

Dear Reviewers,

Thank you for your very thoughtful review of our paper. We provide our responses (Plain text with black and blue colors) to all comments (*italic text*) below. We hope that the revision addresses your concerns.

- Response to Reviewer Comments -

Reviewer #1

-Specific Comments:

1. Most importantly, the authors are still only giving vague directions as to the depths at which particle export needs to be measured. These measurements, however, are absolutely crucial if we are to judge the effectiveness of iron fertilisation as a means of carbon sequestration, and there has been substantial debate in the particle flux literature precisely about this point. Unfortunately, the authors only talk vagues of “intermediate and deep waters” (e.g. Page 4 Line 1, P 26 L 28, and P 27 L 6; clear directions on depths are also lacking in Section 3.1). Measurement depths for particle flux absolutely need to be spelled out explicitly and as much as possible standardised in future experiments so that meaningful comparisons can be made. In the abstract, and on P 29 L 18, the authors indicate that in KIFES, they would want to place one sediment trap “within the mixed layer and another below it”. This is absolutely not appropriate: sediment traps are known to perform poorly within the mixed layer, so the shallowest depth for a trap should be just below the mixed layer, so that mixed layer export can be measured. The second critical depth for a sediment trap is the depth of deepest winter-time convective mixing, which can be estimated from Argo float data for a given study area. In LOHAFEX, that was how the depth of the deep sediment traps at around 400 m was chosen. My recommendation would be that the authors make clear recommendations that particle flux must be measured just below the mixed layer (i.e. 10–20 m below) and at the depth of the winter mixed layer. I am open to alternative suggestions from the authors, but whatever they intend to do absolutely must be spelled out clearly, and the importance of these depths should be explained properly in the manuscript. As it is described at present, I would have real concerns as to whether KIFES would actually collect the correct measurements. It is critical that this manuscript makes appropriate, and explicit, recommendations in this regard.

- Thank you for pointing out this deficiency. We have modified ‘**intermediate and deep waters**’ to ‘**below the winter mixed layer**’. We have added sentences explaining ‘*specific depth for particle flux measurement*’, ‘*the importance of these depths*’, and defining ‘*effectiveness of aOIF*’.

→ Abstract (lines 21–23, page 1):

“However, except in the Southern Ocean European Iron Fertilization Experiment, **the effectiveness of aOIF (i.e., the amount of iron-induced carbon export flux below the winter mixed layer depth)** has been unexpectedly low compared to that achieved by natural phytoplankton blooms.”

→ Abstract (lines 33–36, page 1):

“(4) Employment of neutrally buoyant sediment traps and application of the water-column derived ²³⁴Thorium method at two depths (**i.e., just below the *in situ* mixed layer depth and at the winter mixed layer depth**), with autonomous profilers equipped with an underwater video profiler and a transmissometer.”

→ 1 Introduction (lines 1–4, page 4):

“However, for aOIF to be considered as a useful geoengineering approach (IPCC, 2007), in the long run, the most critical issue is the ‘**effectiveness of aOIF**’. That is, whether a significant portion of the organic carbon produced by aOIF in the surface waters **is exported below the winter mixed layer depth (MLD)** to intermediate/deep layers for long-term (~1,000 years) storage (Fig. 3c) (Lampitt et al., 2008).”

→ 2.1.2 Southern Ocean (lines 2–4, page 6):

“These early aOIF experiments resulted in clear increases in phytoplankton biomass and PP, but the impact on export production (i.e., carbon export from the surface waters **to below the winter MLD**) was not evident (Fig. 3c) (de Baar et al., 2005; Boyd et al., 2007).”

→ 2.4.2 Southern Ocean (lines 2–3, page 13):

“However, it was unclear whether surface-fixed carbon was well and truly delivered **below the winter MLD**.”

