

#### 204 **Growth rate**

205 The growth rate corresponds to the absolute growth rate between two consec-  
 206 utive sampling periods because hard corals are growing slowly. It is estimated  
 207 from the posterior distributions of the differences between  $\hat{y}_t^{(k)}$  and  $\hat{y}_{t-2}^{(k)}$  for every  
 208  $k$  draws (Eq. 3).

$$\text{GrowthRate}(t) = \frac{1}{2000} \sum_{k=1}^{2000} \left( \hat{y}_t^{(k)} - \hat{y}_{t-2}^{(k)} \right), \quad t = 2010, \dots, 2018 \quad (3)$$

209 A total of 2000 draws from the predictive posterior distributions were used to es-  
210 timate the growth rate. The average growth rate and associated 95% credible in-  
211 tervals are estimated using the percentiles (50%, 2.5% and 97.5% respectively).

## 212 **Relative decline**

213 The relative decline expressed the amount of coral loss from disturbance(s).  
214 It was estimated using the maximum predicted coral cover pre-disturbance,  
215  $\hat{y}_{\text{baseline}}$ , at every predictive location and its corresponding value in 2008 (Eq.  
216 4).

$$\text{RelativeDecline} = \frac{\hat{y}_{2008} - \hat{y}_{\text{baseline}}}{\hat{y}_{\text{baseline}}} \quad (4)$$

217 The average decline and associated 95% credible intervals were then categorized  
218 into levels of decline (low, medium, high) using the percentiles (50%, 2.5% and  
219 97.5% respectively).

## 220 **Growth rate thresholds**

221 Absolute growth rates and probability of recovery were used to estimate min-  
222 imum growth rate thresholds that ensured recovery. To do this, we developed

223 logistic models for branching and plate corals with probability of recovery as re-  
224 sponse variables transformed into binary data with 1 when  $\hat{p}_{\text{rec}}$  were greater than  
225 75% chance of recovery and 0 otherwise. Logistic models were fit on the reef  
226 slope south locations for the branching corals in 2014 and reef slope north loca-  
227 tions for the plate in 2016. We assumed that these years matched with the begin-  
228 ning of the exponential phase of coral growth as described in (Ortiz et al., 2018).  
229 The thresholds were defined when the probabilities of recovery (i.e >75% chance  
230 of recovering to at least 80% of pre-2008 coral cover) estimated by the logistic  
231 models were greater than 50%.

## 232 **Implementation**

### 233 **Spatio-temporal Bayesian model**

234 Model outputs were used to predict values of coral cover across the entire Heron  
235 Reef. To do this, the reef was divided into 2,384 locations (Figure 1b) and pre-  
236 dictive values and associated uncertainties were estimated for each of these new  
237 locations and surveyed years. The presence of residual spatial and temporal  
238 auto-correlation was tested using the DHARMA (Hartig, 2019) and glmmTMB  
239 (Brooks et al., 2017) R packages. The spatio-temporal model was implemented  
240 using the r-INLA package (Rue et al., 2017) and run on a high-performance  
241 computer. The best model formulations were retained using visual and statis-

242 tical diagnostics including model fit, residual patterns, basis dimensions, dis-  
243 tributional assumption, and Akaike Information Criterion values (AIC). Auto-  
244 correlation tests, computational details of fitting the models, as well as prior  
245 specifications, are provided in Appendix S1.

### 246 **Indicators of recovery**

247 Indicators were estimated at 2384 predictive locations  $s_i$  across Heron Reef but  
248 interpreted within a smaller area on the western side of the reef (Figure 1b). This  
249 area corresponded to the geographical extent of the data and included 481 loca-  
250 tions. This step ensured the detection of signals from the indicators of recovery  
251 due to a higher uncertainty associated with the spatial predictions at locations  
252 further away from the data (see Appendix S1). Indicators were also estimated  
253 at the habitat scale by averaging the values within the same habitat and associ-  
254 ated uncertainty for a given year. The logistic models were developed using the  
255 R package "brms" (Bürkner, 2017) and thresholds were estimated by averaging  
256 model outputs from 100 model iterations (see Appendix S1).

## 257 **Results and Discussion**

### 258 **Temporal changes of coral communities**

#### 259 **Pre-disturbance (2002-2008)**

260 In the period 2002-2008, before the major environmental disturbance(s), branch-  
261 ing corals were abundant in the southern and northern slopes of Heron Reef  
262 (Figure 2a). The maximum coverage is estimated at 37.3% (27.6-47.4%, 95%  
263 CI) in 2004 and 24.9% (17.1-33.8%) in 2006, respectively. A sizeable relative  
264 decline of branching corals is estimated in these habitats, with 93.4% loss (86.8-  
265 97.3%) in four years for the south slope and 83.5% loss (71.8-91.9%) in five  
266 years for the northern slope.

