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13 Abstract

This chapter focuses on considerations for conducting open-system field experiments in the 14 15 context of ocean alkalinity enhancement (OAE) research. By conducting experiments in realworld marine and coastal systems, researchers can gain valuable insights into ecological 16 dynamics, biogeochemical cycles, and the safety, efficacy, and scalability of OAE techniques 17 under natural conditions. However, logistical constraints and complex natural dynamics pose 18 19 challenges. To date, only a limited number of OAE field studies have been conducted, and guidelines for such experiments are still evolving. Due to the fast pace of carbon dioxide removal 20 (CDR) research and development, we advocate for openly sharing data, knowledge, and lessons 21 learned as quickly and efficiently as possible within the broader OAE community and beyond. 22 Considering the potential ecological and societal consequences of field experiments, active 23 engagement with the public and other stakeholders is desirable while collaboration, data sharing, 24 and transdisciplinary scientific teams can maximize the return on investment. The outcomes of 25 early field experiments are likely to shape the future of OAE research, implementation, and 26 public acceptance, emphasizing the need for transparent and open scientific practices. 27

29 **1. Introduction**

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This chapter addresses considerations for conducting open-system field experiments related to 31 ocean alkalinity enhancement (OAE). We define 'field experiment' or 'field studies' broadly as 32 the addition or manipulation of alkalinity in a natural system that is relevant to OAE, 33 independent of the spatial and temporal scale. We intentionally exclude spatial and temporal 34 scales from our definition to encompass the wide spectrum of OAE methods and approaches. In 35 fact, field experiments are likely to span spatial scales of m² to 100s of km² and last from days to 36 years. Field experiments and studies differ from both 'field trials' and 'field deployments' in their 37 motivation, as both trials and deployments denote the practical application and usage of a 38 specific product, device, or technology. The scientific focus during field trials is likely to be on 39 the efficacy of Carbon Dioxide Removal (CDR) and fine-tuning operational deployment, while 40 field experiments will encompass a broader range of scientific goals and objectives. The nature, 41 logistics, and objectives of field experiments are likely to make them smaller in scale than 42 operational deployments. This will be advantageous, as field experiments that emulate planned 43 44 OAE trials and deployments will help create the scientific framework needed to scale operational OAE safely and responsibly. 45

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The benefits of conducting experiments in natural systems include observing complex ecological 47 48 dynamics and impacts at the ecosystem level, understanding the role of biogeochemical cycles and physical processes that cannot be replicated in other settings, and assessing CDR under real-49 world scenarios. The complexity and breadth of some field experiments will necessitate science 50 that transcends disciplinary boundaries, making collaboration a priority. Success in the field 51 52 faces many challenges due to the inherent complexity of natural systems along with limiting logistical constraints (e.g., permitting, access, social license, infrastructure, life cycle emissions). 53 Despite these challenges, the first OAE field experiments are already underway, many of which 54 are small-scale representations of scalable OAE approaches. There will be much to learn from 55 these early studies, and any knowledge or insights gained should be shared as efficiently and 56 57 openly as possible within the wider OAE community and beyond.

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59 While some OAE field experiments have been completed or are already in progress, many more

are on the horizon. We recommend that three overarching questions be taken into consideration,
especially when in the planning stages:

62 What are the main goals of the experiment?

63 Establishing the objectives of a field experiment early in the planning stage will help guide all aspects of the scientific research plan, including site selection, measurement techniques and 64 approaches, data analysis, and measured outcomes. Potential overarching goals of OAE field 65 experiments include demonstrating functionality, efficacy, process, and/or scalability, 66 67 determining ecological and environmental impacts, developing measurement, reporting, and verification (MRV) protocols, and assessing community engagement. Life cycle assessments 68 69 (LCA) may be a critical learning objective for some projects (e.g., Foteinis et al. 2023), especially those that are examining OAE at the scale of operational deployments. This list of 70 71 overarching goals is not comprehensive, and goals are not necessarily mutually exclusive. For example, larger projects may wish to assess multiple components of an OAE approach while 72 73 smaller projects might be highly focused.

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75 What is the type of alkalinity perturbation?

The type of alkalinity that is added (e.g., aqueous vs. solid, carbonates, hydroxides, oxides, or 76 naturally occurring (ultra-)mafic rocks) will ultimately determine many aspects of the scientific 77 research plan. For example, projects adding ground alkaline minerals (e.g., olivine) to the ocean 78 79 may have different goals and timelines than projects that add aqueous alkalinity (e.g., liquid 80 NaOH) (see Eisaman et al., 2023, this Guide). Priorities for projects adding ground material might include tracking the dissolution of the alkaline material plus monitoring the fate of the 81 dissolved alkalinity and its dissolution co-products (e.g. trace metals), while projects adding 82 aqueous alkalinity will likely be more concerned with the latter. Other important experimental 83 considerations that will be driven by the type of alkalinity perturbation include the concentration 84 of added alkalinity, duration of additions, dilution and advection at the field site, residence time, 85 air-sea equilibration, co-deployed tracers, sampling scheme, and environmental side-effects. 86 These and other research considerations are discussed in more detail below. 87

88 What are the permitting constraints and wider social implications?

Addressing the appropriate regulatory requirements is essential before any field experiment can 89 move forward. Permitting requirements will be influenced by the study location, type of 90 91 alkalinity perturbation, spatial scale, and duration. The use of existing infrastructure (e.g., wastewater discharge sites) and environmental projects (e.g., beach renourishment) may offer 92 ways to facilitate alkalinity perturbations under existing regulatory frameworks. Community 93 engagement and outreach are other areas that will be important to address, especially when the 94 alkalinity perturbation is large and uncontained. Ideally, local communities should be engaged at 95 the earliest possible stage since social license to operate is critical for the success of CDR 96 projects (Nawaz et al., 2022). For a more detailed discussion of legal and social issues see 97 Steenkamp et al. (2023, this Guide) and Satterfield et al. (2023, this Guide). 98

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With these overarching questions in mind, we discuss considerations for OAE field experimentsin more detail below.