→ 2.5 Summary of the significant results from aOIF experiments (lines 27–30, page 15):

“It should also be noted that the rates of bacterial remineralization and grazing pressure on the diatoms were in the same range inside the fertilized patch as outside, which might have assisted the delivery of iron-induced POC **from the ML to deep layers** (Smetacek et al., 2012). These results suggest that to detect **significant carbon exported below the winter MLD** following an increase in PP, at least three conditions are necessary.”

→ 3 Present: Unanswered aOIF questions (line 39, page 15 – line 1, page 16):

“However, the impact on the net transfer of CO₂ from the atmosphere to **below the winter MLD** through the ‘biological pump’ (Fig. 3c) is not yet fully understood or quantified and appears to vary with environmental conditions, export flux measurement techniques, and other unknown factors (Smetacek et al., 2012).”

→ 3.1 Export flux measurement methods (lines 35–37, page 16):

“The application of sediment traps for the determination of the carbon export flux is relatively more biased in the **ML** where ocean currents are generally faster and zooplankton are much more active than deep water. These issues suggest that sediment traps alone may not accurately determine carbon export fluxes within the **ML**.”

→ 3.1 Export flux measurement methods (lines 22–24, page 17):

“LOHAFEX has been the only aOIF experiment so far that has measured particle export using PELAGRA (Particle Export measurement using a LAGRAngian trap) sediment traps based on the NBST system deployed **at two depths of 200 m and 450 m (below the winter MLD)** (Martin et al., 2013).”

→ 3.1 Export flux measurement methods (lines 11–12, page 18):

“NBST systems (e.g., the PELAGRA sediment trap) should be deployed **at two depths (i.e., below both the *in situ* MLD and the winter MLD)** to quantify the aOIF-induced POC flux.”

→ 4 Future: Designing future aOIF experiments (lines 28–31, page 23):

“Eddy centers tend to be subject to relatively slow current speeds, with low shear and high vertical coherence, providing ideal conditions for tracing the same water from the surface **to below the winter MLD**, while simultaneously minimizing lateral stirring and advection (Smetacek et al, 2012).”

→ 4 Future: Designing future aOIF experiments (lines 25–27, page 24):

“Based on previous aOIF experiments, it would, therefore, be important to detect the full phase of a phytoplankton bloom to determine accurately the amount of iron-induced POC exported **out of the winter ML**.”

→ 4 Future: Designing future aOIF experiments (lines 38–41, page 25):

“Direct measurements of carbon export fluxes to determine the effectiveness of aOIF should be conducted by deploying an NBST at two depths: **(1) just below the *in situ* MLD** to detect increases in iron-induced POC in the surface layer along with the calibration of the water-column based ^{234}Th method, and **(2) at the winter MLD to detect iron-induced carbon export fluxes below winter MLD.**”

→ 4 Future: Designing future aOIF experiments (lines 30–33, page 26):

“(4) NBST system and water-column derived ^{234}Th method are employed at two depths **(i.e., just below the *in situ* MLD and at the winter MLD)**, with autonomous profilers equipped with an UVP and a transmissometer to estimate accurately the carbon export flux.”

→ 5.1 Background - Bransfield Basin (lines 8–9, page 27):

“(2) To evaluate the effectiveness of scientific aOIF in terms of atmospheric carbon sequestration (i.e., to identify/quantify significant increases in iron-induced carbon export fluxes **below the winter MLD**) in the SO.”

→ 5.2.4 Year four plan (lines 18–21, page 29):

“(5) Measure iron-induced carbon export fluxes for the regions both inside and outside the center of an eddy structure using NBST systems at two depths **(i.e., just below the *in situ* MLD and at the winter MLD)** along with the calibration of water-column based ^{234}Th measurements and autonomous profilers equipped with a transmissometer and an UVP.”

