

1 **Growing at the limit: reef growth sensitivity to climate and oceanographic changes**  
2 **in the South Western Atlantic**

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24 **Abstract**

25 Whilst the impacts of climatic and oceanographic change on lower latitude reefs are  
26 increasingly well documented, our understanding of how reef-building has fluctuated in  
27 higher latitude settings remains limited. Here, we explore the timing and longevity of  
28 reef-building through the mid- to late Holocene in the most southerly known reef (24°S)  
29 in the Western Atlantic. Reef core data show that reef growth was driven by a single coral  
30 species, *Madracis decactis*, and occurred over two phases since ~6,000 calibrated (cal.)  
31 yr B.P.. These records further indicate that there was a clear growth hiatus from ~5,500  
32 to 2,500 cal. yr B.P., and that there is no evidence of reef accretion on the Queimada  
33 Grande Reef (QGR) over the past 2,000 yrs. It thus presently exists as a submerged  
34 senescent structure colonized largely by non-reef building organisms. Integration of these  
35 growth data with those from sites further north (18°S and 21°S) suggests that Intertropical  
36 Convergence Zone (ITCZ), South Westerlies Winds (SWW) and El Niño-Southern  
37 Oscillation (ENSO) variability and shifts during the Holocene drove changes in the  
38 position of the Brazil-Falklands/Malvinas Confluence (BFMC), and that this has had a  
39 strong regional influence on the timing and longevity of reef growth. Our results add new  
40 evidence to the idea that reef growth in marginal settings can rapidly turn-on or -off  
41 according to regional environmental changes, and thus are of relevance for predicting  
42 high latitude reef growth potential under climate change.

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44 Keywords: marginal reefs, climate change, subtropical reef, reef accretion, South Atlantic

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## 47 **1. Introduction**

48 Due the additive and interacting forces of global climate changes and local  
49 anthropogenic stressors, coral reefs are currently experiencing a collapse without  
50 precedent in recent millennia (e.g., Aronson et al. 2002; Aronson et al. 2004;  
51 Montaggioni, 2005; Bruno and Valdivia 2016, Côté et al. 2016). However, we also know  
52 that independent of anthropogenic forcing, coral reef health and growth at some sites has  
53 fluctuated through the Holocene (e.g., Perry and Smithers 2011; Toth et al. 2012; Toth et  
54 al. 2018; Dehinck et al. 2019, Gischler and Hudson 2019). Emerging geologic evidence  
55 shows the strong influence of regional climatic and oceanographic changes on the timing  
56 and longevity of reef development. Two major areas of interest have arisen in recent years  
57 in relation to understanding reef growth potential as environmental and ecological  
58 conditions change. The first relates to the question of the timescales over which coral reef  
59 structures have developed. There is, for example, now evidence from a few select sites  
60 that the contemporary coral communities on some reefs are actually ephemeral veneers  
61 growing above largely relict physical structures that ceased significant coral reef-building  
62 several thousand years before present (e.g., Toth et al. 2018). This has led to debate about  
63 the extent to which living coral communities are, or are not, actively contributing to on-  
64 going reef-building, or how long the ecological function of a coral reef may subsist after  
65 its geological senescence (Kuffner and Toth 2016, Toth et al. 2018, Perry and Alvarez-  
66 Filip 2018).

67 The second issue concerns how subtle environmental or oceanographic changes may lead  
68 to reef growth turn-on or -off (terminology *sensu* Buddemeier & Hopley 1988), especially  
69 at sites close to the environmental limits of reef development - i.e., in more marginal  
70 marine settings (Perry & Larcombe 2003). In these areas, corals exist close to their

71 environmental thresholds (Kleypas 1996). Thus, their capacity to build reef structures is  
72 likely to be highly dependent on local to regional environmental changes, turning on when  
73 favourable or turning off when conditions deteriorate. Examples from the turbid-zone  
74 inner-shelf areas of Australia's Great Barrier Reef illustrate how distinct cycles of reef  
75 initiation-growth-demise and then, after a hiatus, new reef development have influenced  
76 regional patterns of reef development (e.g., Perry & Smithers 2011). This, in turn, has  
77 implications for understanding how reef-building may respond in these more marginal  
78 settings under climate change. There has been speculation that the latitudinal range of  
79 reef-building corals may expand polewards under ocean warming (e.g., Vergés et al.  
80 2014), and there is evidence from the Pleistocene that this occurred in Western Australia  
81 (Greenstein and Pandolfi 2006). However, our understanding of the timing and sensitivity  
82 of reef-building to oceanographic changes within more marginal reef building zones of  
83 the global oceans is generally very limited.

84 Here, we integrate both new and published reef core data to explore the timing and nature  
85 of reef-building in the most southerly reef complexes in the Western Atlantic.  
86 Specifically, we report age structure data from the Queimada Grande reef (QGR), a high  
87 latitude (24°S) sub-tropical reef located in the Southwestern Atlantic (SWA) (Pereira-  
88 Filho et al. 2019). We then compare our records of QGR growth with those from the  
89 slightly more northerly Abrolhos (18°S) and Anchieta (21°S) reef complexes.  
90 Collectively, these data provide an opportunity to explore regional phases of reef growth  
91 along this high latitude gradient, and to examine the influence of regional climatic and  
92 oceanographic factors on the timing and longevity of reef building over the past ~6000  
93 years.