267 Plate corals dominated the northern slopes (Figure 2b) with maximum coverage  
268 of 26.4 % (19.4-34.0%) in 2004, decreasing to a minimum of 7.7% (5.0-11%) in  
269 2010 (70.8 % 58.3-72.7%, relative decline). Studies by (Haapkylä et al., 2010)  
270 and (Roff et al., 2011) described the extirpation of corals, including plate corals  
271 during the major disease outbreak in 2004-2008 at different sites within these  
272 habitats with a shift in coral composition from plate to bushy corals.



273 **Post-disturbance (2008-2018)**

274 Branching corals mostly grew in the sheltered areas of the Reef Slope South  
275 (Figure 2a) that are protected from high wave exposure by adjacent reefs, includ-  
276 ing Wistari, Sykes, and One Tree Reefs (Connell et al., 1997). In this habitat,  
277 the model estimates a relative increase of 1527 % (720-1587%) from 2008 to  
278 2014, with a maximum amount of cover of 40.2 % (31.9-48.7%). Branching in  
279 the northern slope increased to a maximum of 13.8% (9.0-19.3%) in this period.  
280 Plate corals grew in the southern and northern slopes of Heron Reef to reach  
281 maximum values that were higher than pre-disturbance values (Figure 2b). The  
282 prominent increase for plate corals occurred on the Reef Slope North, which  
283 is more exposed to frequent and higher intensity waves than the southern reef  
284 slope. In the northern slope, plate corals increased to 27.1% (20.9-33.7%) in  
285 2018 and 18.1% (13.3-23.5%) in 2016 for the south habitat.  
286 Through the years of recovery, coral community composition changed with a net  
287 dominance of plate corals in the northern section of the reef. The lack of recov-  
288 ery for branching corals cannot be explained by the nature of the disturbance as  
289 there is no clear evidence of what had impacted this reef section (Haapkylä et al.,  
290 2010; Roff et al., 2011). Differences in wave exposure between north and south  
291 and the high prevalence of branching corals in the southern reef slope before the  
292 decline may have contributed to the rapid southern recovery due to their strong

293 capability of recovery after fragmentation (Lirman, 2000). The study from (Con-  
294 nell et al., 1997) indicated differences between mechanisms of declines and re-  
295 coveries in northern and southern sections of Heron Island. Our model estimates  
296 suggest that plate corals were able to recover from disease outbreaks, but this is  
297 less evident for branching corals on the northern slope. Coral colony size is an  
298 important factor associated with this type of disturbance (Roff et al., 2011), but  
299 the size was not recorded as part of the Heron survey.

## 300 **Spatial indicators of coral recovery**

### 301 **Probability of recovery and associated growth rate**

302 Coral recovery is defined when a reef location reached 80% of its pre-disturbance  
303 value with the pre-disturbance value corresponding to the maximum estimated  
304 cover before 2008. As expected, branching corals were abundant in the south-  
305 ern slope from 2002 to 2005, then decreased until 2008 and recovered gradually  
306 until 2018 (Figure 3a). The rapid recovery is reflected by the dynamics of the  
307 probability of recovery showing less than 75% chance of recovery from 2008  
308 (36.6%, 12.4 - 64.0%) to 2012 (62.6%, 13.9 - 86.7%) and then high probabili-  
309 ties of recovery (greater than the threshold of 0.75) from this year (Figure 3b).  
310 In these locations, the growth rate increased from 4.8% (2.2 - 6.8 %  $y^{-2}$  between  
311 2010-2012 to a maximum of 10.4% (7.5 - 13.3 %  $y^{-2}$ ) until 2014 (Figure 3c).

312 Thereafter, the growth decreased to negative values reaching 4.5 % (6.8 - 2.4%  
313  $y^{-2}$ ) in 2018. This decline in branching growth is likely related to a reduction in  
314 space availability during the years of recovery. The probability of recovery re-  
315 mained low for the northern slope habitat (Figure 3b), associated with a negative  
316 growth rate at the beginning and end of the surveyed years (-4.4%, -6.1 - -2.9%  
317  $y^{-2}$  in 2008 and -1.3%, -2.9 - -0.1%  $y^{-2}$  in 2018) and a positive growth rate rang-  
318 ing between 1-2% between these years (Figure 3c).