2. *The authors in several places talk about “effectiveness” of aOIF, but I don’t remember that being properly defined in the manuscript. As above, I think this does need to be defined, and in my view an appropriate definition would be the amount of additional carbon exported below the winter mixed layer depth as a result of iron addition. The importance of other metrics, such as the amount of carbon sequestered relative to the amount of iron added, might also be mentioned in this context.*

- We added the definition about ‘effectiveness of aOIF’. Please find the [Abstract \(lines 22–23, page 1\)](#) and [1 Introduction \(lines 1–4, page 4\)](#).

3. In the abstract, Line 27, the authors might want to briefly spell out what these questions are. The abstract can probably be shortened by editing the language carefully throughout. I would also recommend that the depths at which carbon sequestration should be monitored are explicitly spelled out in the abstract (and, I reiterate, these depths are not inside the mixed layer and [somewhere] below the mixed layer).

- We have briefly spelled out ‘what these questions are’, shortened the ‘abstract’.

→ Abstract (lines 15–40, page 1):

“Since the start of the industrial revolution, human activities have caused a rapid increase in atmospheric CO₂ concentrations, which have, in turn, had an impact on climate leading to global warming and ocean acidification. Various approaches have been proposed to reduce atmospheric CO₂. The 'Martin (or Iron) Hypothesis' suggests that ocean iron fertilization (OIF) could be an effective method for stimulating oceanic carbon sequestration through the biological pump in iron-limited, high-nutrient, low-chlorophyll (HNLC) regions. To test the Martin hypothesis, 13 artificial OIF (aOIF) experiments have been performed since 1990 in HNLC regions. These aOIF field experiments have demonstrated that primary production can be significantly enhanced by the artificial addition of iron. However, **except in the Southern Ocean European Iron Fertilization Experiment, the effectiveness of aOIF (i.e., the amount of iron-induced carbon export flux below the winter mixed layer depth) has been** unexpectedly low compared to that achieved by natural phytoplankton blooms. These results, including possible side effects have been debated amongst those who support and oppose aOIF experimentation, and many questions **such as effectiveness of scientific aOIF, environmental side effects, and international aOIF law frameworks** remain. In the context of increasing global and political concerns associated with climate change, it is valuable to examine the validity and usefulness of the aOIF experiments. To maximize the effectiveness of aOIF experiments under international aOIF regulations in the future, we suggest a design that incorporates several **components**. (1) Experiments conducted in the center of an eddy structure when grazing pressure is low and silicate levels are high (e.g., in the Southern Ocean south of polar front during early summer). (2) Shipboard observations extending over a minimum of ~40 days, with multiple iron injections (at least 2 (or 3) iron infusions of ~2,000 kg with an interval of ~10–15 days to fertilize a patch of 300 km² and obtain a ~2 nM concentration). (3) Tracing of the iron fertilized patch using both physical (e.g., a drifting buoy) and biogeochemical (e.g., sulfur hexafluoride and photosynthetic quantum efficiency) tracers. **(4) Employment of neutrally buoyant sediment traps and application of the** water-column derived ²³⁴Thorium method at two depths **(i.e., just below the *in situ* mixed layer depth and at the winter mixed layer depth)**, with autonomous profilers equipped with an underwater video profiler and a transmissometer. (5) Monitoring of side effects on marine/ocean ecosystems, including production of climate-relevant gases (e.g., N₂O, dimethyl sulfide, and halogenated volatile organic compounds), **decline in oxygen inventory**, and development of toxic algae blooms, **with optical sensor equipped**

autonomous moored profilers and/or autonomous benthic vehicles. Lastly, we introduce the scientific aOIF experimental design guidelines for a future Korean Iron Fertilization Experiment in the Southern Ocean.”

4. *In the first paragraph of the introduction, I am quite strongly of the opinion that the authors should also state that there is an urgent need to reduce global greenhouse gas emissions.*

- We have added statement.

→ 1 Introduction (lines 2–5, page 3):

“Since the start of the industrial revolution, human activities have caused a rapid increase in atmospheric carbon dioxide (CO₂, **a major greenhouse gas**) from ~280 ppm (pre-industrial revolution) to ~400 ppm (present day) (<http://www.esrl.noaa.gov/>), which has, in turn, led to global warming and ocean acidification, **indicating that there is an urgent need to reduce global greenhouse gas emissions** (IPCC, 2013) (Fig. 1).”

