

Coral Reef Ocean Acidification Monitoring: Development of a US Monitoring Strategy

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ABSTRACT

30 The *Coral Reef Ocean Acidification Monitoring Portfolio (CROAMP) Workshop* was hosted by
31 the NOAA Ocean Acidification Program and the National Coral Reef Institute at the Nova
32 Southeastern University Oceanographic Center in August, 2012. Researchers and project
33 managers from around the world engaged in coral reef ocean acidification monitoring sought to
34 define a suite of metrics to optimally include as part of long-term coral reef monitoring. These
35 metrics comprise a listing of high-level strategic guiding questions towards which an observing
36 network should ideally be configured. These metrics are proposed as best able to discern
37 specific attribution of changes in coral reef ecosystems in response to ocean acidification. The
38 metrics necessarily constrain the observing requirements comprised of a suite of biogeochemical,
39 ecological, hydrologic, and meteorological measurements. We also clarify expectations for what
40 sustained ocean acidification monitoring can and cannot offer. Achieving the full CROAMP
41 requirements identified here will demand extensive leverage of existing national and
42 international initiatives particularly in an environment of increasing fiscal constraint.

43 **I. INTRODUCTION**

44 Ocean acidification poses many challenges to the long-term sustainability of coral reefs primarily
45 (although not exclusively) due to its capacity to reduce the rates of both biocalcification and
46 inorganic precipitation of carbonate minerals. Both individual organism and net community
47 calcification rates have generally been experimentally observed to decrease under expected
48 future OA conditions [e.g., Gattuso et al., 1998; Marubini et al., 2001, 2002; Marshall and Clode,
49 2002; Ohde and Hossain, 2004; Borowitzka, 1981; Gao et al., 1993; Langdon et al., 2000, 2003;
50 Langdon and Atkinson, 2005; Leclercq et al., 2000, 2002; Anthony et al., 2008]. Even under
51 contemporary conditions, field studies have revealed periods of net community dissolution [cite]
52 with the expectation that these periods could increase under continued ocean acidification. Any

53 decline in net community calcification by coral reef organisms would likely compromise the
54 long-term persistence of many coral reef ecosystems because even the growth of healthy,
55 undisturbed coral reefs are known to only slightly outpace rates of loss due to physical and
56 biological erosion [see Glynn, 1997 for review].

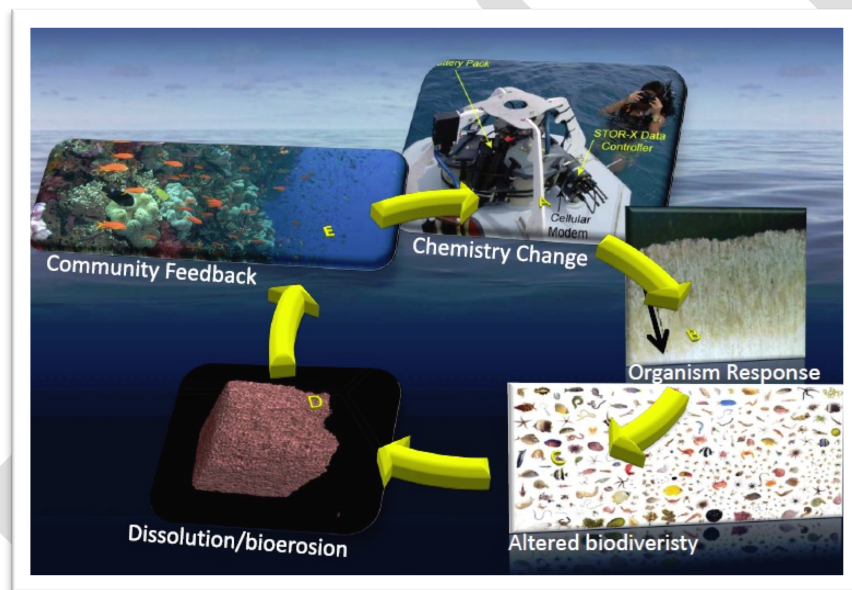
57
58 As US coral reef monitoring initiatives increasingly consider incorporation of ocean acidification
59 aspects into their designs, it's important that these investments be strategically directed towards
60 best informing our understanding of ecological consequences and feedbacks. Frequently,
61 monitoring studies offer only limited capacity to discern specific agents of change. Indeed, a
62 central challenge of field research is interpreting results amidst the sheer complexity of the
63 natural environment where change reflects the net result of multiple factors. Still, there remains
64 a desire to engineer diagnostic monitoring most capable of testing response predictions of coral
65 reef ecosystems to ocean acidification as one means by which to identify potential refugia or
66 "hot spots" and while no metric has been identified which is exclusively sensitive to OA alone,
67 there is consensus that some are better than others. This report details those metrics and outlines
68 the measurement requirements necessary to derive them.