319 Plate corals recovered in the slope habitat in the north and south during the sur-  
320 veyed period (Figure 4a). On the south slope, the probability of recovery was  
321 estimated at 98.3% (86.8 - 99.8%) in 2008 and remained above this value until  
322 2018, despite a low growth rate of 3.0% (1.2 - 7.8%) estimated in 2014 (Figure  
323 4b). On the north slope, the probability of recovery was almost zero for the first  
324 four years of the recovery phase and increased rapidly between 2012 and 2014  
325 to reach more than 75% chance of recovery in 2014-2018 (Figure 4b). In these  
326 locations, the growth rate increased from - 1.3%  $y^{-2}$  (-2.2 - -0.1%  $y^{-2}$ ) in 2008  
327 to 12.2%  $y^{-2}$  (8.7 - 15.8%  $y^{-2}$ ) in 2014 before being negative again (-0.2%  $y^{-2}$ ,  
328 -2.0 - 0.9%  $y^{-2}$ ) in 2016 (Figure 4c). This rapid increase is the signature of plate  
329 corals that use the first years of recovery to build the base of their colony and  
330 then expand exponentially due to high growth rates and potential large sizes as-  
331 sociated with their growth morphology (Ortiz et al., 2021).

332 **Growth rate, relative decline and cover baseline**

333 Overall, branching corals recovered (to within 80% of baseline) at 53.4% of lo-  
334 cations across the north and south reef slopes in 6.90 years ( $\pm 1.25$  SD). The  
335 fastest recovery occurred in six years at 20 locations within the south slope, and  
336 the longest recovery was estimated at ten years at three locations on the north  
337 slope. Most of the reef locations recovered on the south slope (87.1%) com-  
338 pared with 12.9% of locations on the north slope (Figure 5a). On the south slope,  
339 locations that recovered were medium (15-37% relative decline) to highly (>  
340 37% relative decline) impacted by the disturbance(s) (Figure 5b). The cover  
341 baseline was mostly high (> 40% branching cover) and medium (20-40% branch-  
342 ing cover, Figure 5c). Branching corals at most of the locations grew above  
343 the growth rate threshold estimated at  $4.30\% \text{ y}^{-2}$  ( $\pm 0.01$  SD) from the logis-  
344 tic model in 2012 and 2014 (Figure 5d, Appendix S2). Only four years of high  
345 growth rate were needed to recover branching corals in previously abundant  
346 places and medium-highly impacted by disturbance(s). The disturbances highly  
347 impacted branching corals in the north slope (Figure 5b). The low baseline abun-  
348 dance of branching corals in these locations (Figure 5c) is likely related to the  
349 low growth rate (Figure 5d) due to environmental conditions that are not favourable  
350 for this form of corals (Connell et al., 1997; Tanner, 2017). The high decline  
351 in branching (> 37% relative decline) in association with a low growth rate re-

352 sulted in a lack of recovery for most of the locations in the north slope. However,  
353 the medium-high relative decline did not interfere with the recovery of branching  
354 corals in the southern slope because environmental conditions are prone to a fast  
355 growth rate, as shown by the medium-high cover baseline.

356 The recovery of plate corals was more generally spread across the north and  
357 south slopes, with 91.4% of locations showing recovery in 6.87 years ( $\pm 2.43$   
358 SD). The fastest recovery occurred in only two years at eight locations in the  
359 south slope but it took a maximum of ten years to recover for six locations. More  
360 locations recovered in the south slope (58.5%) compared to the north slope (41.5%,  
361 Figure 6a). The highest decline occurred in the north slope with some loss esti-  
362 mated at more than 58% (Figure 6b) in locations with medium (13-27%) and  
363 high ( $> 27\%$ ) cover baseline (Figure 6c). Plate corals at locations the most im-  
364 pacted by the disturbance(s) grew the fastest with a maximum growth rate es-  
365 timated around  $20\% \text{ y}^{-2}$  in 2016 and above the growth rate threshold of  $6.31\%$   
366  $\text{y}^{-2}$  ( $\pm 0.02$  SD) in 2014 (Figure 6d, Appendix S2). Plate corals at locations that  
367 were impacted by disturbance(s) at a medium level grew above this threshold in  
368 2016, and locations impacted at a low level never reached the threshold. These  
369 results emphasize the interplay between relative decline and growth rate by cre-  
370 ating free space for new plate recruits to settle after the disturbance(s) in 2008  
371 and grow until saturation in 2018. In a paper dedicated to tabular *Acropora*,

372 Ortiz et al. (2021) describes the complex mechanism of coral settlement that is  
373 enhanced by the presence of dead colonies. Plate corals were mostly impacted  
374 by coral diseases that are known to preserve the dead skeletons of the corals on  
375 which algae and potential new recruits can settle. They also show high potential  
376 connectivity from locations that were less impacted by the disturbance(s) and  
377 favourable environmental conditions suggested by the high and medium levels  
378 in the cover baseline before the disturbance(s). The last point has also been rele-  
379 vant for the branching corals, where the recovery occurred in locations that were  
380 historically abundant.