5. *P 3 L 12: why is “ocean fertilisation” italicised?*

- We changed to plain text.

6. *P 6 L 10: Presumably you mean “By the end of the 20th century”?*

- We apologize this confusing. Based on **Reviewer #2’s editing**, we have removed this sentence.

7. *P 10 L 26: Correct spelling of the name is “Behrenfeld”*

- We corrected it.

8. *P 16 L35: What is the difference between “tracking” and “quantifying” export flux? I think one will do. Again, please do be explicit throughout the manuscript as to what depths you are thinking about.*

- We modified “through both **tracking and quantifying export flux**” to “through **quantification of export fluxes**”. For ‘specific depths’, please refer **Reviewer #1’s Response 1**.

9. P 18 L 12: *Saying that thorium and sediment traps are “of limited use” in determining the fate of POC is not appropriate; the implication of saying “of limited use” is that these measurements are not very useful. The authors should re-phrase this to say “and, therefore, these methods should ideally be complemented with additional techniques that can measure particle stocks at high depth resolution throughout the water column”.*

- Reviewer is correct. We have revised this sentence.

10. P 18 L 17: *The UVP, and also transmissometer and other optical measurements, actually do not give a particle flux as such, they show the particle concentration. There are ways to estimate fluxes from these measurements using assumptions about particle sinking rates, but in the first instance they provide information about stocks. The authors might also want to refer to the study by Briggs et al. (2011) High-resolution observations of aggregate flux during a sub-polar North Atlantic spring bloom in Deep-Sea Research 1. This paper provides a very nice example of using backscattering and chlorophyll sensors to gain high-resolution data about a particle flux event in a Lagrangian study, so a similar scenario to an iron fertilisation.*

- Thank you for pointing out this deficiency and recommending nice study. We have modified ‘particle flux’ to ‘stock’ or ‘concentration’ and have cited Briggs et al. (2011).

➔ 3.1 Export flux measurement methods (lines 34–36, page 17):

“Total vertical particle volume profiles obtained from the UVP indicated **a maximum concentration** at 75 m ($\sim 0.3 \text{ mm}^3 \text{ L}^{-1}$), with a gradual decrease to 150 m ($\sim 0.15 \text{ mm}^3 \text{ L}^{-1}$).”

➔ 3.1 Export flux measurement methods (line 39, page 17 – line 2, page 18):

“To continuously monitor vertical changes in POC **stocks** following iron addition, EIFEX used a transmissometer, providing high vertical resolution (~ 24 data points per meter) and tracking of the iron-induced **stocks** down to $\sim 3,000$ m, even though, unlike UVPs, transmissometers do not allow classification of particles (Smetacek et al., 2012).”

➔ 3.1 Export flux measurement methods (lines 7–9, page 18):

“Furthermore, recent studies also reported that use of optical spike signals in particulate backscattering and fluorescence, measured from autonomous platforms such as gliders and floats, can provide high-resolution observations of POC flux (Briggs et al., 2011; Dall’Olmo and Mork, 2014).”

11. P 20 L 10: *This sentence isn't quite clear. What does the 20% refer to: a 20% increase of the total SO DMS flux? Or a 20% increase in the 2% of the SO?*

- We apologize for this confusion. We have modified this sentence.

→ **3.2 Considering environmental side effects (lines 34–36, page 19):**

“Estimates derived by the extrapolation of SOFeX-N DMS production results suggested that fertilizing ~2% of the SO **area** over the course of a week would **derive a 20% increase of the total SO DMS flux**, which would lead to a 2°C decrease in air temperature over the SO (Wingenter et al., 2007).”