102 2. Research Methods

103 **2.1 Types of alkalinity addition**

Field experiments of OAE present many challenges. One of the biggest obstacles to success is 104 105 tracking alkalinity added to an open system. Methods for adding alkalinity can be divided into two general approaches: (1) in situ or coastal enhanced weathering from the addition of ground 106 alkaline minerals and rocks with the expectation they will dissolve directly in seawater, and (2) 107 aqueous alkalinity additions, or the addition of 'pre-dissolved' alkalinity to seawater that can be 108 generated in numerous ways including through dissolution reactors and electrochemical 109 techniques (Eisaman et al., 2023, this Guide). Tracking the added alkalinity, and subsequent 110 CDR, under each approach comes with its own unique set of challenges and considerations. 111

Adding ground minerals and rocks to an open system presents two distinct scientific challenges. First, for alkalinity to be considered additional it needs to be attributed to the dissolution of the solid material. This can be accomplished through a range of techniques including measuring the loss of mass of the added material or using geochemical tracers in the receiving waters. Determining dissolution kinetics *in situ* will be particularly important and they are likely to vary between different deployment environments and strategies (e.g., coastal vs open ocean). For

example, the chemistry (e.g., salinity, pH, temperature) of the waters where the mineral is added 118 could vary significantly depending on the environment (e.g., beach face, estuary, continental 119 shelf). Chemical (e.g., seawater conditions such as salinity, pCO₂, and silica concentrations) and 120 physical (e.g., grain size and surface area of the added material) will be critical in determining 121 dissolution rates (Rimstidt et al., 2012; Montserrat et al., 2017; Fuhr et al. 2022). Physical 122 abrasion through wave action and currents is also likely to be an important control on dissolution 123 (Flipkens et al., 2023). Field experiments will help translate dissolution kinetics from laboratory 124 and mesocosm experiments to natural systems, which is not often straightforward due to 125 complicated biogeochemical processes that are hard to replicate ex situ (Morse et al., 2007). 126

127 The second major challenge is common to both solid and aqueous approaches and involves tracking the added alkalinity, which becomes a particularly difficult problem in open-system 128 129 field experiments where water is freely exchanged. Depending on the objectives of the field deployment, this is likely to be a main scientific concern. However, it is important to note that 130 tracking the added alkalinity does not necessarily equate to observing CDR (i.e., an increase in 131 seawater CO₂ stored as bicarbonate or carbonate). Observing an increase in atmospheric CO₂ 132 stored as seawater dissolved inorganic carbon comes with its own set of challenges that are 133 134 discussed in depth by Ho et al. (2023, this Guide).

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Whether or not the alkalinity is derived from in situ mineral dissolution or direct aqueous 136 additions, for OAE to be successful atmospheric CO₂ needs to be taken up by seawater or CO₂ 137 138 effluxes from seawater to the atmosphere need to be reduced. Therefore, understanding the physical mixing and air-sea gas exchange dynamics of the deployment site will be a factor of 139 interest for many field studies. Incorporating physical mixing models with biogeochemical 140 processes will likely be the end goal of many field experiments focused on MRV (Ho et al. and 141 Fennel et al., 2023, this Guide). Choosing sites with minimal mixing of different water masses or 142 with well-defined diffusivities could facilitate tracing released alkalinity and subsequent air-sea 143 CO₂ fluxes. While minimal mixing of different ocean water masses may be desired, higher wind 144 speeds and wave action will increase the rate of air-sea gas exchange and may make CDR easier 145 to measure. Background seawater chemistry will also be important in controlling air-sea gas 146 147 exchange. For example, sites with naturally lower buffering capacities will see greater changes in CO₂ per unit of added alkalinity (Egleston et al., 2010; Hauck et al., 2016). The release of 148

149 conservative tracers will likely be useful for field experiments that wish to track the added150 alkalinity and is discussed in more detail below (Section 2.5).

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Other experimental considerations related to the type of alkalinity perturbation include the 152 duration and location of alkalinity addition, which will be important for environmental and 153 regulatory considerations. Alkalinity can be added once, in timed doses, or continuously. 154 Aqueous alkalinity could be added directly to seawater, but the rate of this addition will likely be 155 important, especially for avoiding secondary precipitation (Hartmann et al., 2023; Moras et al., 156 2022, Fuhr et al., 2022). Compared to experiments based on one-time additions of aqueous 157 alkalinity or fast dissolving solid-phase materials (e.g., Ca(OH)₂), field experiments adding solid 158 minerals with comparatively slow dissolution rates (e.g., olivine) will likely need to consider 159 longer experimental time frames to incorporate the monitoring of mineral dissolution. However, 160 the timescale of each experiment will ultimately depend on the scientific objectives and could 161 last from weeks to years and even decades. Location is another important factor that will 162 influence logistics. For example, amending beach sand with alkaline minerals will present 163 164 different challenges compared to the addition of alkaline material to outfalls that discharge into the ocean. Based on these and other considerations, each field experiment will require specific 165 166 spatial and temporal sampling schemes to be developed. These sampling schemes should be planned well in advance of any perturbation and may require preliminary sampling campaigns to 167 168 fine tune.

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170 **2.2 Alkalinity sources**

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172 OAE via coastal enhanced weathering can be accomplished using a variety of naturally occurring 173 and human-made rocks and minerals (Table 1). The addition of these rocks and minerals is done after they have been ground to a desired grain size with many unique application techniques 174 proposed after the initial grinding step (see Eisaman et al., 2023, this Guide). The simplest 175 application is done via sprinkling the ground material on the ocean surface, although this has 176 177 many disadvantages including sinking and advection of the material before it dissolves (Koehler et al., 2013; Fakharee et al., 2023), although deployment in boat wakes may be viable (Renforth 178 et al., 2017; He and Tyka, 2023). Other application techniques include spreading material in 179

coastal ecosystems such as on beaches, marshes, riverbeds, and estuaries, which have the
potential to enhance dissolution through processes such as physical wave action and favorable
water chemistry. However, the complex physical and biogeochemical processes that promote
enhanced weathering in coastal ecosystems can make field experimentation more complicated by
creating strong spatiotemporal modes of variability in water chemistry. To make results more
broadly applicable, field experiments should attempt to mimic real world alkalinity application
scenarios such as those described above.

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Any field experiments that add ground material to marine ecosystems may consider tracking the fate of that material from the addition site. Experiments could also artificially contain the material using barriers to avoid rapid loss of the ground material via currents, however, this could make the experiment less comparable to real world OAE deployments. Sampling should extend from the water column into areas where the material is added including sediments and pore waters.

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195 Likely environmental impacts associated with coastal enhanced weathering come from the physical impacts of adding finely ground material or the chemical release of trace elements and 196 197 other contaminants. Both processes could have associated risks and/or co-benefits for a range of ecological processes and biogeochemical cycles (Bach et al., 2019). For example, the addition of 198 finely ground material could lead to increased turbidity from the initial addition, subsequent 199 resuspension, or secondary precipitation of particulates in the water column. Additionally, any 200 release of nutrients or heavy metals from the dissolving material could alter primary production 201 or cause harm to biological systems. The bioaccumulation of toxic metals in higher trophic level 202 203 organisms, especially those of commercial importance, is a widespread concern.

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Safety criteria should be put in place that can create a pause in the field experiment or prevent future experiments of the same type from taking place. These guardrails should be developed by the broader OAE community but may include obvious damage or health impacts to ecologically important organisms such as primary producers and keystone species, large and unexpected changes in biogeochemical cycles, and the general deterioration of environmental conditions. Risk-benefit analysis may be particularly useful in determining whether projects can or should move forward and may already be included in regulatory requirements through existing
 frameworks such as environmental impact assessments.