### 381 **Spatial auto-correlation during coral recovery**

382 The presence of spatial and temporal auto-correlation in the long-term data of  
383 Heron Reef has been demonstrated by (Connell et al., 1997). In this study, we  
384 refined this knowledge by quantifying spatial dependence during the recovery  
385 of different forms of corals after accounting for the habitat effect defined by ge-  
386 omorphic zones. Our findings show that the spatial auto-correlation is within  
387 the same range as that estimated by (Connell et al., 1997) (up to 800m) but that  
388 this varies by coral morphology. Spatial ranges were estimated as 1.02km (0.75  
389 - 1.30km, 95% CI), 1.46km (1.07 - 1.90km) and 12.76km (7.24 - 18.97km) for  
390 branching, plate and massive corals, respectively (Appendix S1). The presence

391 of spatial auto-correlation is explained by the mechanisms of coral recruitment  
392 that are related to the supply of larvae from plankton, suitability of substrate for  
393 coral settlement and mortality post-settlement (Connell et al., 1997). Among  
394 these factors, only the supply of larvae from plankton has the potential to act at  
395 the kilometre scale in conjunction with the interaction between habitat and cur-  
396 rents that drive the number of new recruits in these areas (Tebbett et al., 2022).  
397 Connell et al. (1997) and Tanner (2017) suggested that the supply of plankton  
398 is likely to be higher on the southern slope of Heron Reef due to the proxim-  
399 ity to other reefs, which increases coral recovery rates. However, we found that  
400 branching corals drove the recovery on the south slope and plate corals on the  
401 north slope. Further investigations should focus on the interactions between  
402 branching and plate corals within habitats to better understand the influences  
403 of connectivity, demographic traits of coral forms and competition for space in  
404 driving recovery patterns.

405 Analytical approaches to improve the effectiveness of long-term monitoring and  
406 increase the amount of knowledge extracted from the data have been applied  
407 to coral reefs (Kang et al., 2016; Thilan et al., 2019; Mellin et al., 2020). The  
408 quantification of spatial auto-correlation is informative for the design of coral  
409 reef monitoring (Hamilton, 2013). Using a combination of spatio-temporal  
410 modelling and fine-scale data, the estimated values of spatial ranges reveal that

411 surveyed locations separated by 1km and 1.5km can be considered pseudo-  
412 replicated observations for branching and plate corals during recovery, respec-  
413 tively. The presence of redundant information in monitoring data violates the  
414 assumption of independence between observations in traditional statistical tests,  
415 which may bias ecological interpretations (Ver Hoef et al., 2018) about drivers  
416 of coral recovery. We recommend that future surveys that aim to understand  
417 mechanisms underlying coral recovery should ensure that the survey includes  
418 locations within and between a radius of 1.5km allowing for replications and col-  
419 lection of data that are not spatially auto-correlated. Locations should also be  
420 sampled across different habitats, especially in more than one flank of the reef  
421 slope. In this way, monitoring surveys can provide an optimized amount of eco-  
422 logical and spatial information about the recovery patterns of the entire coral  
423 community.

#### 424 **Spatio-temporal modelling for coral reef data**

425 Analyses of the influence of spatial variation during years of recovery allowed  
426 us to identify spatial patterns of recovery for different types of corals within a  
427 reef. The application of spatio-temporal models to this unique dataset reveals  
428 that consideration of interactions between space and time is essential in order  
429 to predict recovery patterns and investigate the fine-scale variability of coral



430 dynamics (Appendix S1). Such interactions are challenging to compute, even  
431 using Bayesian approaches, but improvements in the field of computational sci-  
432 ence and applied statistics will ease their inclusion in future statistical modelling  
433 frameworks (Wikle and Zammit-Mangion, 2022). Moreover, these computa-  
434 tional improvements will enable the scaling-up of the approach to more than one  
435 reef.