## 94 **2. Methods**

## 95 2.1. Study area

96 In the SWA, modern reefs occur from the Amazon river mouth south to 24°S (Moura et  
97 al. 2016, Pereira-Filho et al. 2019) and are built by coralline algae, bryozoans and an  
98 impoverished coral fauna (~23 coral and 5 hydrocoral species) dominated by massive and  
99 encrusting species (e.g. *Mussismilia* spp., *Montastrea cavernosa*, *Siderastrea* spp. and  
100 *Madracis decactis*), several of which are endemic (Leão et al. 2016, Pereira-Filho et al.  
101 2019). While coral reefs occur in North Atlantic subtropical areas (e.g., Florida, Bermuda,  
102 Bahamas; Toth et al. 2018), only one subtropical coral reef, the Queimada Grande Reef  
103 (QGR), is known in the SWA (Figure 1A). Contrasting with its Atlantic tropical  
104 counterparts with their mixed coral, coralline algal and bryozoan frameworks, the QGR  
105 framework was built by only one widely distributed coral species, *Madracis decactis*  
106 (Pereira-Filho et al. 2019). We also note that whilst the currently depauperate living corals  
107 at the site (i.e., only two species *M. decactis* and *Mussismilia hispida*) have a coverage  
108 comparable with that another SWA tropical reefs, in the QGR they do not appear to be  
109 supporting modern reef accretion/accumulation (Pereira-Filho et al. 2019). The QGR  
110 covers ~320,000 m<sup>2</sup> between 10-20 m depths (Figure 1B-C), forming a relatively flat  
111 plateau that fringes the subtidal rock shore (~10-20 m depth) on the leeward (western)  
112 side of the Queimada Grande Island. The island derives from the regressive erosion of  
113 the Serra do Mar scarp which comprises igneous and metamorphic rocks from the Ribeira  
114 mobile belt (granite, granulite, migmatite and gneiss) ~750-450 Ma BP (Almeida et al.  
115 1973). The Queimada Grande Island marine habitats are under a semi-diurnal tidal  
116 influence (~1m of amplitude during spring tidal changes) and are strongly influenced by  
117 the South Atlantic Central Water with sea surface temperatures (SST) ranging from 18 to  
118 28 °C (Bio-ORACLE, <https://www.bio-oracle.org/>). Three morphological palaeo-reef  
119 zones can be distinguished across the reef structure. Although depth and in-water

120 conditions made core recovery challenging, short percussion cores were recovered from  
121 each “zone” to assess internal reef structure and age. These zones are referred to here for  
122 descriptive purposes as: i) proximal reef, ii) mid reef and iii) distal reef. Most coral  
123 colonies were preserved in place (i.e., cylindrical dead coral branches vertically disposed  
124 and filled by trapped sediment) (Figure 1C).

## 125 **2.2. Queimada Grande Reef mapping**

126 Side Scan Sonar surveys included contiguous transects parallel to the island’s rocky  
127 shores between 0 and 40 m depths (Figure 1B). An Edgetech 4100 system with a 272TD  
128 towfish was operated at 100 kHz with 200 and 400 m swaths. Acoustic data were  
129 processed using SonarWis Map4 software; geo-referenced mosaics were exported as  
130 GeoTiff images with 1 m/pixel resolution into ArcGIS 9.2. Morphological attributes such  
131 as area and depth were treated as shapes. The main bottom features identified (i.e., rocky  
132 shore, coral reef and rhodolith bed) were confirmed by ~30h of SCUBA diving  
133 deployments.

## 134 **2.3. Framework description and radiocarbon dating**

135 We obtained three percussion cores with 100% recovery [each ~130 cm - the length of  
136 which was limited by safety constraints for emplacing cores in this way in these water  
137 depths] along a cross-reef transect that included samples from the proximal reef (14 m  
138 water depth), mid reef (13 m) and distal reef (15 m) (Figure 2). Subsamples (n = 15) were  
139 collected in a 30-cm-interval from each core, labeled, and treated with HCl to remove  
140 superficial contamination.

141 Radiocarbon dating was performed on coral samples in excellent taphonomic condition  
142 (i.e., less than ~20% bioerosion and/or infilling; c.f. Toth et al. 2018). These samples were  
143 washed, dried, and reacted under vacuum with 100% phosphoric acid, yielding carbon  
144 dioxide, which was cryogenically purified and reduced to graphite (Vogel et al. 1984).

145 Graphite  $^{14}\text{C}/^{13}\text{C}$  ratios were measured using a CAIS 0.5 MeV accelerator mass  
146 spectrometer (AMS) at the Center for Applied Isotope Studies (University of Georgia).  
147 Measured  $^{14}\text{C}/^{13}\text{C}$  ratios were corrected relative to the international reference material  
148 NBS SRM 4990 (Oxalic Acid I) and corrected for carbon isotope fractionation based on  
149  $\delta^{13}\text{C}$  values (expressed in ‰ relative to the V-PDB international reference) measured on  
150 separate aliquots using an isotope ratio mass spectrometer (IRMS) (analytical precision  
151  $< 0.1\%$ ). Uncalibrated dates are given in radiocarbon years before 1,950 (years AD),  
152 using the  $^{14}\text{C}$  half-life of 5,568 years. The error is quoted as two-sigma standard deviation  
153 and reflects both statistical and instrumental errors. Calibration of radiocarbon date was  
154 performed using the software CALIB Version 8.20 and the calibration curve Marine20  
155 (<http://calib.org>, accessed 2021-01-10). We used the 10 nearest point  $\Delta R -99 \pm 103.0$  as  
156 the best current estimate of variance in the local open water marine reservoir effect. The  
157 median probability age is given as the best estimate of calibrated age. One dating was  
158 also obtained from organic matter from sediment. In this case, the calibration curve  
159 IntCal20 was applied. Sixty-three radiocarbon dating data from Abrolhos and Anchieta  
160 tropical reefs (i.e.,  $18^\circ\text{S}$  and  $21^\circ\text{S}$ , respectively) were compiled from Bastos et al. (2018)  
161 and Dechnik et al. (2019).