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Aqueous and slurry-based additions of alkalinity provide different benefits and challenges 214 compared to solid forms of alkalinity feedstock. One of the primary benefits of aqueous 215 additions is that the alkalinity has been pre-dissolved, avoiding the often slow dissolution 216 kinetics of minerals and rocks in seawater. Aqueous alkalinity can be generated by two main 217 mechanisms (1) the dissolution of alkaline rocks and minerals in reactors, and (2) 218 electrochemical processes that generate alkalinity by splitting seawater or other brine streams 219 into an acid and base (Eisaman et al., 2023, this Guide). For some materials, such as Ca(OH)₂ 220 and Mg(OH)₂, dissolution slurries are formed and a combination of particulate and aqueous 221 alkalinity can be dosed into seawater. Any particulates that are dosed from the slurry need to 222 dissolve, meaning dissolution kinetics in seawater will be critical. However, the dissolution of 223 these materials tends to be much quicker than with rocks and minerals (Table 1). There are 224 important processes that need to be considered when adding aqueous alkalinity, including the 225 226 unintended precipitation of calcium carbonates due to locally elevated saturation states (Hartmann et al., 2023; Moras et al., 2023). 227

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Field experiments that use aqueous or slurry-based alkalinity additions will need to assess the 229 230 impacts on seawater chemistry at the source of addition and across a dilution radius. Depending on the type of experiment and magnitude of additions this dilution radius could extend upwards 231 of kilometers, but the magnitude of the perturbation to carbonate chemistry would become 232 smaller the further away from the alkalinity source (He & Tyka, 2023). The potential 233 234 environmental impacts from aqueous type alkalinity additions will be similar to those discussed 235 for coastal enhanced weathering, but also include extreme localized changes in carbonate chemistry. 236

238	Table 1.	Types of	falkalinity	sources	and	consideration	is for	each.
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Alkalinity Source	Solid/Aqueous	Dissolution kinetics	Dissolution co- products
NaOH	Aqueous	Instantaneous but can	Alkalinity, Na ⁺

		induce brucite (Mg(OH) ₂) precipitation when NaOH elevates pH >9. Brucite re- dissolves relatively quickly in most cases.	
Manufactured and natural Mg derived alkalinity sources (e.g.,, brucite)	Solid or aqueous slurry	Relatively fast but a combination of dissolution rates both in the receiving and dosing waters.	Alkalinity, limited amounts of nutrients and trace metals (generally less than silicates), Mg ²⁺ .
Silicates (e.g. olivine, basalt, wollastonite)	Solid	Relatively slow dissolution kinetics, but rates are different between silicates.	Alkalinity, silicate, trace metals. Materials need to be individually assessed prior to their use.
Manufactured lime- derived alkalinity sources (e.g. quicklime, ikaite))	Solid or aqueous slurry	Relatively fast but different kinetics between lime products.	Alkalinity, limited amounts of nutrients and trace metals (generally less than silicates), Ca ²⁺ . Materials need to be individually assessed prior to their use.
Iron and steel slag	Solid	Components within steel slag that provide alkalinity (e.g. CaO) dissolve relatively fast but different iron and steel slag contain different amounts.	Alkalinity, Ca ²⁺ , Mg ²⁺ , silicate, phosphate, and trace metals. Materials need to be individually assessed prior to their use.
Natural and synthetic carbonates (e.g. calcite, aragonite)	Solid	They don't dissolve under common surface ocean carbonate chemistry conditions. Dissolution rates can be higher in microenvironments such as corrosive sediment porewaters	Alkalinity, phosphate in some mined sources, dissolved inorganic carbon.

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240 **2.3 Considerations for site selection**

241 Careful consideration should be given to site selection and experimental design to make sure the 242 study adequately address the specific research questions and goals. Some aspects of the field site that will be important include ecosystem- and site-specific characteristics, the prevailing 243 meteorological and oceanographic conditions, and natural spatiotemporal variability. Logistical 244 considerations for site selection include physical access, permitting, availability of electricity, 245 ship time, and consideration of the local community. These considerations will grow with the 246 scale of field experiments and will likely be first-order determinants of where field experiments 247 take place. For example, proximity to a marine institute (for land-based approaches) or access to 248 a research cruise (for open ocean approaches) may be desirable. Logistics will ultimately 249 determine where operational OAE deployments take place and early field experiments will help 250 to elucidate important issues including the impacts of life cycle emissions on CDR. 251

OAE field experimentation requires careful assessment of the field site prior to alkalinity 252 additions to provide foundational knowledge of the site characteristics. Scientific considerations 253 for site selection can be broken down into three categories, the (1) physical, (2) chemical, and (3) 254 biological properties of each site. Important considerations for each category are provided in Box 255 1. To facilitate baseline assessments and site selection we propose Table 2 as guidance for 256 257 relevant parameters to measure. We note that this list is broad, however it is not exhaustive and specific field sites may require the monitoring of different or additional parameters. Furthermore, 258 259 some of the listed parameters may be more applicable to specific OAE approaches. Preliminary knowledge of the field site will inform both the experimental design as well as interpretation of 260 data and experimental outcomes. Due to the large investments in cost and time required to collect 261 baseline data, locations with a wealth of pre-existing scientific data may be considered. This 262 baseline data could be available in the peer-reviewed literature and/or from publicly available 263 264 coastal and open ocean time-series (e.g., Sutton et al., 2019).

Box 1. Scientific considerations for field experiments.

Physics:

- What are the expected dilution rates of the added alkalinity?
- What is the site turbulence and how will this impact alkalinity additions (e.g., keeping particles in suspension)?
- What is the natural light penetration and what impacts could increases in turbidity have on this?
- What is the residence time of water in the surface ocean or mixed layer and how does this relate to the estimated air-sea equilibration time?
- What is driving air-sea gas exchange?
- Will changes in turbidity impact the albedo of the experimental site?
- What is the potential for the lateral export and exchange of alkalinity and other materials?
- Is there the potential for physical disturbance (e.g. impacts of alkalinity additions on physical water mass parameters such as density or the physical impacts of adding undissolved minerals to the benthos)?

• Where will the alkalinity signal be most observable (e.g., pore water vs. water column)?

Chemistry:

- What are the natural carbonate chemistry conditions?
- What modes of variability (e.g., daily, seasonal, interannual) impact seawater chemistry?
- How will variations in seawater chemistry impact signal to noise?
- How will seawater chemistry impact mineral dissolution rates?
- Is there potential to disturb the natural concentrations of macro- or micronutrients or toxic metals through dissolution by-products?
- How do anthropogenic sources of alkalinity interact with (and potentially modify) natural sources and sinks of alkalinity?

Biology:

- What organisms (benthic and pelagic) are present in the study area and what are their relative sensitivities to fluctuations in seawater carbonate chemistry (if known)?
- Are there culturally or commercially important species present?
- Are there endangered or rare species present? Is the site a nursery and/or nesting ground? Are there keystone species and/or important primary producers present? These considerations will likely be part of the permitting process.
- Are there times of the day or seasons with elevated species or ecosystem sensitivities?
- What are the trophic dynamics in the environment, and how might the food web be impacted (e.g., shifts in predator/prey relationships)? What are the cascading implications for the ecosystem as a whole? Might effects be transferred beyond the study site via migratory species?
- Could particulates (e.g., ground rock) cause physical damage prior to dissolution?