436 We acknowledge that the high volume of data needed in order to fully exploit the  
437 benefits of spatio-temporal statistical models is another limiting condition in the  
438 field of coral reef research. Additional work is needed to estimate the minimum  
439 amount of data that will enable the implementation of spatio-temporal models to  
440 more than one reef. The development of new coral-reef monitoring techniques,  
441 including the use of machine learning (González-Rivero et al., 2020), citizen sci-  
442 ence (Santos-Fernandez et al., 2021) and combined approaches (Peterson et al.,  
443 2020) to boost collection, processing and exploration of reef data and their de-  
444 mocratization are rapidly being adopted by research and governmental institu-  
445 tions across the Indo-Pacific. In combination with advanced modelling tech-  
446 niques able to handle large and complex datasets, outputs from data-driven ap-  
447 proaches should be systematically integrated into the reef management toolbox  
448 (Zurell et al., 2022). They are keys to providing rapid and up-to-date information  
449 to reef managers, supporting the development of adaptive strategies and assess-

450 ment of management interventions.

## 451 **Conclusions**

452 The spatial mismatch between the large spatial scale of climate-driven distur-  
453 bances and the finer spatial scale of management interventions (Cumming et al.,  
454 2017; Bellwood et al., 2019) forces the development and implementation of new  
455 types of measures to support coral recovery (Anthony et al., 2020). In this study,  
456 we developed new indicators of coral recovery that have been estimated based  
457 on the fine spatial scale variability of coral changes within a reef and the spatio-  
458 temporal structures of data. The resulting predictive maps of indicators of coral  
459 recovery across Heron Reef show clear zonation of recovery probabilities that  
460 is different between coral morphology and related to the decline from distur-  
461 bance(s) and historical abundance.

462 If management interventions ignore this information and only use locations  
463 where long-term monitoring sites are located, the benefits of interventions may  
464 be lost because of uninformed spatial prioritization (Anthony et al., 2020). This  
465 is especially important considering that existing monitoring only represents 40%  
466 of the environmental regimes of the GBR (Mellin et al., 2020). Importantly, be-  
467 cause the scale for management of local stressors is only a fraction of the global  
468 scale of influence of climate change, managers are likely to be forced to increas-

469 ingly consider prioritization of reef areas with high intrinsic resilience capacity  
470 (GAME et al., 2008). Predictive maps from spatio-temporal models have the po-  
471 tential to fill a gap by gathering information from existing knowledge underlying  
472 coral recovery, learning from data to infer at unobserved locations and develop-  
473 ing useful indicators for decision-making.

474 Data Science, including the combination of machine learning algorithms to  
475 rapidly process a large amount of information and statistical modelling to de-  
476 velop robust ecological knowledge, has the potential to radically change the way  
477 of managing coral reef. The use of this approach is still in its infancy in Aus-  
478 tralia, but current research efforts bridge the gap between data scientists, coral  
479 reef ecologists and reef managers to provide more comprehensive information  
480 about the decline of the condition of coral reef habitats in the Great Battier Reef  
481 and support the development of mitigation interventions under future climate  
482 scenarios. Importantly, while there is an increasing trend of incorporating cli-  
483 mate change into spatial prioritisation, serious gaps still exist in current method-  
484 ologies (Jones et al., 2016). This study provides a contribution to this increas-  
485 ingly challenging field by developing methodologies that recognise discrete and  
486 long-term impacts on ecosystem recovery potential. Our approach can be easily  
487 integrated into broader spatial prioritisation frameworks that respond to spatial  
488 and temporal scales of the processes being managed.

## 489 **Data archiving statement**

490 Data are already published and publicly available, with those items properly  
491 cited in this submission. Data sets utilized for this research are as follows: <https://doi.pangaea.de/10.1594/PANGAEA.907025> (Roelfsema et al.,  
492 <https://doi.org/10.6084/m9.figshare.14034320.v1>  
493 (Roelfsema et al., 2019) and <https://doi.org/10.6084/m9.figshare.14034320.v1>  
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## Figure captions

Figure 1: Locations of the data and spatial predictions. a) Dots show the locations of surveyed 100m sub-sites across Heron Island Reef and geomorphic zones b) Predictive locations used in the spatio-temporal model. The boxed area indicates the area used to interpret the indicators of coral recovery. Data available from the Pangea Digital Repository: <https://doi.pangaea.de/10.1594/PANGAEA.907025>.

Figure 2: Long-term trajectories of cover estimated by the model at the habitat scale for a) branching, b) plate, and c) massive corals. The dots and error bars denote the observed values and associated 95% confidence intervals. The line and shaded areas are the model estimates showing the mean and 95% credible intervals estimated from the model posterior distributions. Note that the y-axis is on a different scale for the massive corals in panel c.