69

70 **II. CROAMP WORKSHOP**

71 A working group of nearly fifty cross-disciplinary multinational federal and academic scientists
72 was hosted by the NOAA Ocean Acidification Program and the National Coral Reef Institute at
73 the Nova Southeastern University Oceanographic Center, Center of Excellence for Coral Reef
74 Ecosystem Science, Dania Beach, FL on August 28-29, 2012. The workshop comprised 11

75 panel discussions detailing the current state-of-science and offering proposed monitoring metrics
76 centered around five thematic areas: direct chemical changes, organism response, biodiversity
77 impacts, dissolution/bioerosion, and community-scale feedbacks. In addition, seven capacity
78 presentations were offered reviewing current and developing monitoring research efforts around
79 the world across multiple agency and academic efforts. A series of breakout sessions worked to
80 develop a portfolio of proposed metrics to be recommended for coral reef ecosystem monitoring
81 which would offer valuable insights into how coral reef ecosystem may be changing in response
82 to ocean acidification.



83
84 **Figure1. Ocean acidification is anticipated to impact coral reefs in several ways. CROAMP**
85 **discussed monitoring for changes across five overarching thematic areas: A. detecting**
86 **direct chemical changes with the coral reef waters attributable to OA, B. detecting specific**
87 **organism responses to OA, C. tracking changes in biodiversity most susceptible to OA, D.**
88 **quantifying changes in dissolution and bioerosion rates, E. discerning changes and**
89 **feedbacks to whole-community metabolism.**

90

91 The changing status of these metrics over time should aid in assigning specific attribution to OA,
92 though it was recognized that there will likely be multiple factors driving observed changes we
93 must be vigilant of co-variants. These workshop outcomes are intended to inform national and
94 international long-term OA monitoring efforts within coral reef ecosystems.

95

96 **III. The CROAMP Framework**

97 The CROAMP Framework was developed around the guiding principle that the observing
98 requirements should be derived from high-level overarching questions addressing how key
99 aspects of coral reef ecosystems might be anticipated to change over-time based upon the current
100 state-of-the-science. Within each thematic area, the following was developed:

101 *Strategic Monitoring Goal (SMG)* – These represent the high-level overarching goals towards
102 which an observing portfolio is intended to answer. It indicates “*what*” we seek to know about
103 the system with regards to its ultimate response to ocean acidification. Each strategic monitoring
104 goal includes units of measure or is explicitly reported as unitless as applicable.

105 *Tactical Monitoring Metric (TMM)* – A TMM represents one or more specific tasks necessary to
106 achieve each of the strategic monitoring goals. These detail “*how*” we will acquire the requisite
107 information including the methods, associated measured parameters, and optimal frequency of
108 measure. As is the case with the Strategic Monitoring Goal, each Tactical Monitoring Metric
109 also includes units of measure. While various methods may be available to meet the tactical
110 requirements, the intent of this document is not to provide an exhaustive detail of all methods

111 that could be employed, but simply to offer valuable examples. Other important considerations
112 in developing each of the TMM's included the feasibility of adoption into proposed monitoring
113 initiatives and what site criteria would be most amendable towards deriving the proposed TMM.
114 Finally, a critical aspect which was discussed as part of each TMM is the number and importance
115 of co-variants which represent factors other than OA which will convolute assigning specific
116 attribution. In nearly all cases, multiple co-variants were identified and are reflected in the
117 amount of secondary measurements which would need to be obtained simultaneously.



118
119 **Figure 2. A guiding principle of CROAMP is that observing requirements are determined**
120 **by first carefully formulating tactical metrics that specifically inform strategic**
121 **requirements within each thematic area. Therefore, the primary emphasis of CROAMP**
122 **was upon developing the strategic requirements in a deliberate “top-down” fashion.**

123

124 *1. Direct Chemistry Changes in Response to Ocean Acidification*

125 By far the most certain consequence of ocean acidification are the direct chemistry changes in
126 response to rising atmospheric CO₂ concentration which are reflected in surface waters as
127 reduction in pH, an enrichment in total dissolved inorganic carbon, and a decrease in the
128 availability of carbonate ion. Such changes are already apparent in the extended time-series
129 observations at the Bermuda Atlantic Time-Series (BATS) and Hawaii Ocean Time-Series.
130 While changes in the hydrologic cycle, ocean circulation, and primary productivity may augment
131 these rates regionally over decadal time-scales, generally the seasonally detrended changes
132 observed in the surface waters (upper 50m) at these stations closely approximated near
133 equilibrium with rising atmospheric CO₂. However, there is reason to suspect that such rates
134 may not precisely equate to the rates observed within many coral reef environments due to local
135 feedback and buffering processes. The relative influence local processes will have on the
136 overlay ocean acidification rate will be dependent upon several factors including water mass
137 residence time, benthic community composition, and net community metabolism. These
138 feedbacks will be considered more closely in the next section.