- **Table 2.** Parameters that could be considered in assessing sites for OAE field experiments.
- 267 Importantly, some parameters summarized below may require a baseline assessment over
- 268 sufficiently long time frames to cover the intrinsic variability of physical, chemical, and
- 269 biological parameters in the studied system. For example, baseline assessment of marine food
- 270 web structure will likely require a prolonged monitoring effort before (and after) the OAE
- 271 deployment to have a higher chance of detecting OAE-induced effects on marine biota.

Parameter	Rationale	Potential pathway for assessment
Dilution rate	 Exposure risk to alkalinity and mineral dissolution products. Detectability of OAE-induced chemical changes. 	Tracer release experiment (section 2.5).
Turbulence	- Physical energy input to keep ground particles near the sea surface during dissolution.	Microstructure profiler.
Residence time of perturbed patch in surface ocean	- Determination of residence time of an OAE-perturbed patch in the surface to assess whether there is enough time for air-sea equilibration with the atmosphere.	Risk assessment for incomplete air-sea CO ₂ exchange (He and Tyka, 2023; Bach et al., 2023).
Transboundary transport	- Determination of whether there is a high risk for OAE-derived chemicals to be transported into sensitive areas (e.g. marine protected areas, other state territories) in high concentrations. May be useful for residence time as well.	 Tracer release experiment Virtual Lagrangian particle tracking. Utilizing natural tracers observable via remote sensing (e.g., CDOM or Gelbstoff). Mixed layer depth.
Light penetration	- Determination of light environment to assess to what extent the addition of particulate alkalinity source could impact turbidity.	Light loggers, turbidity, CTD casts.
Carbonate chemistry conditions	 Baseline of mean conditions and variability to assess how much change OAE must induce to become detectable. Determination if OAE-related changes are likely to affect marine organisms. 	Dickson et al. (2007) and ocean acidification literature. Schulz et al., (2023, this Guide)

Macronutrients	- Assessment of whether the designated system is prone to macronutrient fertilization via OAE. (Note that not all OAE approaches would introduce macronutrients into the ocean system).	Standard photometric approaches (Hansen and Koroleff, 1999). Experimental assessment of limiting elements.
Micronutrients	- Assessment of whether the designated system is prone to micronutrient fertilization via OAE. (Note that not all OAE approaches would introduce micronutrients into the ocean system).	GEOTRACES cookbook. (https://www.geotraces.org/m ethods-cookbook/) Experimental assessment of limiting elements.
Marine food web structure	- Assessment of the planktonic and/or benthic food web structure prior to testing an OAE deployment.	There is a whole range of surveying tools that could be applied depending on the size and abundance of organisms. Applied methods could range from OMICS (including eDNA), optical observations, acoustics, and flow cytometry.
Risk of damaging organisms by adding ground minerals	- Providing knowledge of whether organisms could be physically harmed, for example through covering them with mineral powder.	Same range of methods as for the food web assessment.
Endangered species	- Clarification if endangered species could be present at the designated field site.	Same range of methods as for the food web assessment. Plane or drone surveys can help to confirm sightings of larger organisms and there may be online resources to be utilized (e.g., WhaleMap). Furthermore, local knowledge should be sought after from the diverse range of stakeholder groups. For example, consultation with indigenous communities, fishermen, local authorities, and environmental agencies.
Foraging/breeding ground	- Clarification if the designated field site is an important	Same range as for endangered species assessments.

	breeding/foraging area for migratory organisms.	
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273 **2.4. Measurement considerations**

274 What to measure and the type of instrumentation needed will ultimately depend on the site, scale, and goals of each individual experiment and should be considered on a case-by-case basis. For 275 example, depending on the alkalinity source utilized (Table 1), it may (e.g., in the case of 276 277 olivine) or may not (e.g., in the case of NaOH) be a priority to measure trace metal or nutrient 278 concentrations. In addition to alkalinity type, the experimental scale will also dictate 279 measurement considerations. For example, if the scale of the perturbation is small or the signal is 280 very dilute, environmental impacts will not likely be measurable far from where the perturbation takes place. If there is a large addition of alkalinity, especially in a semi-enclosed system, both 281 282 environmental impacts and changes in chemistry will be easier to detect. Ultimately, when OAE is done at a larger scale (e.g., millions of moles alkalinity) it is likely that large changes in 283 284 seawater chemistry will wish to be avoided to reduce environmental impacts and avoid secondary precipitation. This presents an interesting challenge to conducting field experiments, 285 286 as the dilution of alkalinity and ultimately CO₂ signal will make MRV more challenging (Ho et al., 2023, this Guide). 287

Seawater carbonate chemistry measurements will be central to most sampling schemes. To cover 288 the appropriate spatial and temporal scales, traditional bottle sampling will likely have to be 289 combined with state of the art in situ sensors (Bushinsky et al., 2019; Briggs et al., 2020; Ho et 290 al., this Guide). Bushinsky et al. (Figure 1; 2019) provides a comprehensive overview of the 291 spatiotemporal capabilities of existing carbonate chemistry sensors and platforms, and care 292 293 should be taken to make sure sensors are appropriate for measurements in seawater. The appropriate methods and protocols for sampling and analysis are outlined in other chapters in this 294 295 guide (Schulz et al., 2023, this Guide) and in the Guide to Best Practices (Dickson et al., 2007). Some general considerations for field experiments include appropriately characterizing the 296 297 natural variability that occurs at the field site through space and time. While total alkalinity titrations should remain a priority, at least two carbonate chemistry parameters (e.g., total 298 299 alkalinity, dissolved inorganic carbon, pH, or pCO₂) should be measured for each sample. It is

- 300 important to note that the combination of pCO₂ and pH is not ideal when calculating CO₂
- 301 chemistry (e.g., using CO2SYS) due to the elevated errors when combining those parameters in
- determining the rest of the carbonate chemistry system in seawater (Lee and Millero, 1995).
- 303 Currently, commercially available autonomous sensors exist for pH and pCO₂, with sensors in
- development for both TA and DIC (Fassbender et al., 2015; Briggs et al., 2020; Qiu et al., 2023).
- 305 While autonomous sensors generally have greater uncertainty than bottle samples coupled with
- 306 laboratory analysis, they will likely play an important role in sampling schemes to help cover
- 307 adequate spatial and temporal resolution in naturally variable marine systems.

While monitoring the background variability and subsequent additions of alkalinity will be 308 309 critical, scientists may also wish to directly measure fluxes of carbon at the field study site (Ho et al., this Guide). The direct measurement of carbon fluxes can be accomplished via different 310 311 methods including benthic and floating chambers, eddy covariance and other benthic boundary layer techniques, and mass balances. These techniques have benefits and drawbacks, including 312 having to enclose the natural system (e.g., chambers) and elevated uncertainty that could be 313 outside of the expected changes due to the perturbation (e.g., eddy covariance). Benthic chamber 314 315 measurements may be particularly important to quantify the dissolution of minerals and rocks 316 added to sediments. Ultimately, any measurements of fluxes due to OAE activities will likely need to be coupled with numerical modeling to estimate the overall drawdown of atmospheric 317 CO_2 (Fennel et al., 2023, this Guide). 318

Field experiments should be informed by other scientific studies as much as possible (e.g., studies based on laboratory experiments, mesocosm studies, natural analogs, and numerical modelling). While not necessarily directly translatable to natural systems (Edmunds et al., 2016; Page et al., 2021), these types of studies can provide first order assessments on safety and efficacy, helping to prevent unintended harmful ecological side effects when conducting large scale perturbations.