Figure 3: Spatial and temporal changes in branching corals. a) Predicted coral cover estimated at unobserved locations between 2002-2018, b) Temporal changes of the probability of recovery at the habitat scale. The dotted line shows the threshold of successful recovery and c) Estimated growth rate at the habitat scale across years.

Figure 4: Spatial and temporal changes in plate corals. a) Predicted coral cover

estimated at unobserved locations between 2002-2018, b) Temporal changes of the probability of recovery at the habitat scale. The dotted line shows the threshold of successful recovery and c) Estimated growth rate at the habitat scale across years.

Figure 5: Indicators of recovery for branching corals. a) Predictive locations on the reef slope on which branching corals recovered or not using the 75% chance of recovery across 2008-2018 as threshold, b) Associated levels of relative decline estimated from the overall distribution within the reef slope of the small area, c) Associated levels of baseline cover estimated from the overall distribution within reef slope of the small area and d) Temporal changes of the branching absolute growth rate in  $y^{-2}$  between 2010 and 2018 for the south reef slope. Line colours denote the presence and absence of recovery at the predictive locations, and the solid black line shows the growth rate threshold of recovery estimated by the logistic model.

Figure 6: Indicators of recovery for plate corals. a) Predictive locations on the reef slope on which branching corals recovered or not using the 75% chance of recovery across 2008-2018 as threshold, b) associated levels of relative decline estimated from the overall distribution within the reef slope of the small area, c) Associated levels of baseline cover estimated from the overall distribution within reef slope of the small area and, d) Temporal changes of the branching absolute growth rate in  $y^{-2}$  between 2010 and 2018 for the south reef slope. Line colours

denote the presence and absence of recovery at the predictive locations, and the solid black line shows the growth rate threshold of recovery estimated by the logistic model.