139
140 A first-order question any ocean acidification monitoring observatory needs to be able to answer
141 is if the rates of ocean acidification are consistent with the regional oceanic average, or are they
142 decoupled. If they are decoupled then answering “why” will likely be addressed in the
143 subsequent thematic areas. To evaluate “if” they are decoupled demands that one monitoring not
144 only carbonate chemistry dynamics within the reef environment of interest, but also have careful
145 constraint on the rate of changes of neighboring oceanic waters where the relative influence of
146 coastal and benthic processes is negligible. This allows one to normalize the observed rates of
147 change within the reef environment to the broader regional rates of change.

148

149

What is the annual rate of change in seawater carbonate chemistry within reef environments relative to comparative rates of changes in neighboring open ocean surface waters? ($\Omega \text{ yr}^{-1}$, $\mu\text{atm pCO}_{2,\text{sw}} \text{ yr}^{-1}$, $\mu\text{mol DIC kg}^{-1} \text{ yr}^{-1}$, $\mu\text{mol TA kg}^{-1} \text{ yr}^{-1}$, $\text{pH}_{\text{Total}} \text{ yr}^{-1}$)

Annual mean carbonate chemistry (CO2sys) within reef environment. ($\mu\text{mol kg}^{-1}$)

- Primary Measured Parameters: Temp ($^{\circ}\text{C}$), Salinity (unitless), TCO_2 ($\mu\text{mol kg}^{-1}$), TAlk ($\mu\text{mol kg}^{-1}$), $\text{pCO}_{2,\text{sea}}$ (μatm)
- Secondary Measured Parameters: pH (Total), $\text{pCO}_{2,\text{air}}$ (μatm), Total P ($\mu\text{mol kg}^{-1}$), Total Si ($\mu\text{mol kg}^{-1}$)

Annual mean carbonate chemistry (CO2sys) within neighboring open ocean surface waters. ($\mu\text{mol kg}^{-1}$)

- Primary Measured Parameters: Temp ($^{\circ}\text{C}$), Salinity (unitless), TCO_2 ($\mu\text{mol kg}^{-1}$), TAlk ($\mu\text{mol kg}^{-1}$), $\text{pCO}_{2,\text{sea}}$ (μatm)
- Secondary Measured Parameters: pH (Total), $\text{pCO}_{2,\text{air}}$ (μatm), Total P, Total Si

150

151 **Figure 3. The CROAMP requirements for long-term monitoring of the *direct changes in***
 152 ***chemistry in response to OA.***

153 Estimates of the rate of change in aragonite saturation state (Ω_{arg}) within tropical surface waters
 154 is on the order of 0.1 Ω_{arg} per decade. While such rates are readily discernible from multi-
 155 decadal time-series stations within oceanic waters using sampling frequencies as coarse as
 156 monthly, the variability within reef environments can be an order of magnitude greater
 157 demanding much high sampling frequencies.TO ADD QUANTITATIVE ANALYSIS OF
 158 TIME SERIES DATA TO EVALUATE MINIMUM FREQUENCY.

159 For monitoring within the reef environments, subdaily (3-hourly) is probably preferred to avoid
 160 aliasing the data but bi-weekly discrete sampling is probably the minimum sampling frequency.

161 If adequate preliminary data is available, an appropriate frequency should be determined using a

162 Fourier series. The oceanic reference station will depend on the open ocean ecosystem chosen
163 for comparison, but in many cases, seasonal to monthly measures may be adequate.

164

165 Site criteria... To skillfully detect changes in response to OA it will be important to select sites
166 where local biogeochemistry is reasonably constrained and coastal processes are a minimal
167 influence. However, it is recognized that at least a few sites should be chosen to monitor such
168 changes within the context of local anthropogenic influences for comparative purposes. At those
169 sites, additional effort may be required to characterize influencing coastal processes. This
170 tactical monitoring index is best suited to fixed time-series stations as it is unlikely that stratified
171 sampling could be conducted at sufficient temporal frequency to avoid aliasing the data for
172 regression analysis. However, in systems of relatively lower short-term variability (e.g. low
173 residence time systems) it may be possible to derive this index from stratified sampling provided
174 it was maintained for several decades.

175 co-variants - Most of these measures will co-vary with temperature, salinity, and any processes
176 which might alter the CO₂SYs.

177

178 *Methods:*

179 Provided a measure of any two of these four parameters, the other two can theoretically be
180 solved. To do so, the CO₂ System needs to be solved using selected dissociation constants. The
181 carbonate equilibria calculations should use the Mehrbach et al. [1973] formulations of the K₁
182 and K₂ dissociation constants as refit by Dickson and Millero [1987]. The sensitivity with

183 which the saturation states and CO₂-3 can be calculated is dependent on the parameter
184 measured; pH and pCO₂ yield the greatest uncertainty, followed by the TA, DIC pair. While
185 triple constraint is advised within coastal systems, if only two parameters are to be measured, pH
186 or pCO₂ with DIC yields the greatest precision, closely followed by pH or pCO₂ and TA. All
187 measurements should follow procedures and protocols outlined in the Best Practices guides
188 (Dickson et al., 2007; Riebesell et al., 2010).