#### 162 **2.4. Optically stimulated luminescence dating**

163 Luminescence dating was carried out at the Luminescence and Gamma  
164 Spectrometry Laboratory (LEGaL) of the Institute of Geosciences of the University of  
165 São Paulo. The reef cores were opened under subdued red/amber-light conditions to avoid  
166 bleaching of natural luminescence signals, and at least 40g of sediments were picked from  
167 between the coral framework with a small spoon ( $n = 5$ ). Quartz grains retrieved from  
168 these sediment samples were used for optically stimulated luminescence (OSL). Samples  
169 were collected at the same depth interval as the  $^{14}\text{C}$  dated coral samples and quartz

170 concentrates were prepared following standard procedures as described in Aitken (1998).  
171 The 180–250  $\mu\text{m}$  grain size fraction was obtained by wet sieving, followed by chemical  
172 treatment with 27%  $\text{H}_2\text{O}_2$  and 10%  $\text{HCl}$  to remove organic matter and carbonates,  
173 respectively. Subsequently, samples were treated with 48–51%  $\text{HF}$  for 40 min to remove  
174 feldspar and outer rinds of quartz grains dosed by alpha particles. Due to the small amount  
175 of sample after the chemical treatment, we decided not to separate quartz from heavy  
176 minerals (the absence of feldspar was tested with infrared light), since the signal of heavy  
177 minerals are generally one order of magnitude lower than that of quartz (Krbetschek et  
178 al. 1997, Del Rio et al. 2019).

179 The OSL dating method using the single-aliquot regenerative (SAR) dose protocol  
180 (Murray and Wintle 2000, Wintle and Murray 2006) was carried out on quartz aliquots.  
181 Luminescence measurements were performed in a Lexsyg Smart TL/OSL reader  
182 equipped with blue and infrared LEDs, filters for light detection in the UV band and beta  
183 radiation source ( $^{90}\text{Sr}/^{90}\text{Y}$ ) with dose rate of  $0.116 \text{ Gy s}^{-1}$  for steel discs. Dose recovery  
184 tests were performed on aliquots bleached for 3 h under a solar simulator lamp. Dose  
185 recovery tests were carried out with a pre-heat temperature of  $220 \text{ }^\circ\text{C}$  and given a dose of  
186 5 Gy. The calculated-to-give dose ratio obtained for the dose recovery test was 0.91. The  
187 equivalent dose for each sample was determined by measuring at least 24 quartz aliquots.  
188 Exponential fitting of dose-response data was carried out only when the recycling ratio  
189 was in the 0.9–1.1 range, and recuperation was  $< 5\%$  and feldspar contamination was  
190 negligible. Feldspar contamination in each aliquot was appraised by repeating the first  
191 regeneration dose and using infrared stimulation at  $60 \text{ }^\circ\text{C}$  before blue stimulation.  
192 Aliquots were rejected when an infrared stimulated luminescence signal was detected, or  
193 when the OSL signal was depleted following infrared stimulation. The equivalent dose



194 was determined through the central age model (Galbraith et al. 1999) using at least 13  
195 accepted quartz aliquots per sample.

196 To calculate dose rates, mean specific activities ( $\text{Bq kg}^{-1}$ ) of  $^{238}\text{U}$ ,  $^{232}\text{Th}$ , and  $^{40}\text{K}$   
197 were assessed through gamma ray spectrometry using a high purity germanium (HPGe)  
198 detector with energy resolution of 2.1 KeV and relative efficiency of 55%, encased in an  
199 ultra-low background shield. Samples were kept in sealed plastic containers for at least  
200 28 days for radon equilibration before gamma ray spectrometry. The conversion factors  
201 provided by Guérin et al. (2011) were used for beta and gamma dose rates' calculation.  
202 The cosmic dose rate was evaluated using sample depth, altitude, longitude and latitude  
203 (Prescott and Stephan 1982). Moisture content was considered as the maximum water  
204 saturation (water weight/dry sample weight) for each sample, by saturating the samples  
205 used in dose rate calculation. The OSL ages were calculated by dividing the equivalent  
206 dose value by the dose rate value (Table 1).

## 207 **2.5. Sediment analysis**

208 In the three reef cores, sediment samples ( $n=72$ ) were collected manually in a 4  
209 cm vertical spacing, using a small spoon. Only sediment trapped in the reef framework  
210 among coral colonies (10-15 ml) was sampled. All samples were wet sieved in a 0.5 mm  
211 mesh to remove larger coral fragments resulting from core processing. After sieving,  
212 samples were oven-dried, weighted and treated with HCl and  $\text{H}_2\text{O}_2$  to remove carbonates  
213 ( $\text{CaCO}_3$ ) and organic matter (OM), respectively. Samples were washed with distilled  
214 water, dried and weighed after chemical treatment in order to calculate carbonate and OM  
215 contents. Grain-size analyses were performed for all samples, after chemical treatment,  
216 using a Malvern Mastersizer 2000 laser granulometer coupled with a Hydro 2000MU  
217 dispersion unit (distilled water used as dispersion medium) and a built-in ultrasound  
218 device. Descriptive statistics parameters were calculated according Pearson's moments

219 method. Size fractions are given in 0.125 phi interval (Wentworth, 1922). Since silt and  
220 fine sand were the dominant size fractions in all samples, the ratio (silt + fine sand)/silt  
221 was calculated in order to evaluate the variation of these fractions without auto  
222 correlation.