Other measurements that may be useful during OAE field experiments are outlined in Table 2. It is important to note that this list is not meant to be exhaustive, and measurement selection will have to be made on a case-by-case basis. Considering the difficulties of tracking water masses in an open system, the next section is a more detailed discussion on tracers for monitoring mixing and dilution of water within the OAE field experiment site. Tracking added alkalinity will be
critical to determine the impacts and efficacy of alkalinity enrichments and may be one of the
biggest challenges facing OAE field experiments.

332 **2.5 Dual tracer regression technique**

If the goal is to track alkalinity additions and measure their effects on carbon fluxes (i.e., net 333 334 ecosystem production or air-sea exchange), a dual tracer regression method can be used (e.g., Albright et al. 2016 & 2018). This approach uses the change in ratios between an active tracer 335 336 (alkalinity) and a passive tracer (dye, artificial gas tracer, Table 3) to assess the fraction of added alkalinity taken up or released by biogeochemical processes in the system. Passive tracers do not 337 338 affect fluid dynamics and are passively advected by the surrounding flow field. The use of passive tracers, such as dye tracers (e.g., rhodamine, fluorescein) or artificial gas tracers (e.g., 339 SF6, CF3SF5) that do not occur in nature helps eliminate background noise. Additional 340 considerations include how many tracers to use and what information each tracer provides (Table 341 342 3).

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During a dual tracer experiment, changes in the active tracer (alkalinity) result from mixing, 344 345 dilution, and biogeochemical activity, whereas changes in the passive tracer are due solely to mixing and dilution. By comparing the alkalinity to dye ratios before (e.g., upstream) and after 346 (e.g., downstream) the water mass interacts with a study area, it is possible to isolate the change 347 in alkalinity that is due to biogeochemical processes such as calcium carbonate precipitation and 348 349 dissolution (Figures 1 & 2). This technique is an extension of Friedlander et al. (1986) and may have applications in other areas of research pertinent to marine CDR, such as nutrient or 350 pollution assessments and the uptake of industrial or agricultural waste. The primary 351 experimental criteria for the dual tracer technique are that the active and passive tracers are 352 added in a fixed ratio and at a fixed rate, in areas where there is a dominant flow direction, 353 dispersion or dilution. 354



Figure 1. Rhodamine dye flowing over a coral reef flat study site during a study in One Tree

358 Island, Australia (Albright et al. 2016). NaOH was used as an active tracer to raise alkalinity, and

359 rhodamine was used as a passive tracer to account for mixing and dilution. Changes in the

360 alkalinity-dye ratios were used to isolate the change in alkalinity flux that was associated with an

361 increase in net community calcification on the reef flat.



Figure 2. Theoretical representations of the null (H0) and alternative (H1) hypotheses for a dual tracer regression experiment where NaOH was used as a source of alkalinity and rhodamine dye

- 365 was used as a passive tracer (from Albright et al. 2016). (a). In H0, the benthic community does
- 366 not take up added alkalinity. Here, the change in alkalinity between the upstream and
- 367 downstream transects would not be systematically related to the dye concentration, and the ratio
- ³⁶⁸ of the alkalinity–dye relationship, r, would not be expected to change between the upstream and
- downstream locations (that is, $r_{up} = r_{down}$). (b). In H1, an uptake of added alkalinity occurs by the
- benthic community. Here, areas with more alkalinity (and more dye) change at a different rate
- than areas with less alkalinity (and less dye), resulting in a change in the alkalinity–dye slope
- 372 (that is, $r_{up} > r_{down}$).
- 373

- **Table 3.** Passive tracers that are available and commonly used for use in field experiments and
- 375 considerations for each. Additional tracers may be useful that are not listed in this table,
- including Helium 3 and Tritium.

Tracer	Туре	Pros	Limitations	Lifespan
Rhodamine	Fluorescent dye	Sensor-based, high frequency (>4 Hz) detection, platform flexibility, detection from space and/or the sky for surface releases.	Optically degrades and absorbs to particles, not good for longer-term studies, not as good signal to noise/detection limits as inert gas tracers.	Several days to weeks
Fluorescein	Fluorescent dye	Sensor-based, high frequency (>4 Hz) detection, platform flexibility, detection from space and/or the sky for surface releases.	Degrades optically - not good for longer-term studies (>24h)	<24 h
SF6	Artificial gas	Inert; capable of being measured at very low concentrations; able to quantify mixing and residence time; good for large- scale ocean tracer release experiments.	Lower frequency detection and less flexibility with platforms, requires discrete measurement. High global warming potential.	years
Trifluorom ethyl sulfur pentafluori	Artificial gas	Good for large- scale ocean experiments.	Difficult to obtain, lower frequency detection and less flexibility with	years

de		autonomous platforms,	
(CF_3SF_5)		requires discrete	
		measurement. High global	
		warming potential.	

2.6 Detecting change and the importance of controlled experiments

379

Separating an experimental 'signal' from the background 'noise' inherent in natural systems can 380 be challenging, especially in field experiments where replication may not be practical (Carpenter, 381 1990). Gaining baseline knowledge on the physical, chemical, and biological components of the 382 study site should be a priority. There is often considerable natural variability in marine systems, 383 and especially in coastal systems, due to fluctuations in biological activity, hydrodynamics, 384 seasonal and/or interannual influences, and others (Bates et al., 1998; Bates 2002; Hagens and 385 Middelburg, 2016; Landschützer et al., 2018; Sutton et al., 2019; Kapsenberg and Cyronak, 386 2019; Torres et al., 2021). Fully characterizing this variability could take many years, which may 387 388 create significant barriers to experimental progress in the field. Therefore, we recommend that any potential modes of spatiotemporal variability be recognized and evaluated while planning 389 390 field experiments. For instance, in coastal systems with river and groundwater inputs it will be important to know the impact that freshwater has on carbonate chemistry. 391 392

Where possible, conducting controlled experiments will help to maximize the ratio of signal to 393 394 noise, thereby improving statistical power to detect experimental effects. The pros and cons of replicating experimental controls in space versus time should be taken into consideration. For 395 396 many field experiments (and natural analogs; see Subhas et al., 2023, this Guide), sample size will be inherently limited (e.g., one, or few study sites); therefore, conducting controls in time 397 (e.g., every third day) may be the best option. For studies with limited (or no) replication, there 398 are statistical methods that can be used to isolate effects pre- and post-treatment (Carpenter, 399 1990). Numerical simulations and machine learning based network design are potentially 400 401 valuable tools to optimize observational networks to detect experimental change.