189

190

191 *2. Community Metabolism Response and Feedbacks to Ocean Acidification*

192 Changes in community-scale processes including net community productivity (P), respiration
193 (R), calcification (G), and dissolution (D) can significantly impact the local surrounding
194 chemistry thereby overprinting the affects of OA in some cases. Understanding the interactions
195 between coral reef communities and the surrounding chemical environment is critical towards
196 improving our understanding with regards to how OA is unfolding within these systems and
197 what local processes might prove dominant.

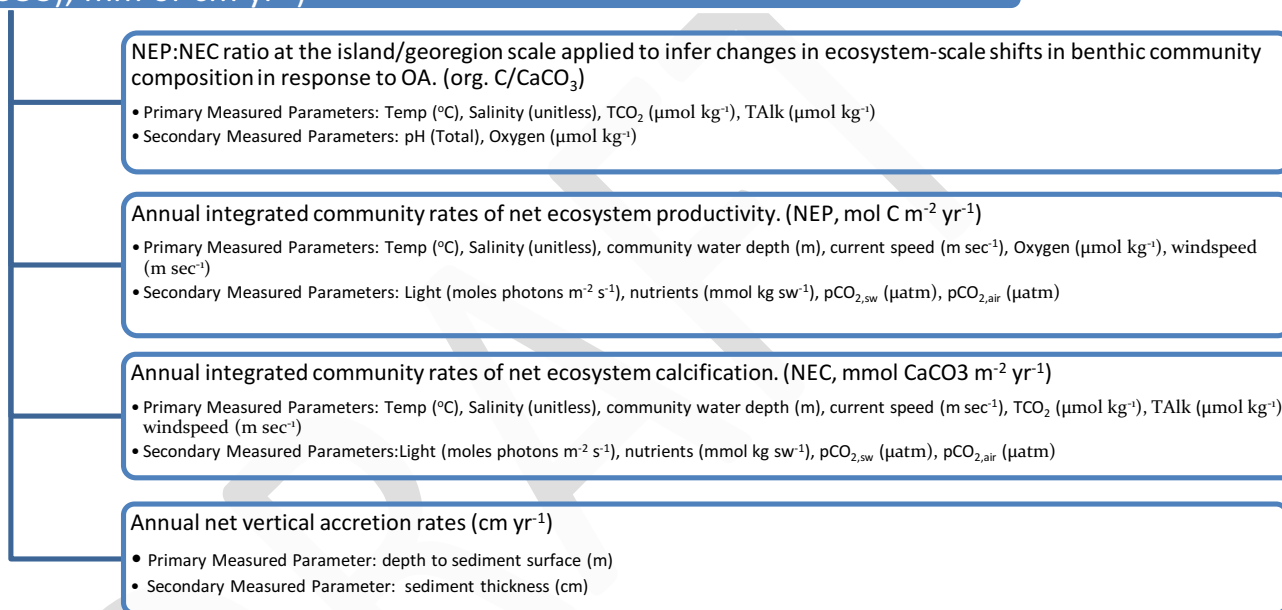
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199

200 Ocean acidification may profoundly affect the basic ecological interactions which structure coral
201 reef ecosystems, providing even greater challenges for local management strategies aimed at
202 retaining or regaining coral dominance. A recent meta-analysis of available experimental data

203 suggests that such ‘phase shifts’ might increase in response to OA. Specifically, as calcification
 204 is reduced, algae (non-calcifying) and sea grass may benefit [Hendriks et al., 2010].

What are the long-term trends in annual integrated net community metabolic performance? (e.g. mol C m⁻² yr⁻¹, mol CaCO₃ m⁻² yr⁻¹, NEP:NEC (C/CaCO₃), mm or cm yr⁻¹)



205

206 **Figure 4. The CROAMP requirements for long-term monitoring *Community Metabolism***
 207 ***Response to OA.***

208

209 An alternative approach to paired TA and DIC measures might be tracking shifts in the
 210 regression slope of pH vs O₂. While simpler to implement than the paired measurements, this

211 approach remains largely speculative at this point and needs to be better documented. Good
212 quality autonomous instruments are increasingly available for short duration (e.g. weeks to a few
213 months) deployments. High quality pH measures are best obtained through discrete sample
214 collection according to best practices. Spectrophotometric determination can then be performed
215 post-collection. Discrete samples need to be measured within an hour of collection.

216

217 Site criteria - To skillfully detect community changes in response to OA it will be important to
218 select sites where local biogeochemistry is reasonably constrained and coastal processes are a
219 minimal influence. However, it is recognized that at least a few sites should be chosen to
220 monitor such changes within the context of local anthropogenic influences for comparative
221 purposes.