# Figures

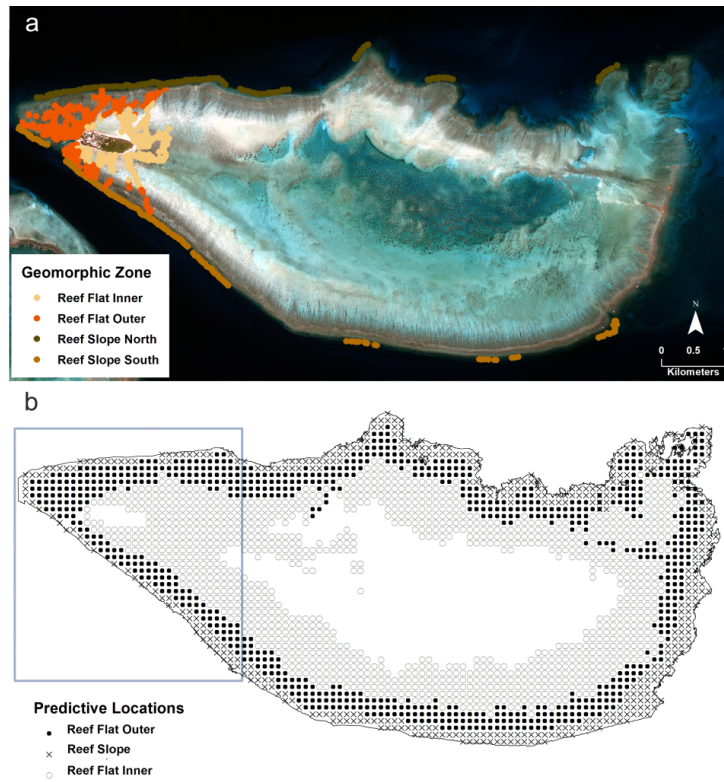


Figure 1

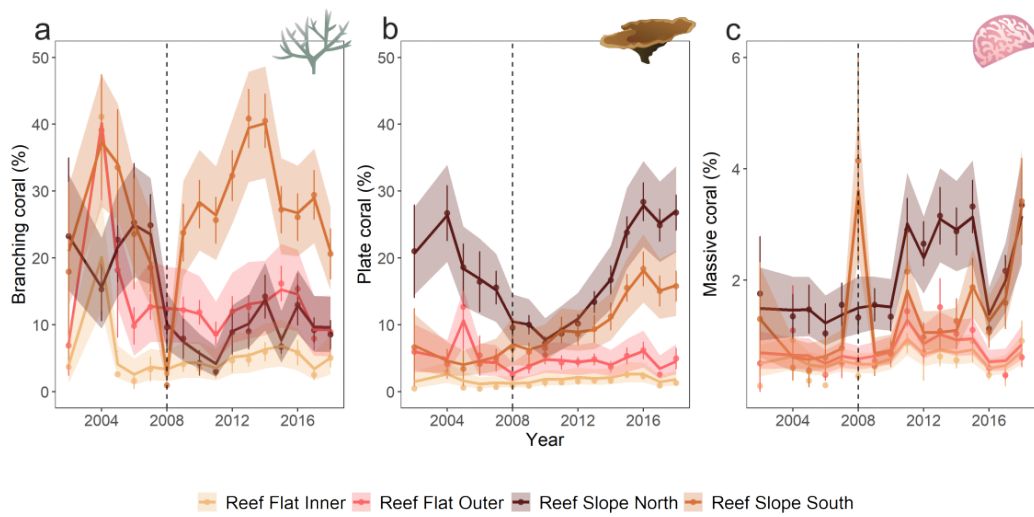


Figure 2

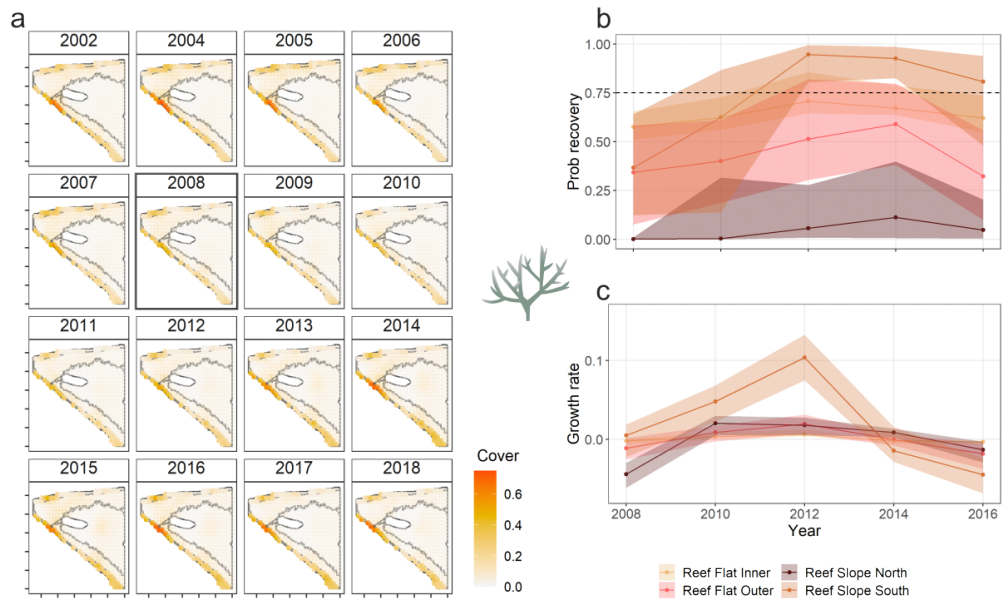