### 223 **3. Results**

224 Side scan sonar mapping revealed that the major bottom features of the QGR can be  
225 delineated into three palaeo reef zones on the basis of their morphology, relative depth  
226 and position relative to the shoreline: an distal reef zone (distal from the rock shore) with  
227 an inclination of  $\sim 30^\circ$  at the deepest portion (between depths of 15 to 25 m relative to  
228 present MSL); an proximal reef zone, located close to the shore at a depth of  $\sim 14$  m; and  
229 between these an mid reef zone, at a depth of  $\sim 13$  m (Figure 2). Radiocarbon dating of  
230 coral clasts in cores also clearly demonstrates that the present reef surface is essentially a  
231 relict feature, and that whilst the cores did not capture the reef initiation phase, it is clear  
232 that the reef was actively accreting prior to 6,000 cal. yr. B.P.. Analysis of cores from  
233 each of these palaeo reef zones indicates: i) that the QGR is constructed entirely of one  
234 cylindrical branched coral species *Madracis decactis*; and ii) that the accumulating reef  
235 framework comprises intermixed units of in-place and detrital coral rubble, with non-  
236 reefal units between (defined by the lack of any coral material) (Figure 3).

237 Cores from the interpreted proximal and mid reef zones identify two apparently distinct  
238 phases of reef growth since  $\sim 6000$  cal. yr. B.P.. These occurred between 5,749-5,677 cal.  
239 yr. B.P. and between 2,380-2,586 cal. yr. B.P., with a non-depositional hiatus in between  
240 (median probability ages are presented as the best estimate of  $^{14}\text{C}$  calibrated ages, the  
241 complete data are available in Table 2 and Figure 3). In the proximal reef core, dates  
242 ranged between 5,749 (base of core) to 2,533 cal. yr. B.P. (top of core), but with no  
243 evidence of framework accumulation between  $\sim 5,677$  and 2,501 cal. yr. B.P.. This period

244 of time is represented by a time condensed sediment interval ~15 cm thick devoid of coral  
245 material (Figure 3). Sediments recovered along the core (i.e., among coral branches and  
246 deposited during the reef hiatus interval) are composed mostly of carbonates whose  
247 abundance ranged between 55-75%, organic matter content varied from 1 to 5%, with one  
248 peak coinciding with lower carbonate contents at ~60 cm (Figure 4). While organic matter  
249 recovered from 56 cm returned a  $^{14}\text{C}$  age of 4,730 cal. yr. B.P. ( with a  $\delta^{13}\text{C}$  value of -  
250 24.14 ‰ suggesting a continental origin from C3 plants), quartz OSL ages from the same  
251 portion indicate that sedimentation occurred around  $3,602 \pm 388$  years ago [i.e., optically  
252 stimulated luminescence (OSL) age analysis, see Figure 4 and Tables 1 and 2]. This  
253 ~1,000 years lag between terrestrial carbon fixation and sediment deposition suggests  
254 distant sources of fine sediment portions deposited ~3,600 years ago.

255 In the “mid reef” core, dates ranged between 5,556 (base of core) and 2,489 cal. yr. B.P.  
256 (at core top) (Table 2). No in place coral occurred in the core between a depth of ~40-80  
257 cm, and this depositional interval also indicates a period lacking reef framework  
258 accumulation between ~5,500 and ~2,500 cal. yr. B.P. (Figure 3). The only  $^{14}\text{C}$  datable  
259 carbonate in this non-reefal unit were serpulids tubes (i.e., sessile polychaete) which  
260 returned an age of ~3,300 cal. yr. B.P.. Sediments were composed mostly of carbonates  
261 that ranged between 50-80%. Organic matter content varied from 1 to 8%, higher  
262 percentages coinciding with lower carbonate contents at 40 cm (Figure 4). Sediments  
263 from the non-reefal unit, where serpulids were found, returned a broadly comparable OSL  
264 age of  $2,931 \pm 229$  years ago (Figures 3 and 4 and Table 1) and suggests higher rates of  
265 sedimentation during periods unfavorable to the coral growth.

266 The distal reef core was collected at 15 m depth. It similarly contained in place coral  
267 skeletons all of which dated to the post-hiatus period identified in the mid and proximal  
268 reef cores above (dates ranged from 2,366 and 2,478 cal. yr. B.P.). Although some age

269 inversions could be suggested by calibrated ages, they are not significant considering the  
270 error bars of radiocarbon dating and may be indicative of a mix of both in place but also  
271 some transported (from shallower zone) coral clasts. In this reef zone (i.e., the distal reef),  
272 where 1 m of the core comprises approximately 100 years of deposition, three distinct  
273 sedimentary horizons are discernible characterised by an increase in sand and a decrease  
274 in silt contents, and may indicate a period of depositional interruption, or again event  
275 horizons (Figure 3 and 4). Sediments seem to be have trapped soon after reef framework  
276 accumulation based on difference between coral growth (i.e., C14) and sedimentation  
277 (i.e., OSL) (Tables 1 and 2). Carbonate content is in the range of 45-60%, while organic  
278 matter content varied from 2.5 to 15%, with higher concentration coinciding with lower  
279 carbonate contents (Figure 4).