222

223 *3. Dissolution and Bioerosion Response and Feedbacks to Ocean Acidification*

224 Some species exhibit a natural tolerance or local environmental conditions can compensate for
225 global OA effects. It has now been well established that coral calcification rates are, in fact, a
226 function of multiple factors including light, temperature, carbonate chemistry and nutrient
227 supply.

228 Beyond the potential direct effects for coral calcification, OA poses additional concerns for coral
229 reef ecosystems. These include dissolution of coral reef sediments which often contain
230 appreciable amounts of more soluble carbonate minerals [Morse et al., 2006]. Sediment
231 dissolution may outpace carbonate production on many reefs by 2030 [Yates and Halley, 2006].

232 Furthermore, current-day areas where upwelling causes chemistries comparable to future OA
 233 conditions (i.e. a tripling of atmospheric CO₂) exhibit poorly cemented and highly bioeroded
 234 coral reefs [Manzello et al., 2008]. These effects may compromise coral reef framework integrity
 235 and resilience in the face of other acute threats such as coral bleaching, diseases, increases in
 236 storm intensity, and rising sea level [e.g., Silverman & Calderia, 2009]. Indeed, in CO₂ enriched
 237 waters of the Galapagos Islands, reef structures were completely eroded to rubble and sand in
 238 less than 10 years following one acute warming disturbance [i.e., 1982-83 El Nino event:
 239 Manzello, 2009].

240

What are the long-term trends in mechanical breakdown/dissolution rates of CaCO₃? (g CaCO₃ m⁻² yr⁻¹)

Annual mean alkalinity anomalies relative to neighboring open ocean surface waters (μmol kg⁻¹).

- Primary Measured Parameters: Temp (°C), Salinity (unitless), TAlk (μmol kg⁻¹)
- Secondary Measured Parameters: Total P (μmol kg⁻¹), Total Si (μmol kg⁻¹)

Bioerosion rates at specific sites. (g CaCO₃ m⁻² yr⁻¹)

- Primary Measured Parameters: Bioeroder colonization density (number m⁻²) and rate (number m⁻² yr⁻¹) of experimental substrate
- Secondary Measured Parameters: Epibenthic bioeroder density (individuals m⁻²), Bioeroder size frequency distribution (number of size classes m⁻²)

Metabolic CaCO₃ dissolution rates. (g CaCO₃ m⁻² yr⁻¹)

- Primary Measured Parameters: Temp (°C), Salinity (unitless), TAlk sediment profiles (μmol kg⁻¹), TAlk sediment-seawater flux (μmol m⁻² day⁻¹), Mass loss (g CaCO₃) of experimental substrate
- Secondary Measured Parameters: Light (moles photons m⁻² s⁻¹), nutrients (mmol kg sw⁻¹), total organic carbon (TOC, mg m⁻³)

Architectural complexity. (m m⁻¹)

- Primary Measure Parameters: Rugosity (m m⁻¹)
- Secondary Measured Parameters:

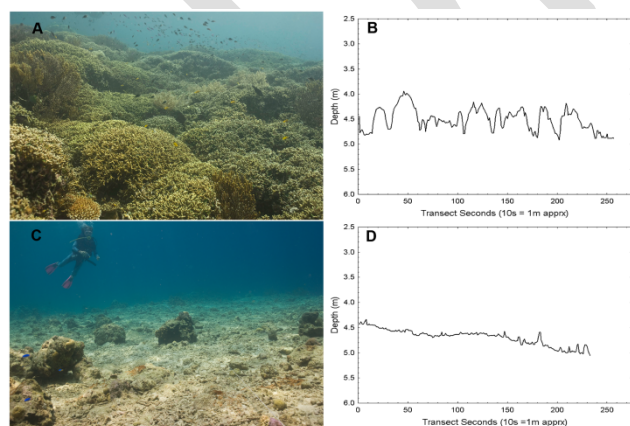
Sediment mineralogy/composition (mol %)

- Primary Measured Parameter: carbonate mineral absolute (g CaCO₃ g⁻¹ sediment) and relative abundance (calcite:aragonite:high-MgCO₃), mol fraction MgCO₃ (%)
- Secondary Measured Parameter: grain size distribution (# m⁻²)

241

242 **Figure 5. The CROAMP requirements for long-term monitoring of the *Dissolution and***
243 ***Bioerosion Response* in response to OA.**

244 A measure of small-scale variations or amplitude in the height of a reef surface. Rugosity is a
245 measure of architectural complexity which is presumed to be an indicator of the amount of
246 habitat available for colonization by benthic organisms, and shelter and foraging area for fish.
247 Tracking rugosity over time can be a useful metric to examine structural changes in reefs as the
248 framework building corals die, rates of bioerosion overtake accretion, and the reefs eventually
249 flatten [18] although care must be taken to account for acute disturbance events (e.g. storm
250 effects and mass wasting) when assigning attribution to long-term trends. Structural complexity
251 is classically measured with a reef rugosity index, which is the ratio of a straight line transect to
252 the distance a flexible chain of equal length travels when draped over the reef substrate, however
253 recent advances in digital reef rugosity (DRR) have greatly reduced that labor required to obtain
254 a rugosity measure while making it a more objective and repeatable measure useful in statistical
255 analysis. Dustan P, Doherty O, Pardede S (2013) Digital Reef Rugosity Estimates Coral Reef
256 Habitat Complexity. PLoS ONE 8(2): e57386. doi:10.1371/journal.pone.0057386