Figure 3

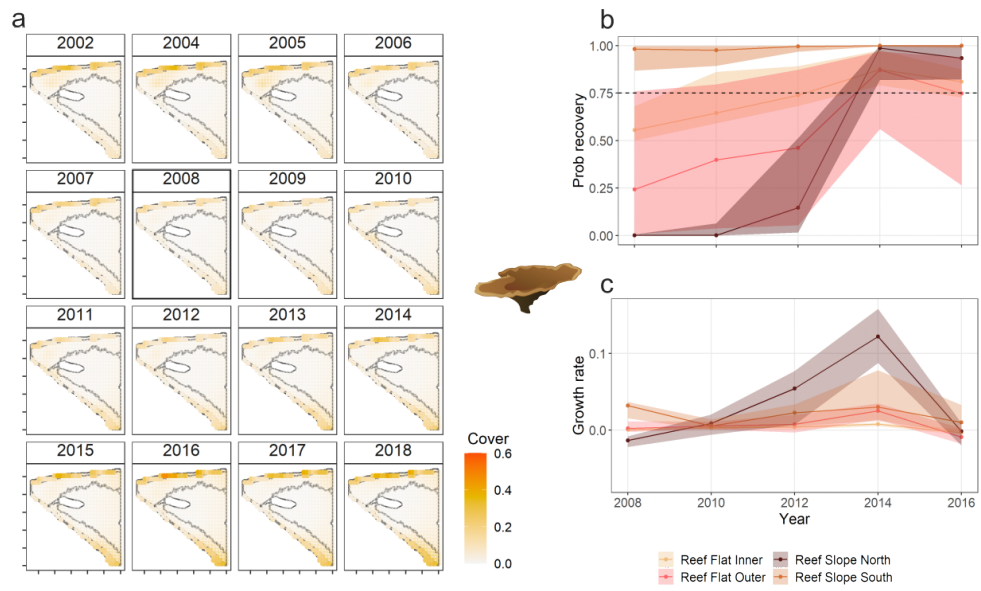


Figure 4

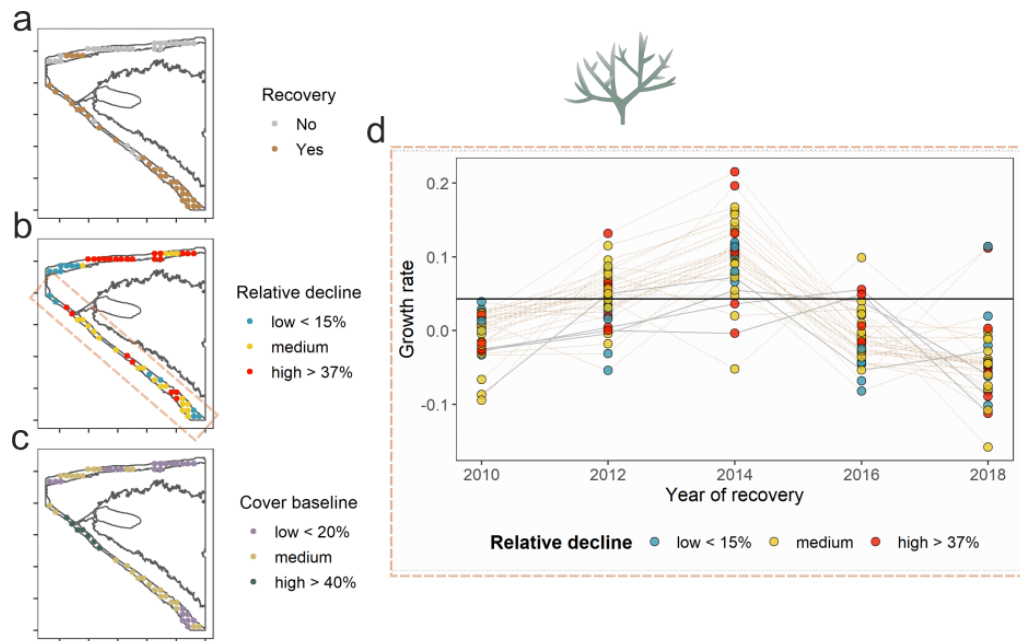


Figure 5



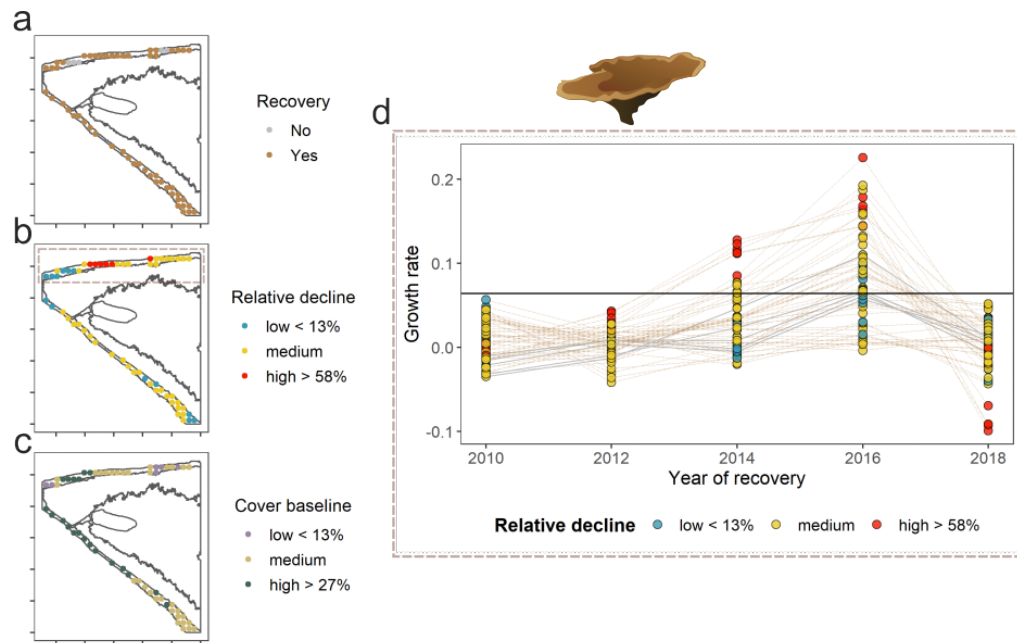


Figure 6