#### 280 **4. Discussion**

281 Core records from the sub-tropical QGR show clear evidence for a Mid-Holocene  
282 cessation in framework accumulation at this site. Although we could not determine when  
283 reef growth initiated at these sites, our data indicate that the reef was clearly accreting  
284 prior to 6,000 cal. yr. B.P. when the regional sea level was in a late stages of rise before  
285 reaching a highstand at ~5,500 cal. yr. B.P. (~4 m above the present) (Angulo et al., 1999,  
286 2006, 2013, Toniolo et al. 2020) (Figure 5). Nevertheless, the current depth of the palaeo-  
287 reef surface and coral ages from its surface indicate that the QGR never reached sea level  
288 (Figure 5). Thus, regional sea-level rise (SLR) was not the major driver of vertical growth  
289 in terms of providing space for reef accommodation such as has been reported for reefs  
290 northward from QGR (i.e., Anchieta and Abrolhos reefs; 20°S and 18°S, respectively)  
291 (Dechnik et al. 2019, Vasconcelos et al. 2019) (Figure 5) and many other tropical reefs.  
292 Furthermore, the relatively limited amount of RSL rise that occurred over the period of  
293 interest here argues against sudden jumps in sea level leading to either reef drowning or

294 back-stepping (sensu Blanchon et al. 2002). Instead, our data add evidence to support the  
295 idea that reef-building in marginal settings is strongly influenced by regional climatic and  
296 oceanographic changes. These changes impact both the timing and longevity of reef  
297 development, with reefs turning on or turning off when conditions are favorable or  
298 unfavorable. In the QGR records our data suggest four episodes of reef development and  
299 demise: i) an initial reef accretion phase (RAP) up to ~5,500 cal. yr. B.P.; ii) a hiatus  
300 phase between ~5,500 and ~2,500 cal. yr. B.P.; iii) a second phase of reef framework  
301 accumulation at ~2600 to ~2400 cal. yr. B.P.; and, iv) the post ~2,300 y B.P. period during  
302 which no further reef accretion occurred. Interestingly, the Mid to Late Holocene reef  
303 growth phases described here have also been reported for other regions such as the  
304 Southern Pacific (Woodroffe et al. 2010, Perry and Smithers 2011), the Northwestern  
305 Pacific (Hamanaka et al. 2012), the western coast of Panama (Toth et al. 2012), and the  
306 tropical southwestern Atlantic (Dechnik et al. 2019).

307 SLR has previously been evoked as having influence on Mid to Late Holocene  
308 reef growth in SWA tropical areas (e.g., at Anchieta Reef and Abrolhos Reef, 20°S and  
309 18°S, respectively). However, this period was also accompanied by other climatic and  
310 oceanographic changes that may explain changes in reef accretion phases in high latitude  
311 settings such as the Queimada Grande subtropical reef. For instance, RAP I may relate to  
312 a southward displacement of the Inter-Tropical Convergence Zone (ITCZ), coupled with  
313 Southern Westerly Wind (SWW) weakening (Mariani et al. 2017), changes which have  
314 been attributed to the movement of the South Atlantic Subtropical Dipole (SASD)  
315 (Wainer et al. 2014). Specifically, shifts in the SWW are known to have altered the  
316 position of the Brazil-Falklands/Malvinas Confluence (BFMC) during the Holocene  
317 (Voigt et al. 2015). The BFMC is one of the world's most energetic oceanographic  
318 features and a major driver of freshwater, sediment and nutrient dispersion from the La

319 Plata River (Peterson and Stramma 1991, Brandini et al. 2000, Gu et al 2019). Currently,  
320 the south-flowing Brazil Current (BC) converges with the north-flowing  
321 Falklands/Malvinas Current (FMC) at  $\sim 38^{\circ}\text{S}$ , and the La Plata discharge influence  
322 reaches  $\sim 25^{\circ}\text{S}$  (Gyllencreutz et al. 2010) (Figure 6A). From 8,700 to 5,500 cal. yr. B.P.,  
323 as a consequence of SWW weakening, the BFMC shifted from its Mid Holocene  
324 northernmost position (Figure 6B) southward, resulting in warmer SSTs at  $24^{\circ}\text{S}$  (Voigt  
325 et al. 2015, Gu et al. 2019). These relatively warm conditions, combined with lower El  
326 Niño-Southern Oscillation (ENSO) frequencies, prevailed until  $\sim 5,500$  cal. yr. B.P.  
327 (Hodell and Kanfoush 2001, Voigt et al. 2015, Gu et al. 2019), conditions that we suggest  
328 favoured reef development at QGR and probably also influenced other Southwestern  
329 Atlantic tropical reefs (Dechnik et al. 2019) (Figure 6C). Wider evidence for this warmer  
330 South Atlantic SST period occur in cores from  $53^{\circ}\text{S}$ , which exhibit foraminifera  
331 abundance peaks up to 6,500 cal yr. B.P. (Figure 6D). This warm period was also marked  
332 by latitudinal expansion of Pacific South America lowland thermophilous trees (Moreno  
333 2004).