12. P 21 L 12: *The statement about domoic acid levels needs a reference. In several places throughout this paragraph, the authors rely on the conclusions reached by Trick et al. 2010 – however, I don't think that this study gives a reliable guide to possible impacts of Pseudonitzschia blooms. Trick et al. chiefly conducted bottle experiments that yielded a range of several orders of magnitude in cell quotas of DA. Their conclusions were then chiefly based on extrapolating their highest measured cell quota in a bottle incubation with quite unrealistic estimates of likely surface biomass levels to estimate a possible in-water DA concentration. If the authors do wish to cite this paper, I would strongly recommend that these caveats to their conclusions should be pointed out explicitly.*

- Thank you for your suggestion. As Reviewer #1's comments, we have added 'these caveats to their conclusion' in Section 3.2. We have also added some references.

→ **3.2 Considering environmental side effects (lines 1–4, page 21):**

“Trick et al. (2010) suggested that large-scale OIF may induce DA accumulation with developing toxic *Pseudo-nitzschia* blooms. However, large uncertainties remain as Trick et al. (2010) simply extrapolated DA concentration based on bottle incubation experiments with HNLC surface waters to the DA production expected from large-scale OIF.”

13. P 24 L1: *better to say “along with sufficient levels of solar radiation” instead of using “receipt”*

- Thank you. Based on Reviewer #2's editing, we have modified this sentence as follows:

→ **4 Future: Designing future aOIF experiments (lines 36–39, page 23):**

“To the south of the SO PF, phytoplankton blooms usually occur during early summer (i.e., from late December to early January) due to an increase in the nutrient flux from subsurface waters **induced by winter mixing**, along with **the favorable light conditions provided by a shoaling of ML** (Moore and Abbott, 2002).”

14. *P 26 Line 39: MIT and WHOI are entirely separate institutions, despite their collaborative PhD programme.*

- We modified [MIT-WHOI](#) to [WHOI](#).

Reviewer #2

-General Comments:

This third version of review article by Yoon et al. presenting a summary of results from all ocean iron fertilization experiments (OIF) carried to date and introducing plans for a new OIF experiment in the Bransfield Basin under the leadership of the Korean Polar Research Institute. Despite some improvement in the presentation of results from previous experiments, this new version falls short of addressing the issues and comments of both reviewers: the discussion on why previous experiments led to different outcomes is still superficial and lacks a robust analysis of previous results/data and methods. As a consequence, the answers to the questions of why? where/when? and how? still rest mostly on broad statements (see comments below and in the annotated manuscript) and it is not clear how the plans for KIFES are anchored and improve on the lessons learned from previous experiments. finally I believe the issue of C sequestration (which seems to be the main aim in KIFES) is also largely informed by model simulations, therefore, how additional experiments might inform ocean biogeochemical models and affect their results might be an important aspect when presenting the scientific rationale.

- Thank you very much for all your valuable comments. We have addressed all the comments as shown in the revised manuscript. Please see below for a description and/discussion of how we addressed your comments. For annotated editing, we have reflected almost your comments and edits.

-Specific Comments:

1. p. 5, § 1: *The hypothesis mentioned in Coale et al. (1996) are: 1) rapid loss of iron from the patch. 2) Insufficient light after subduction of the patch. 3) Zooplankton Grazing. 4) Limitation by other micro-nutrients or silicate.*

- We have modified 'hypothesis' as follows:

→ 2.1.1 Equatorial Pacific (lines 3–7, page 5):

“Four hypotheses were advanced to explain the weak responses observed: (1) the possibility of unforeseen micro-nutrient (e.g., zinc, cadmium, and manganese) or macro-nutrient (e.g., silicate) limitations, (2) the short residence time of bioavailable iron in the surface patch **due to colloidal aggregation and/or sinking of larger particles containing iron, (3) insufficient light brought about by subduction of the patch, and (4) high grazing pressure by zooplankton (Martin et al., 1994; Cullen, 1995; Coale et al., 1996; Gordon et al., 1998).**”