257

258

259 *4. Organism Response to Ocean Acidification*

260 Other expected impacts of OA include a potential lowering of the thermal thresholds for
261 bleaching [Anthony et al., 2008], the impairment on early life stages of corals such as reduced
262 fertilization success, reduced larval settlement, and reduced growth and survival rates of newly
263 settled corals [Albright et al., 2008; Cohen and Holcomb, 2009; Albright et al., 2010, Morita et
264 al., 2010; Suwa et al., 2010]. These species-level impacts are particularly relevant given the
265 potential extension of Endangered Species Act protection for additional scleractinian coral reef
266 species as this Act provides for potent conservation actions for species at risk of extinction, and
267 impacts on population bottlenecks such as recruitment may pose more direct extinction risk than
268 calcification and growth impacts.

269
270 The affects of OA on some non-calcifying organisms may indeed be just as serious as the affects
271 to calcifiers but are largely unexplored.

272
273 In addition, predictions of coral reef ecosystem responses to OA are further complicated due to
274 the fact that reef organisms secrete species-specific types of calcium carbonate mineralogies (i.e.,
275 aragonite, calcite, and magnesium calcite) which exhibit a range of solubilities.

276

What are the long-term regional changes in growth rates of target taxa in response to OA?

Growth rates of select calcifiers.

- Primary Measured Parameters: Coral skeletal linear extension rate (mm yr^{-1}), density (g cm^{-3}), and mass (g)
- Secondary Measured Parameters: *Halimeda* spp. (or other calcified algal taxa) growth rate (g yr^{-1}), Other taxa (e.g. forams, urchins) growth rate (g yr^{-1})

Growth and productivity rates in select non-calcifiers. ($\text{g colony}^{-1} \text{yr}^{-1}$)

- Primary Measured Parameters: seagrass spp. plant mass (g), shoot density ($\# \text{m}^{-2}$), and photosynthetic rate ($\mu\text{g C } \mu\text{g chlorophyll } \alpha^{-1} \text{ hr}^{-1}$); fleshy algal spp. plant mass (g) and photosynthetic rate ($\mu\text{g C } \mu\text{g chlorophyll } \alpha^{-1} \text{ hr}^{-1}$)
- Secondary Measured Parameters: Temp ($^{\circ}\text{C}$), Light (moles photons $\text{m}^{-2} \text{s}^{-1}$), nutrients (mmol kg sw^{-1})

277

What are the long-term regional changes in 'recruitment' rates of target taxa in response to OA.

Crustose coralline algae accumulation rate (CCA, mm yr^{-1}).

- Primary Measured Parameters: CCA accumulation rate on standardized substrates (mm yr^{-1})
- Secondary Measured Parameters: Temp ($^{\circ}\text{C}$), Light (moles photons $\text{m}^{-2} \text{s}^{-1}$), nutrients (mmol kg sw^{-1}), current speed (cm sec^{-1}), grazing rate ($\text{g grazer}^{-1} \text{d}^{-1}$)

Calcifying epiphytes accumulation rate. ($\mu\text{g Chl } a \text{ cm}^{-2} \text{yr}^{-1}$)

- Primary Measured Parameters: seagrass epiphyte density ($\mu\text{g Chl } a \text{ cm}^{-2}$)
- Secondary Measured Parameters: Temp ($^{\circ}\text{C}$), Light (moles photons $\text{m}^{-2} \text{s}^{-1}$), nutrients (mmol kg sw^{-1}), current speed (cm sec^{-1}), grazing rate ($\text{g grazer}^{-1} \text{d}^{-1}$)

278

What are the long-term regional changes in physiological 'condition' of individual organisms response to OA?

Plan carbon/nutrient status.

- Primary Measured Parameters: Seagrass tissues C:N (moles C moles N⁻¹), Algal tissues C:N (moles C moles N⁻¹)
- Secondary Measured Parameters: Temp (°C), Light (moles photons m⁻² s⁻¹), nutrients (mmol kg sw⁻¹), current speed (cm sec⁻¹)

Organism mineral content.

- Primary Measured Parameters: calcifier structural mineralogy (polymorph)
- Secondary Measured Parameters: Temp (°C), Salinity (unitless), TCO₂ (μmol kg⁻¹), TAlk (μmol kg⁻¹), pCO_{2,sea} (μatm)

Coral tissue thickness.