402

403 **3. Additional considerations**

Permitting. Addressing regulatory requirements is critical prior to conducting field experiments. 404 The spatial and temporal scale of the field trial, as well as the specific considerations of the 405 deployment site (e.g., protection status) will determine permitting requirements. Engaging with 406 this process early is advised - for example, understanding who the permit-granting authorities are 407 for a given area and timelines for associated regulatory processes. In some cases, the use of 408 existing infrastructure (e.g., wastewater discharge sites) and environmental projects (e.g., beach 409 renourishment) may offer ways to streamline experiments, although permitting will be governed 410 by existing regulations. For a detailed discussion on legal considerations, see Steenkamp and 411 Webb (2023, this Guide). 412

413 Community engagement and social considerations of field experiments. The likelihood of harmful ecological consequences from OAE field experiments remains unclear and will 414 ultimately depend on the technology and temporal and spatial scale of the experiment. Field 415 experiments evaluating CDR approaches carry the risk of unintended consequences and impacts 416 over large spatial scales, so appropriate scaling (e.g., starting small) is necessary (NASEM, 417 2022). In response to these unknowns, researchers should follow the key components for a code 418 419 of conduct for marine CDR research, e.g., as outlined by Loomis et al. (2022), which details best 420 practices that encourage responsible research amongst both the public and private sectors. 421

Social license to operate is critical for the success of CDR projects and researchers have an 422 obligation to involve the full community of people (public and stakeholders) who may be 423 424 impacted by the research (Nawaz et al., 2022; Cooley et al., 2023). Therefore, public outreach is important both before and during field experimentation. The study site will determine the 425 potential for community engagement. Coordinating with local and/or regional organizations who 426 are connected to relevant stakeholders (for example, your local SeaGrant office if in the United 427 States) will be helpful. For additional discussion on social considerations of OAE field trials, see 428 Satterfield et al. (2023, this Guide). 429

430 Collaboration and data/information sharing. Considering the inherent challenges to OAE field 431 experiments (cost, permitting, access, logistics, environmental safety), fostering interdisciplinary 432 and collaborative teams will help ensure the greatest return on investment. Examples of ways to 433 foster collaboration include, developing test-bed field sites that are open to participation from

diverse groups, making efforts to include groups who may not traditionally have access to and/or 434 the capacity for field campaigns, and including travel support in grant applications to support 435 external collaborators. Making concerted efforts to share information, resources, and ideas will 436 allow researchers to combine knowledge and resources in ways that might not have been 437 possible when working alone, thereby advancing OAE technology and science at a faster pace. 438 When publishing in peer-reviewed literature, uploading data to publicly available data 439 repositories and publishing in open access journals following best practices should be prioritized 440 (Jiang et al., 2023, this Guide). 441

Inclusivity and transparency during OAE field trials are crucial to ensure that knowledge gained 442 443 is fed back into scientific and other communities efficiently, iteratively informing and refining the next generation of experiments. Some field experiments will mimic plans for real world OAE 444 deployments and should therefore be done in collaboration with relevant stakeholders across 445 science, industry, policy, and communities. To foster collaboration and technology transfer, we 446 advocate for a centralized platform and/or organization to share data and information in this 447 rapidly evolving field. This might look like a centralized, freely accessible platform for early 448 449 and/or 'real-time' information sharing (i.e., before publication) that can facilitate faster information exchange within the research community (e.g., data sharing, permitting issues). Two 450 existing options that could help fill this gap are the OA information exchange 451 (https://www.oainfoexchange.org/index.html) and the Ocean Visions community 452 (https://community.oceanvisions.org/dashboard). It may prove useful to designate core working 453 groups of experts in various aspects of CDR that investigate specific needs and priorities and 454 work to synthesize and share existing knowledge in the context of field experiments. This 455 approach has been adopted by other scientific disciplines in high priority, rapidly evolving, and 456 highly collaborative fields, greatly benefiting the scientific community at large (e.g., the Coral 457 Restoration Consortium, https://www.crc.world/ - and associated working groups). Coordinating 458 field trials with research groups conducting laboratory and mesocosm experiments, studying 459 natural analogs, and undertaking modeling efforts will help strengthen the interpretation and 460 extrapolation of results. 461

462 **4.1 Conclusion and Recommendations**

Given that few OAE field studies have been conducted to date, there is much to learn from the 463 earliest experiments with respect to experimental design, measurement and monitoring, 464 deployment considerations, environmental impact, and more. Early experiments will only engage 465 with a fraction of the temporal and spatial scales involved in full-scale operational OAE, and 466 longer-term and larger-scale studies will become increasingly important to reveal scale-467 dependencies as the field develops. It is important that marine CDR research is hypothesis-468 driven, structured, deliberate, and well-planned to best inform future decision-making about 469 OAE techniques and deployments. Careful consideration of the physical, chemical, and 470 biological components of the study area will help inform the experimental approach. The use of 471 baseline studies (both previous and contemporary to the OAE deployment) and controls will help 472 to maximize signal-to-noise ratios and identify experimental effects. The timescale of OAE field 473 experiments should not be underestimated, especially when considering permitting and the data 474 needed to capture the baseline variability in natural systems. 475

Considering the urgent timeline required for humanity to meet our climate goals, field
experiments need to move forward swiftly yet deliberately. To ensure the success of OAE,
diverse perspectives from research, industry, policy, and society must converge, demanding
transdisciplinary thinking and a commitment to open and transparent science. Central to this
ambitious undertaking are the early field experiments, results from which will ultimately
determine the successes and failures of OAE projects and technologies.

482 4.2 Key recommendations for field experiments relevant to the research of ocean alkalinity 483 enhancement

1. Ensure inclusivity and transparency (community engagement, data sharing, etc) for 484 OAE field experiments to both advance the field as quickly as possible, and to ensure 485 the field progresses in a socially responsible manner. 486 2. Assess the potential risks and benefits for any perturbation. Proceed according to the 487 code of conduct and precautionary principles. 488 3. Develop methods to track signal versus noise in highly variable environments, 489 including robust baseline studies to characterize underlying variability (biological, 490 chemical, physical), and the inclusion of controlled experiments such as chamber 491

492	incubations to isolate treatment effects.
493	4. Consider logistical constraints and opportunities.
494	
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496	The contact author has declared that none of the authors has any competing interests.
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498	LTB is scientific advisor to Submarine, a start-up service provider for monitoring, reporting, and
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501	
502	
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513 **References**