2. p. 10: *Given that the patch was subducted during IronEx 1, the response of the patch in terms of impact of aOIF on biogeochemistry, is not particularly relevant. I recommend the authors focus on the results of IronEx II. That would help shorten section 2.4.1 and keep the discussion to the point.*

- IronEx-1 was the first aOIF experiment to test Martin's hypothesis. As we mentioned specific objectives of this review in Introduction – “**to provide a thorough overview of the aOIF experiments conducted over the last 25 years**”, the results of IronEx-1 should be included in this review, even though the magnitude of the biogeochemical response in IronEx-1 was not as large as expected due to the subduction of iron-fertilized patch. Therefore, we would like to keep the description on the results of IronEx-1 in Section 2.4.1.

- We have added the information on “*the patch was subducted during IronEx-1*” in Section 2.4.1 as follows:

➔ **2.4.1 Equatorial Pacific** (lines 22–24, page 10):

“While the elevated IronEx-1 Fv/Fm ratios promptly disappeared, **suggesting** rapid iron loss **due to the subduction of the fertilized patch and/or adsorption onto colloidal particles** (perhaps indicative of insufficient iron supply),”

3. p. 11-12, lines 36-38 and 1-3, respectively: *I am not sure what point the authors want to make with this paragraph.*

- We apologize for this confusion. We have revised this paragraph.

➔ **2.4.2 Southern Ocean** (lines 31–34, page 11):

“Satellite observations were used to investigate the changing spatial and temporal distribution of chlorophyll-a concentration in response to iron fertilization in the fertilized patches compared to the surrounding waters; **for example, SOFeX-N/-S found elevated chlorophyll-a concentrations in fertilized patches after iron addition through satellite images (Fig. 7d)** (Boyd et al., 2000; Coale et al., 2004; Westberry et al., 2013).”

4. p. 15, Section 2.5: *I would remove discussion of the FEP in the NA, since it is not relevant in this context (see also annotated manuscript).*

- We have removed ‘*discussion of the FeeP experiment in the NA*’ in **Section 2.2.4, Section 2.4.4, and Section 2.5.**

5. p. 15, Section 2.5, § 1 & 2: *These paragraphs only provide background information that is already given in previous sections. I would remove and focus on the significant results as mentioned in the heading.*

- We have removed these paragraphs in Section 2.5.

6. p.15, lines 35-38: *The authors argue that Si is necessary for bloom formation. Looking at results from SOFeX-N in Fig. 6 and 7, it does not seem to be necessarily the case. As I have mentioned in a previous review, this might be due to the fact that the authors compared concentrations, when they should actually compare standing stocks (i.e. integrated values for the mixed layer).*

- We apologize for this confusion. SOFeX-N/SAGE/LOHAFEX were commonly conducted in the initial low silicate concentrations. However, unlike SAGE and LOHAFEX, during SOFeX-N, silicate limitation in the iron-fertilized waters was temporarily relieved through lateral mixing with the surrounding waters that had relatively higher silicate concentrations (Coale et al., 2004), and it contributed to taxonomic shift toward diatom-dominated communities (from 16% to 44% of total phytoplankton community). Please refer the SOFeX-N results in Section 2.4.2 (lines 13–17, page 12). Finally, we modified this paragraph as follows:

➔ **2.5 Summary of the significant results from aOIF experiments (lines 28–38, page 14):**