- Primary Measured Parameters: tissue thickness (mm)
- Secondary Measured Parameters: Temp (°C), Light (moles photons m⁻² s⁻¹), nutrients (mmol kg sw⁻¹), current speed (cm sec⁻¹)

279

280 **Figure 6. The CROAMP requirements for long-term monitoring of the *Organism Response***
281 **in response to OA.**

282

283 *5. Changes in Coral Reef Biodiversity to Ocean Acidification*

284 While coral reefs are considered the most biologically diverse of all marine ecosystems this
285 diversity varies greatly across spatial and environmental gradients. How this variation influences
286 reef resilience and adaptation to climate change and OA is poorly understood. Much of the
287 biomass and biodiversity of reefs lies within the complex architecture of the reef matrix
288 [Ginsburg, 1983; Small et al., 1998; Sala and Knowlton, 2006; Knowlton et al., 2010]. The
289 potential effects of OA on this community of organisms, collectively known as the cryptobiota
290 [Macintyre et al., 1982], remains largely unknown.

291

292 Understanding the biodiversity and community structure of coral reef ecosystems is therefore
293 necessary in recognizing and predicting shifts in community structure in response to OA.

294

295

296

What are the long-term trends in biodiversity as apparent from changes in benthic composition, community structure, and ecological function.?

Biomass, population and trophic structure of cryptobiota.

- Primary Measured Parameters: ID/richness (OTU, Organismal Taxonomic Unit), absolute (OTU #) and relative abundance (%), density (# m⁻²), diversity index (unitless), evenness (unitless), sessile invertebrate percent cover (%), rugosity (m m⁻¹), benthic cover (%)
- Secondary Measured Parameters: Temp (°C), Salinity (unitless), TCO₂ (μmol kg⁻¹), TAlk (μmol kg⁻¹), Light (moles photons m⁻² s⁻¹), nutrients (mmol kg sw⁻¹), current speed (cm sec⁻¹)

Population structure of corals.

- Primary Measured Parameters: ID/richness (OTU, Organismal Taxonomic Unit), absolute (OTU #) and relative abundance (%), density (# m⁻²), diversity index (unitless), evenness (unitless), percent cover (%), rugosity (m m⁻¹)
- Secondary Measured Parameters: Temp (°C), Salinity (unitless), TCO₂ (μmol kg⁻¹), TAlk (μmol kg⁻¹), Light (moles photons m⁻² s⁻¹), nutrients (mmol kg sw⁻¹), current speed (cm sec⁻¹)

Population structure of algae.

- Primary Measured Parameters: ID/richness (OTU, Organismal Taxonomic Unit), absolute (OTU #) and relative abundance (%), density (# m⁻²), diversity index (unitless), evenness (unitless), percent cover (%), rugosity (m m⁻¹)
- Secondary Measured Parameters: Temp (°C), Salinity (unitless), TCO₂ (μmol kg⁻¹), TAlk (μmol kg⁻¹), Light (moles photons m⁻² s⁻¹), nutrients (mmol kg sw⁻¹), current speed (cm sec⁻¹)

Changes in the population structure of urchins.

- Primary Measured Parameters: ID/richness (OTU, Organismal Taxonomic Unit), absolute (OTU #) and relative abundance (%), density (# m⁻²), diversity index (unitless), evenness (unitless), mean test size (cm), rugosity (m m⁻¹), test size frequency distribution (cm), benthic cover (%)
- Secondary Measured Parameters: Temp (°C), Light (moles photons m⁻² s⁻¹), nutrients (mmol kg sw⁻¹), current speed (cm sec⁻¹)

297

298 **Figure 7. The CROAMP requirements for long-term monitoring of the *Coral Reef***
299 ***Biodiversity* response to OA.**

300

301 **IV. Existing Ocean Acidification Monitoring Capacity**

302 **V. Existing Ocean Acidification Monitoring Capacity**

303 **VI. Summary**

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309 the ability and rate of calcium carbonate production by many marine calcifiers is anticipated to
310 decrease while rates of bioerosion and dissolution could increase. Such affects would have
311 important consequences for the role and function of coral reefs and compromise the ecoservices
312 they afford us.

313 **References.**

314 **Tables.**

315 Summary of Measurements

<u>Measurement (units)</u>	<u>Description</u>	<u>Methods</u>	<u>References</u>
<i>Physicochemical</i>			
<i>Nutrients (mmol kg sw⁻¹)</i> –	These include all dissolved inorganic nutrients (ammonium, silicate, nitrate, nitrite, and phosphate).	Discrete water sample collection followed by post-collection analysis according to best-practices.	
Oxygen (μmol kg ⁻¹) –		Discrete water sample collection followed by post-collection analysis according to best-practices.	
pCO _{2,air} (μatm) –			
pCO _{2,sw} (μatm) –			
pH (Total) –			
Salinity (unitless) –	Salinity is a fundamental measurements for interpreting carbonate chemistry and must be obtained contemporaneously with any measurement of CO ₂ SYN.		
TAlk (μmol kg ⁻¹) –			
TCO ₂ (μmol kg ⁻¹) –			
Temperature (oC) –	Temp. is a fundamental measurements for interpreting carbonate chemistry and must be obtained contemporaneously with any measurement of CO ₂ SYN. Furthermore, it imparts a fundamental control on biology and thus needs to be a key measurement ecological monitoring.	Autonomous temperature sensors (e.g. CTD). Do not rely solely on manufacturers' stated accuracy; regular lab calibrations and at-sea comparisons are recommended	
Total organic carbon (TOC, mg m ⁻³) –			