- Albright, R., Caldeira, L., Hosfelt, J., Kwiatkowski, L., Maclaren, J. K., Mason, B. M., ... &
- Caldeira, K. (2016), Reversal of ocean acidification enhances net coral reef calcification. Nature,
 531(7594), 362-365.
- 517 Albright, R., Takeshita, Y., Koweek, D. A., Ninokawa, A., Wolfe, K., Rivlin, T., ... & Caldeira,
- 518 K. (2018), Carbon dioxide addition to coral reef waters suppresses net community calcification.
- 519 Nature, 555(7697), 516-519.
- 520 Bach, L. T., Gill, S. J., Rickaby, R. E., Gore, S., & Renforth, P. (2019), CO₂ removal with
- 521 enhanced weathering and ocean alkalinity enhancement: potential risks and co-benefits for
- 522 marine pelagic ecosystems. Frontiers in Climate, 1, 7.
- 523 Bach, L.T., Ho, D.T., Boyd, P.W. & Tyka, M.D. (2023), Toward a consensus framework to
- ⁵²⁴ evaluate air–sea CO2 equilibration for marine CO2 removal. Limnology & Oceanography
- 525 Letters https://doi.org/10.1002/lol2.10330.
- 526 Bates, N. R. (2002), Seasonal variability of the effect of coral reefs on seawater CO2 and air—
- sea CO₂ exchange, Limnology and Oceanography, 47, 43–52.
- 528 Bates, N. R., Takahashi, T., Chipman, D. W., & Knap, A. H. (1998), Variability of pCO₂ on diel
- to seasonal timescales in the Sargasso Sea near Bermuda, Journal of Geophysical Research:
- 530 Oceans, 103, 15567–15585.
- 531 Briggs, E. M., De Carlo, E. H., Sabine, C. L., Howins, N. M., & Martz, T. R. (2020),
- 532 Autonomous ion-sensitive field effect transistor-based total alkalinity and pH measurements on a
- barrier reef of Kane'ohe Bay. ACS Earth and Space Chemistry, 4(3), 355-362.
- 534 Bushinsky, S. M., Takeshita, Y., & Williams, N. L. (2019), Observing changes in ocean
- carbonate chemistry: our autonomous future. Current climate change reports, 5, 207-220.
- Carpenter, S. R. (1990), Large-scale perturbations: opportunities for innovation. Ecology, 71(6),
 2038-2043.
- 538 Cooley, S. R., Klinsky, S., Morrow, D. R., & Satterfield, T. (2023), Sociotechnical

- considerations about ocean carbon dioxide removal. Annual Review of Marine Science, 15, 41-66.
- 541 Dickson, A. G., Sabine, C. L., & Christian, J. R. (2007), Guide to best practices for ocean CO2
 542 measurements. North Pacific Marine Science Organization.
- 543 Edmunds, P. J., Comeau, S., Lantz, C., Andersson, A., Briggs, C., Cohen, A., ... & Carpenter, R.
- 544 C. (2016), Integrating the effects of ocean acidification across functional scales on tropical coral
- 545 reefs. Bioscience, 66(5), 350-362.
- 546 Egleston, E. S., Sabine, C. L., & Morel, F. M. M. (2010), Revelle revisited: Buffer factors that
- 547 quantify the response of ocean chemistry to changes in DIC and alkalinity, Global
- 548 Biogeochemical Cycles, 24.
- Eisaman, M., Geilert, S., Renforth, P., Bastianini, L., Campbell, J., Dale, A., Foteinis, S., Grasse,
- 550 P., Hawrot, O., Löscher, C., Rau, G., and Rønning, J. (2023), Assessing technical aspects of
- ocean alkalinity enhancement approaches, In: Guide to Best Practices in Ocean Alkalinity
- 552 Enhancement Research (OAE Guide 23), edited by: Oschlies, A., Stevenson, A., Bach, L.,
- 553 Fennel, K., Rickaby, R., Satterfield, T., Webb, R., and Gattuso, J.-P., Copernicus Publications,
- 554 State Planet.
- 555 Fassbender, A. J., Sabine, C. L., Lawrence-Slavas, N., De Carlo, E. H., Meinig, C., & Maenner
- Jones, S. (2015), Robust sensor for extended autonomous measurements of surface ocean
- dissolved inorganic carbon. Environmental Science & Technology, 49(6), 3628-3635.
- 558 Fennel, K., Long, M. C., Algar, C., Carter, B., Keller, D., Laurent, A., Mattern, J. P., Musgrave,
- 859 R., Oschlies, A., Ostiguy, J., Palter, J., and Whitt, D. B (2023), Modeling considerations for
- research on Ocean Alkalinity Enhancement (OAE), In: Guide to Best Practices in Ocean
- Alkalinity Enhancement Research (OAE Guide 23), edited by: Oschlies, A., Stevenson, A.,
- 562 Bach, L., Fennel, K., Rickaby, R., Satterfield, T., Webb, R., and Gattuso, J.-P., Copernicus
- 563 Publications, State Planet.
- 564 Flipkens, G., Fuhr, M., Fiers, G., Meysman, F. J. R., Town, R. M., & Blust, R. (2023), Enhanced
- 565 olivine dissolution in seawater through continuous grain collisions, Geochimica et

- 566 Cosmochimica Acta, 359, 84-99.
- 567 Foteinis, S., Campbell, J. S., and Renforth P. (2023), Life Cycle Assessment of Coastal
- 568 Enhanced Weathering for Carbon Dioxide Removal from Air. Environmental Science &

569 Technology, 57(15), 6169-6178

- 570 Friedlander, S.K., Turner, J.R. & Hering, S.V. (1986), A new method for estimating dry
- deposition velocities for atmospheric aerosols. J. Aerosol Sci. 17, 240–244.
- 572 Fuhr, M., Geilert, S., Schmidt, M., Liebetrau, V., Vogt, C., Ledwig, B., & Wallmann, K. (2022),
- 573 Kinetics of Olivine Weathering in Seawater: An Experimental Study, Frontiers in Climate, 4.
- 574 Hagens, M. & Middelburg, J. J. (2016), Attributing seasonal pH variability in surface ocean
- waters to governing factors, Geophysical Research Letters, 43, 2016GL071719.
- Hansen, H. P., & Koroleff, F. (1999), Determination of nutrients. Methods of seawater analysis,
 159-228.
- 578 Hartmann, J., Suitner, N., Lim, C., Schneider, J., Marín-Samper, L., Arístegui, J., ... & Riebesell,
- 579 U. (2022), Stability of alkalinity in Ocean Alkalinity Enhancement (OAE) approaches-
- 580 consequences for durability of CO₂ storage. Biogeosciences, 1-29.
- Hauck, J., Köhler, P., Wolf-Gladrow, D., & Völker, C. (2016), Iron fertilisation and century-
- scale effects of open ocean dissolution of olivine in a simulated CO₂ removal experiment,
- 583 Environmental Research Letters, 11, 024007.
- He, J., & Tyka, M. D. (2023), Limits and CO2 equilibration of near-coast alkalinity
 enhancement. Biogeosciences, 20(1), 1726-4189.
- 586 Ho, D. T., Bopp, L., Palter, J. B., Long, M. C., Boyd, P., Neukermans, G., & Bach, L. (2023),
- 587 Monitoring, Reporting, and Verification for Ocean Alkalinity Enhancement, In: Guide to Best
- 588 Practices in Ocean Alkalinity Enhancement Research (OAE Guide 23), edited by: Oschlies, A.,
- 589 Stevenson, A., Bach, L., Fennel, K., Rickaby, R., Satterfield, T., Webb, R., and Gattuso, J.-P.,
- 590 Copernicus Publications, State Planet.