334 The QGR then “turned off” from  $\sim 5,500$  through to  $\sim 2,500$  cal. yr. B.P., a change  
335 we interpret as a function of deteriorating (for coral growth) oceanographic conditions in  
336 this marginal marine setting. Evidence from elsewhere in the Southern Ocean would  
337 suggest that this is likely to have been driven by significant SST cooling and ice expansion  
338 further south (Figure 6E). Specifically, during this period, there is evidence for an  
339 apparent decoupling of the ITCZ and SWW, as a consequence of AMOC stability  
340 (Mariani et al. 2017). This resulted in the northward migration of Antarctic winter sea ice  
341 as a consequence of weakened polar maritime air masses (Hodell and Kanfoush 2001),  
342 and an increasing importance of ENSO (El Niño and La Niña) in modulating climate  
343 (Toth et al. 2012, Mariani et al. 2017). Terrestrial records from this time also show

344 vegetation being replaced by cold-resistant rain forest trees (Moreno 2004). Negative  
345 SASD records (Wainer et al. 2017), absence of foraminifera up to ~1,830 cal. yr. B.P. in  
346 cores from 53°S (Hodell and Kanfoush 2001) (Figure 6D-F) and Sr/Ca records on  
347 vermitid shells from 23°S to 26°S (Toniolo et al. 2020) corroborate this interpretation.  
348 Interestingly, a similar period of reef growth hiatus in the Northwestern Pacific (i.e.,  
349 ~29°N) is also explained by cold SSTs as consequences of changes in oceanographic  
350 regional patterns (i.e., a Kuroshio Current weakening) (Hamanaka et al. 2012). Sediment  
351 coarsening (higher contributions of sand) coincides with peaks in the carbonate fraction  
352 in the QGR, suggesting an increase in wave/current energy during the hiatus (Figure. 3).  
353 While increases in the carbonate fraction seem related to reef erosion, terrigenous sand  
354 increases are likely associated with remobilization of sediments from the inner shelf by  
355 offshore transport. Indeed, the reef growth hiatus at QGR is concurrent with the lowest  
356 rates of sediment retention by a coastal sand barrier located in its nearest estuarine system  
357 (Guedes et al. 2011). Additionally, differences between sediment depositional ages (OSL  
358 dated at 3,602 years ago) and organic matter radiocarbon ages (4,741 cal. yr. B.P.) during  
359 the reef growth hiatus period suggest distant sources for the fine organic sediments.  
360 Indeed, values of  $\delta^{13}\text{C}$  (i.e., -24 to -23‰, see Table 2) indicates C3 continental plants  
361 (Myer 2003) that would be consistent with influence of the South America drainage  
362 system during the Mid to Late-Holocene, including Plata River plume.

363         The second short phase of reef growth at QGR (RAP II) occurred from ~2,500 to  
364 2,300 cal. yr. B.P.. Relative sea level fell ~4 m around this time towards present levels  
365 (Figure 4), probably elevating light penetration on the submergent reef surface. However,  
366 waning La Niña and reduced El Niño strength linked with weakened SWW, as well as a  
367 positive SSAD, were likely key drivers of this period of renewed reef growth, which is  
368 also evident in Abrolhos reef (18°S) (Figure 6C and F-G). Reduced terrigenous sediment

369 content in cores also suggests a reduction in wave/current energy at QGR around ~2,500  
370 years B.P. (Figure 4). However, the synergistic effects of these conditions, which were  
371 favorable to coral growth, lasted a very short time (~200-300 years) and were followed  
372 by a period with the highest ENSO variability recorded during the Holocene (i.e., since  
373 ~2,000 cal. yr. B.P.) (Figure 6G). While SSE winds during La Niña would have increased  
374 the northward penetration of the La Plata plume and colder conditions, El Niño is linked  
375 to higher precipitation around 24°S. This shift in ENSO frequency corresponds to a  
376 coarsening of the sand fraction in the cores (Figure 4). Late Holocene reef turn-on has  
377 also been reported in other areas that are directly affected by ENSO (i.e., the southwestern  
378 and northeastern Pacific), although at these locations reef accretion has generally  
379 persisted until present.

380         Since ~2,000 cal. yr. B.P., the QGR has thus existed in an essentially senescent  
381 state as a submerged structure colonized by non-reef building organisms. These include  
382 fleshy and turf algae, the soft coral *Palythoa caribaeorum*, sponges (e.g., *Aplysina fulva*,  
383 *A. caissara*, and *Scopalina ruetzleri*), tunicates, and colonies of the hermatypic corals  
384 *Mussismilia hispida*, *Madracis decactis* and crustose coralline algae (Pereira-Filho et al.  
385 2019), although the latter are not sufficiently abundant to add framework to the structure.  
386 *M. hispida*, which is the most common coral of these contemporary reefs (i.e., ~15% of  
387 benthic coverage) was not recovered in our cores. This may reflect later colonization by  
388 *M. hispida*, which is at its very southern limits of occurrence (also corroborated by recent  
389 molecular data, Peluso et al. 2018). In contrast, the only coral recovered in cores - i.e.,  
390 *Madracis decactis* - is widely distributed throughout the Atlantic Ocean (from ~32°N to  
391 ~27°S down to 30 m depth). It is thus readily acclimated to suboptimal conditions and  
392 capable of exploiting favourable environmental shifts at its limits of occurrence (i.e.,  
393 subtropical regions) to sustain localised reef development.