Total P ($\mu\text{mol kg sw-1}$) –			
Total Si ($\mu\text{mol kg sw-1}$) –			
Ecological			
<i>[Coral, algae, urchin and algae] absolute (OTU #) and relative abundance (%) –</i>			
<i>[Coral, algae, urchin and algae] benthic cover (%) –</i>			
<i>[Coral, algae, urchin and algae] diversity index (unitless) –</i>			
<i>[Coral, algae, urchin and algae] ID/richness (OTU, Organismal Taxonomic Unit) –</i>			
<i>[Coral, algae, urchin and algae] percent cover (%) –</i>			
<i>[Coral, algae, urchin and algae] density ($\# \text{ m}^{-2}$) –</i>			
<i>[Coral, algae, urchin and algae] evenness (unitless) –</i>			
<i>Algal tissues C:N (moles C moles N-1) –</i>			
<i>Bioeroder colonization density (number m^{-2}) –</i>			
<i>Bioeroder colonization rate (number $\text{m}^{-2} \text{ yr}^{-1}$) –</i>			
<i>Bioeroder size frequency distribution (number of size classes m^{-2}) –</i>			
<i>Biomass –</i>			
<i>Calcifier structural</i>			

<i>minerology (polymorph) –</i>			
<i>Coral skeletal density (g cm⁻³) –</i>			
<i>Coral skeletal linear extension rate (mm yr⁻¹) –</i>			
<i>Coral skeletal mass (g) –</i>			
<i>Coral tissue thickness (mm) –</i>			
<i>Crustose coralline algae accumulation rate (CCA, mm yr⁻¹) –</i>			
<i>Density of individuals (per m²) –</i>			
<i>Epibenthic bioeroder density (individuals m⁻²) –</i>			
<i>Fleshy algal spp. Photosynthetic rate (µg C µg chlorophyll a⁻¹ hr⁻¹) –</i>			
<i>Fleshy algal spp. plant mass (g) –</i>			
<i>Grazing rate (g grazer⁻¹ d⁻¹) –</i>			
<i>Halimeda spp. (or other calcified algal taxa) growth rate (g yr⁻¹) –</i>			
<i>Identification –</i>			
<i>Mass loss of experimental substrate (g CaCO₃) –</i>			
<i>Other taxa (e.g. forams, urchins) growth rate (g yr⁻¹) –</i>			
<i>Percent cover (%) –</i>			
<i>Seagrass epiphyte density (µg Chl a cm⁻²) –</i>			

<i>Seagrass spp. photosynthetic rate ($\mu\text{g C } \mu\text{g chlorophyll a}^{-1} \text{ hr}^{-1}$) –</i>			
<i>Seagrass spp. plant mass (g) –</i>			
<i>Seagrass spp. shoot density ($\# \text{ m}^{-2}$) –</i>			
<i>Seagrass tissues C:N (moles C moles N⁻¹) –</i>			
<i>Substrate type –</i>			
Hydrodynamic and Structural			
<i>Carbonate mineral absolute abundance ($\text{g CaCO}_3 \text{ g}^{-1} \text{ Sediment}$) –</i>			
<i>Carbonate mineral relative abundance (calcite:aragonite:high-MgCO₃) –</i>			
<i>Community water depth (m) –</i>			
<i>Current speed (m sec^{-1}) –</i>			
<i>Depth to sediment surface (m) –</i>			
<i>Grain size distribution ($\# \text{ m}^{-2}$) –</i>			
<i>Mole fraction MgCO₃ (%) –</i>			
<i>Rugosity (STD depth, m) –</i>	<i>A measure of small-scale variations or amplitude in the height of a reef surface. Rugosity is a measure of architectural complexity which is presumed to be an indicator of the amount of habitat available for colonization by benthic organisms, and shelter and foraging area for fish.</i>	digital reef rugosity (DRR)	Dustan et al., (2013)

<i>Sediment thickness (cm) –</i>			
<i>Talk sediment profiles ($\mu\text{mol kg}^{-1}$) –</i>			
<i>Talk sediment-seawater flux ($\mu\text{mol m}^{-2} \text{day}^{-1}$) –</i>			
<i>Meteorological</i>			
<i>Light (moles photons $\text{m}^{-2} \text{s}^{-1}$) –</i>			
<i>Windspeed (m sec^{-1}) –</i>			

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