- Jiang, L.-Q., Subhas, A., Basso, D., Fennel, K., and Gattuso, J.-P. (2023), Data reporting and
- 592 sharing for ocean alkalinity enhancement research, In: Guide to Best Practices in Ocean
- 593 Alkalinity Enhancement Research (OAE Guide 23), edited by: Oschlies, A., Stevenson, A.,
- Bach, L., Fennel, K., Rickaby, R., Satterfield, T., Webb, R., and Gattuso, J.-P., Copernicus
- 595 Publications, State Planet.
- 596 Kapsenberg, L., & Cyronak, T. (2019), Ocean acidification refugia in variable environments.
- 597 Global Change Biology, 25(10), 3201-3214.
- Landschützer, P., Gruber, N., Bakker, D. C. E., Stemmler, I., & Six, K. D. (2018), Strengthening
- seasonal marine CO₂ variations due to increasing atmospheric CO₂, Nature Climate Change, 8,
 146–150.
- Lee, K., & Millero, F. J. (1995), Thermodynamic studies of the carbonate system in seawater.
- Deep Sea Research Part I: Oceanographic Research Papers, 42(11-12), 2035-2061.
- Loomis, R., Cooley, S. R., Collins, J. R., Engler, S., & Suatoni, L. (2022), A Code of Conduct Is
 Imperative for Ocean Carbon Dioxide Removal Research. Frontiers in Marine Science, 9.
- Montserrat, F., Renforth, P., Hartmann, J., Leermakers, M., Knops, P., & Meysman, F. J. R.
- 606 (2017), Olivine Dissolution in Seawater: Implications for CO2 Sequestration through Enhanced
- Weathering in Coastal Environments, Environmental Science & Technology, 51, 3960-3972.
- Moras, C. A., Bach, L. T., Cyronak, T., Joannes-Boyau, R., & Schulz, K. G. (2022), Ocean
- alkalinity enhancement–avoiding runaway CaCO₃ precipitation during quick and hydrated lime
- dissolution. Biogeosciences, 19(15), 3537-3557.
- Morse, J. W., Arvidson, R. S., & Lüttge, A. (2007), Calcium carbonate formation and
 dissolution, Chemical Reviews, 107, 342-381.
- Nawaz, S., Peterson St-Laurent, G., & Satterfield, T. (2023), Public evaluations of four
- approaches to ocean-based carbon dioxide removal. Climate Policy, 23(3), 379-394.
- National Academies of Sciences, Engineering, and Medicine. (2022), A Research Strategy for

- 616 Ocean-based Carbon Dioxide Removal and Sequestration. Washington, DC: The National
- 617 Academies Press. doi.org/10.17226/26278.
- 618
- 619 Page, H. N., Bahr, K. D., Cyronak, T., Jewett, E. B., Johnson, M. D., & McCoy, S. J. (2022),
- 620 Responses of benthic calcifying algae to ocean acidification differ between laboratory and field
- settings. ICES Journal of Marine Science, 79(1), 1-11.
- 622 Qiu, L., Li, Q., Yuan, D., Chen, J., Xie, J., Jiang, K., Guo, L., Zhong, G., Yang, B., &
- Achterberg, E. P. (2023), High-Precision In Situ Total Alkalinity Analyzer Capable of Month-
- Long Observations in Seawaters, ACS Sensors, 8, 2702-2712.
- Renforth, P., & Henderson, G. (2017), Assessing ocean alkalinity for carbon sequestration,
- 626 Reviews of Geophysics, 55, 636-674.
- Rimstidt, J. D., Brantley, S. L., & Olsen, A. A. 2012. Systematic review of forsterite dissolution
 rate data, Geochimica et Cosmochimica Acta, 99, 159-178.
- 629 Satterfield, T., Nawaz, S., and Boettcher, M. (2023), Social Considerations and Best Practices for
- 630 Engaging Publics on Ocean Alkalinity Enhancement, In: Guide to Best Practices in Ocean
- Alkalinity Enhancement Research (OAE Guide 23), edited by: Oschlies, A., Stevenson, A.,
- Bach, L., Fennel, K., Rickaby, R., Satterfield, T., Webb, R., and Gattuso, J.-P., Copernicus
- 633 Publications, State Planet.
- 634 Schulz, K. G., Bach, L. T., and Dickson, A. G. (2023), Seawater carbonate system considerations
- 635 for ocean alkalinity enhancement research, , In: Guide to Best Practices in Ocean Alkalinity
- Enhancement Research (OAE Guide 23), edited by: Oschlies, A., Stevenson, A., Bach, L.,
- 637 Fennel, K., Rickaby, R., Satterfield, T., Webb, R., and Gattuso, J.-P., Copernicus Publications,
- 638 State Planet.
- 639 Steenkamp, R. C. and Webb, R. M. (2023), Legal Considerations relevant to the Research of
- 640 Ocean Alkalinity Enhancement, In: Guide to Best Practices in Ocean Alkalinity Enhancement
- Research (OAE Guide 23), edited by: Oschlies, A., Stevenson, A., Bach, L., Fennel, K., Rickaby,
- R., Satterfield, T., Webb, R., and Gattuso, J.-P., Copernicus Publications, State Planet.
- 643 Subhas, A. V., Lehmann, N., and Rickaby, R. (2023), Natural Analogs to Ocean Alkalinity
- 644 Enhancement, In: Guide to Best Practices in Ocean Alkalinity Enhancement Research (OAE
- Guide 23), edited by: Oschlies, A., Stevenson, A., Bach, L., Fennel, K., Rickaby, R., Satterfield,
- T., Webb, R., and Gattuso, J.-P., Copernicus Publications, State Planet.
- 647 Sutton, A. J., Feely, R. A., Maenner-Jones, S., Musielwicz, S., Osborne, J., Dietrich, C.,

- 648 Monacci, N., Cross, J., Bott, R., Kozyr, A., Andersson, A. J., Bates, N. R., Cai, W.-J., Cronin, M.
- 649 F., De Carlo, E. H., Hales, B., Howden, S. D., Lee, C. M., Manzello, D. P., McPhaden, M. J.,
- 650 Meléndez, M., Mickett, J. B., Newton, J. A., Noakes, S. E., Noh, J. H., Olafsdottir, S. R.,
- 651 Salisbury, J. E., Send, U., Trull, T. W., Vandemark, D. C., & Weller, R. A. (2019), Autonomous
- 652 seawater pCO2 and pH time series from 40 surface buoys and the emergence of anthropogenic
- trends, Earth System Science Data, 11, 421–439.
- Torres, O., Kwiatkowski, L., Sutton, A. J., Dorey, N., and Orr, J. C. (2021), Characterizing Mean
- and Extreme Diurnal Variability of Ocean CO₂ System Variables Across Marine Environments,
- 656 Geophysical Research Letters, 48.