394            Similar reef-building phase-shifts also occurred in the SWA's largest and most  
395 speciose reefs (i.e., Abrolhos Reefs, 17–19.5° S). At this more northerly location (~1,000  
396 km north of QGR) corals were also the major framework builders up to ~2,000 cal. yr.  
397 B.P. (Bastos et al. 2018). However, in contrast to QGR, reef growth has continued at  
398 Abrolhos over the last ~2,000 years, although it has not been driven by corals but by  
399 bryozoans. The absence of on-going reef accretion at QGR thus probably reflects its  
400 reliance on sufficient densities of the sole Holocene reef-builder *M. decactis* to drive  
401 framework accumulation. We also noted that the Holocene sequences reported here are  
402 thinner than those reported in other South Atlantic tropical reefs (e.g., Vasconcelos et al.  
403 2018, Bastos et al. 2018, Dechnick et al. 2019), which would be expected given the very  
404 southerly location of the QGR (i.e., 24°S). The lack of contemporary reef-building at this  
405 site, combined with the evidence of past reef growth phases, illustrates how even subtle  
406 shifts in climatic and oceanographic conditions in such marginal settings can rapidly turn-  
407 on or -off reef growth, an observation of relevance for predicting high latitude reef growth  
408 potential under climate change.

409

## 410 **5. Conclusion**

411 Our data show that SWA reefs growth has experienced distinct phases of turn-on and -  
412 off throughout the Mid- to Late-Holocene period. Beyond the well know influence of  
413 SLR on reef growth, our data suggest a strong influence of regional climate changes in  
414 driving SWA reef development over the last ~6,000 yrs B.P.. Thus, this study contributes  
415 to a broader understanding of marginal reefs by adding new evidence to the idea that reef  
416 growth potential in these marine settings can rapidly change according to environmental  
417 changes. Clearly, the logistical issues that prevented longer and more extensive core  
418 recovery necessitate some caveats on our interpretation, however the consistency with the

419 timing of reef growth transitions in slightly more northerly reefs in this region, provide  
420 support to the patterns we identify. More widely it is important to emphasise that  
421 knowledge on high latitude reef development generally, but of South Atlantic reef  
422 development specifically, remains limited and important questions are still unanswered.  
423 For instance, there has been recent debate regarding the resilience of the SWA reef-  
424 building corals, with some evidence indicating high susceptibility to global warming  
425 (Duarte et al. 2020), but other evidence suggesting that corals in SWA marginal marine  
426 settings may be highly resilient when facing this modern challenge (Mies et al. 2020).  
427 Here, we add geological pieces to this intricate puzzle by showing that SWA reefs  
428 alternated between reef development and demise throughout the last ~6,000 years.

429

#### 430 Declaration of Competing Interest

431 The authors declare that they have no known competing financial interests or personal  
432 relationships that could have appeared to influence the work reported in this paper.

433

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445 Table 1: Optically stimulated luminescence age analysis. PR – Proximal Reef, MR –  
 446 Mid Reef, and DR – Distal Reef  
 447

Sample (Core depth)	Accepted aliquots	Equivalent Dose (Gy)	Overdispersion (%)	K (%)	<sup>238</sup> U (ppm)	<sup>232</sup> Th (ppm)	Cosmic dose rate (Gy.ky-1)	Total dose rate	Age (years)
PR(60cm)	19/25	1.78	24.4	0.31	0.618	3.463	0.0744	0.494	3,602±388
MR(60cm)	13/15	1.54	18	0.40	0.594	2.497	0.0744	0.525	2,931±229
DR (30cm)	3/5	1.28	27.2	0.50	0.54	2.230	0.0744	0.518	2,472±514
DR (60cm)	20/26	1.59	9	0.72	1.043	2.526	0.0744	0.726	2,189±185
DR (90cm)	11/12	1.54	13.5	0.50	0.61	1.970	0.0744	0.576	2,674±229

448

449

450 **Table 2:** Radiocarbon ages and  $\delta^{13}\text{C}$  and  $\delta^{18}\text{O}$  values and Optically Stimulated  
451 Luminescence dating parameters. Calibration of radiocarbon datings was performed  
452 using the software CALIB Version 8.20 and the Marine20 calibration curve  
453 (<http://calib.org>). We used the ten nearest points  $\Delta R -99 \pm 103.0$  as the best estimate of  
454 the local open water marine reservoir effect and referring to 2 sigma calibrated range. The  
455 median probability age is given as the best estimate of calibrated age. \* Organic matter  
456 from sediment (in this case the IntCal20 calibration curve was used), \*\* non-coral sample  
457 (i.e., Serpulid tubes)  
458