“This effect was particularly noticeable as diatoms became the dominant species during IronEx-2, SOIRE, EisenEx, SEEDS-1, SOFeX-S, EIFEX, and SERIES. Diatom-dominated blooms induced >4.5-fold increases in chlorophyll-a concentrations and accounted for >65% of the chlorophyll-a increase (Boyd et al., 2000; Gervais et al., 2002; Coale et al., 2004; Smetacek et al., 2012). **The shift to a diatom-dominated community appears to be related to initial availability of silicate (i.e., initial silicate was >~5 μM in all the experiments listed above).** However, as silicate concentrations decreased to <2 μM due to removal by **phytoplankton**, diatom blooms rapidly declined. SAGE and LOHAFEX had low initial levels of silicate (<2 μM). **As a consequence**, pico and nano-phytoplankton dominated **their** communities and diatom growth was limited by the lack of available silicate. **However during SOFeX-N, initial silicate limitation (<~3 μM) in the iron-fertilized waters was temporarily relieved through lateral mixing with the surrounding waters that had relatively higher silicate concentrations (Coale et al., 2004), which contributed to a taxonomic shift toward diatom-dominated communities (from 16% to 44% of total phytoplankton community).**”

7. p. 16, lines 34-56: the authors make no mention of model simulations. There are several studies on efficiency and potential negative impact from model studies. This should be mentioned when dealing with this topic.

- Thank you for suggestion. We have added as follows:

➔ 3 Present: Unanswered aOIF questions (line 39, page 15 – line 16, page 16):

“However, the impact on the net transfer of CO₂ from the atmosphere to **below the winter MLD** through the ‘biological pump’ (**Fig. 3c**) is not yet fully understood or quantified and appears to vary with environmental conditions, export flux measurement techniques, and other unknown factors (Smetacek et al., 2012). **There have also been a wide range of the estimates of atmospheric CO₂ drawdown resulting from large-scale and long-term aOIF based on model simulations (Joos et al., 1991; Peng and Broecker, 1991; Sarmiento and Orr, 1991; Kurz and Maier-Reimer, 1993; Gnanadesikan et al., 2003; Aumont and Bopp, 2006; Denman, 2008; Jin et al., 2008; Zahariev et al., 2008; Strong et al., 2009; Sarmiento et al., 2010).** While it is generally agreed that OIF effectiveness needs to be determined through **quantification of export fluxes**, there has been no discussion about which export flux measurement techniques are the most effective. **Meanwhile**, concern has been expressed regarding possible environmental side effects in response to iron addition (Fuhrman and Capone, 1991). These side effects include the production of greenhouse gases (e.g., N₂O and CH₄) (Lawrence, 2002; **Jin and Gruber, 2003**; Liss et al., 2005; Law, 2008; **Oschlies et al., 2010**), the development of hypoxia/anoxia in the water column (Sarmiento and Orr, 1991; **Oschlies et al., 2010; Keller et al., 2014**), and toxic algal blooms (e.g., *Pseudo-nitzschia*) (Silver et al., 2010; Trick et al., 2010). **These unwanted side effects** could lead to negative climate and ecosystem changes (Fuhrman and Capone, 1991; **Sarmiento and Orr, 1991; Jin and Gruber, 2003; Schiermeier, 2003; Oschlies et al., 2010**). **Model studies suggested that the unintended ecological and biogeochemical consequences in response to large-scale aOIF might cancel out the effectiveness of aOIF. For example, aOIF enhanced N₂O production may have offset (up to ~40%) the benefits of CO₂ sequestration in the EP (Sarmiento and Orr, 1991; Jin and Gruber, 2003; Oschlies et al., 2010; Hauck et al., 2016).**”

- For more details on model studies on efficiency and potential negative impact, please refer CH₄ (lines 35–37, page 18), N₂O (lines 8–12, page 19), DMS (lines 38–39, page 19), and Oxygen (lines 18–23, page 20) part in 3.2 Section and Section 4 (lines 25–34, page 22).

8. Figure 6, panel b: Given that the size of the initial fertilized patches varied between experiments, it would be more informative to have the target iron concentrations in seawater rather than the total amount used.

- Thank you. We have changed Figure 6b from ‘total amount used’ to ‘target iron concentrations’.

→ Figure captions (line 28, page 55):

“(b) **First target iron concentrations (nM).**”

9. Section 3.1: I am entirely sure what the aim of this paragraph, broad presentations of methods used, or a discussion on the result (export fluxes) from previous experiments? While both issues might be relevant the authors do not really tackle the issues: why export fluxes varied so much in all experiments? Where the methods used appropriate?