RADIOCARBON AGES								
Core	Lab Code	Material	Core Depth (cm)	$\delta^{13}\text{C}\text{‰}$	$\delta^{18}\text{O}\text{‰}$	$^{14}\text{C}$ years BP	Years Cal BP	Median Probability
Proximal Reef	UGAMS35182	Coral	Top	-1.23	-2.15	2,840 ( $\pm 20$ )	2,247 – 2,805	2,533
	UGAMS35183	Coral	30	-1.40	-1.75	2,810 ( $\pm 20$ )	2,198 – 2,759	2,501
	UGAMS35184	Coral	60	-1.27	-1.49	5,390 ( $\pm 25$ )	5,412 – 5,940	5,677
	UGAMS 39087*	Sediment	60	-24.14	-	4,210 ( $\pm 25$ )	4,576 – 4,859	4,730*
	UGAMS 35185	Coral	90	-1.45	-2.10	5,430 ( $\pm 25$ )	5,458 – 5,982	5,719
	UGAMS 35186	Coral	120	-1.22	-1.74	5,460 ( $\pm 20$ )	5,475 – 6,004	5,749
Mid Reef	UGAMS 35177	Coral	Top	-1.12	-1.60	2,800 ( $\pm 20$ )	2,188 – 2,749	2,489
	UGAMS35178	Coral	30	-1.34	-1.96	2,890 ( $\pm 20$ )	2,307 – 2,857	2,586
	UGAMS35179**	Serpullid	60	2,09	0.15	3,530 ( $\pm 20$ )	3,069 – 3,666	3,367**
	UGAMS35180	Coral	90	-1.51	-1.56	5,220 ( $\pm 20$ )	5,223 – 5,765	5,491
	UGAMS 35181	Coral	100	-1.18	-1.67	5,280 ( $\pm 20$ )	5,301 – 5,837	5,556
Distal Reef	UGAMS35187	Coral	Top	-2.26	-1.30	2,710 ( $\pm 20$ )	2,089 – 2,691	2,380
	UGAMS35188	Coral	30	-1.19	-1.85	2,700 ( $\pm 20$ )	2,074 – 2,683	2,366
	UGAMS35189	Coral	60	-1.79	-1.40	2,750 ( $\pm 20$ )	2,137 – 2,715	2,431
	UGAMS35190	Coral	90	-1.00	-1.27	2,790 ( $\pm 20$ )	2,178 – 2,742	2,478
	UGAMS35191	Coral	100	-1.73	-1.76	2,780 ( $\pm 20$ )	2,167 – 2,734	2,467

459

460

461 **Figure Legends**

462

463 **Figure 1** – A) South Atlantic coral reefs for which Holocene reef growth is relatively  
464 well know. B) Sonogram of the Queimada Grande Reef (QGR). Hard substrates (i.e.,  
465 rocky shore and coral reef) are indicated by higher reflectance sign while rhodolith beds  
466 (intermediate bottom between hard and soft bottom) are indicated by lower reflectance.  
467 White rectangle indicates the coring location, while dashed white line shows the limit of  
468 the QGR, with its characteristic reflectance pattern. C) Image illustrating the QGR reef  
469 framework build major by *Madracis decactis* encrusted by coralline algae at the top.  
470 Space between vertical coral branches is filled with sediments.

471 **Figure 2** – A) Detailed sonogram where percussion cores were obtained (asterisks).  
472 White rectangle corresponds to the same area indicated in Figure 1B. Rocky shore and  
473 rodolith bed were interpreted according their reflectance sign and by ground truth  
474 SCUBA diving. B) Schematic cross-section of core locations.

475 **Figure 3** - Stratigraphic position and calibrated AMS radiocarbon ages [years Before  
476 Present (B.P.)] of the coral samples (excepted at 60 cm mid reef where serpulid worm  
477 tube was dated) and OSL ages of sediment (years ago). Vertical axis indicates the core  
478 depth.

479 **Figure 4** –Vertical profiles of CaCO<sub>3</sub>, organic matter content (%) and grain size  
480 parameters. Horizontal dashed lines indicate sediment OSL age (years ago) or  
481 radiocarbon age from organic matter content (cal years B.P.). A) Proximal Reef, B) Mid  
482 Reef and C) Distal Reef.

483 **Figure 5** – Holocene Relative Sea Level curve for the Southwestern Atlantic after Angulo  
484 et al., (1999, 2006, 2013) and Toniolo et al. 2020. Reef growth phases are represented

485 according to calibrated radiocarbon datings of corals from Anchieta (green), Abrolhos  
486 (brown) and Queimada Grande Reef (blue) (18°S, 21°S and 24°S, respectively). Error  
487 bars correspond to two-sigma standard deviation from the mean of the probability  
488 distribution of radiocarbon ages, and reflects both statistical and instrumental  
489 uncertainties.

490 **Figure 6** – A) Schematic position of the Brazilian Falkland/Malvinas Confluence  
491 (BFMC) and the major oceanographic features; i.e., South Atlantic Current (SAC),  
492 Falkland/Malvinas Current (FMC), Brazilian Current (BC) and Brazilian countercurrent  
493 (BCC). Sea surface temperature data correspond to February 1998 (obtained from NOAA  
494 products [https://www.ospo.noaa.gov/data/sst/mean\\_mon/February.98.monmean.gif](https://www.ospo.noaa.gov/data/sst/mean_mon/February.98.monmean.gif) website,  
495 [https://www.ospo.noaa.gov/data/sst/mean\\_mon/February.98.monmean.gif](https://www.ospo.noaa.gov/data/sst/mean_mon/February.98.monmean.gif)). B) BFMC  
496 northernmost position between 8,700 and 5,500 BP, based on Gu et al. (2019).  
497 Temperature information is merely schematic and was produced hypothesizing the  
498 northward displacement of the current fronts presented in A in order to illustrate our  
499 hypothesis. C) Rapid accretion phases of the Southwestern Atlantic reefs (i.e., Abrolhos,  
500 Anchieta and Queimada Grande Reefs) (i.e., calibrated median probability and two-sigma  
501 standard deviation), shaded vertical grey bars indicate reef accretion phases and hiatus  
502 (dark and light, respectively) D) Foraminifera abundance (%) (53°S) from Hodell and  
503 Kanfoush (2001). E) February sea surface temperature estimated from diatom  
504 assemblages from Hodell (2001). F) South Atlantic Subtropical Dipole index during the  
505 Holocene from Wainer et al. (2014). G) El Niño Southern Oscillation (ENSO)  
506 reconstructions from El Junco Lake (Conroy et al., 2008).

507

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