- The results of export flux varied with the environmental condition (e.g., silicate concentrations, grazing pressure rates within the mixed layer, and bacterial respiration), experimental duration, and export flux measurement techniques. Based on Reviewer #1’s comments in the second revision (**Please refer Response 7 & 8 in author’s response version 2**), we had added the discussion about the effectiveness of OIF ‘why export fluxes varied so much in all experiments’ in Section 2.5 (line 26, page 14 – line 32, page 15) and Section 3.1 (line 8, page 17 – line 9, page 18).

- For ‘Where the methods used appropriate’, we had also showed the limitations in each method (e.g., discrepancy between the ^{234}Th and sediment traps which was caused by the under- or over-sampling of POC into the drifting traps during SOIREE and SEEDS-1 and limited ability to fully scan the vertical profile of POC fluxes in both sediment traps and ^{234}Th methods during LOHAFEX) and methods used additionally to resolve these limitations (e.g., the deployment of neutrally buoyant sediment trap which could reduce the bias by sediment traps during LOHAFEX, the employment of optical sensors such as underwater video profile or transmissometers during LOHAFEX and EIFEX to monitor vertical changes in sinking particles with high resolution, and deployment of autonomous carbon explorers up to 18 months after initial deployment during SOFeX-N to continuously monitoring iron-induced POC fluxes after aOIF) in Section 3.1 (line 8, page 17 – line 9, page 18). Based on these methods, we concluded that the combination of multiple approaches is essential to accurately observe the POC produced in response to iron addition and its fate (Please refer the last paragraph of Section 3.1: lines 10–17, page 18 and ‘What’ of Section 4: line 38, page 25 – line 11, page 26). Also, we have added the mention about ‘where the export flux should be detected’ based on Reviewer #1’s comments. Please refer the **Reviewer #1’s Response 1**.

10. p. 17, lines 23-29: *The authors seem to imply that ^{234}Th is more reliable than sediment traps. This is not necessarily true: the ^{234}Th method is sensitive to the type of model used (steady-state vs. non-steady state, C/ ^{234}Th ratios, mixing, vertical resolution and uncertainties in measurements especially when sampling deeper layers). Further, the ^{234}Th is not appropriate to measure deep fluxes. While deep fluxes (below the winter mixed layer down to the sediments) are mentioned in p.18, there is no real discussion of this highly relevant issue (which is also related to the monitoring of long-term effects).*

- We have added the limitations of the ^{234}Th approach.

→ 3.1 Export flux measurement methods (lines 4–7, page 17):

“However, the water column-based ^{234}Th method is sensitive to the characterization of the POC to ^{234}Th ratio on sinking particles and/or the choice of ^{234}Th flux models (Buesseler et al., 2006). Therefore, sampling to estimate the POC to ^{234}Th ratio should be conducted below MLD to accurately detect downward carbon export flux into intermediate/deep waters.”

- For discussion on ‘*monitoring of long-term effects*’, we had mentioned the autonomous carbon export, allowing the continuous observation of POC fluxes during and after aOIF experiment in Section 3.1 (lines 2–17, page 18) and Section 4 (lines 1–11, page 26). We have also added the long-term monitoring methods for possible side effects after aOIF experiments as follows.

→ 4 Future: Designing future aOIF experiments (lines 21–24, page 26):

“The possible side effects after an aOIF experiment can be continuously monitored from optical sensors equipped autonomous moored profiler and/or autonomous benthic vehicle (e.g., crawler, which is capable to perform a long-term benthic oxygen measurements for ~12 months) (Dunne et al., 2002; Purser et al., 2013; Wenzhöfer et al., 2016).”

→ 4 Future: Designing future aOIF experiments (lines 33–35, page 26):

“(5) The side effects on marine/ocean ecosystems, including **decline in oxygen contents and the production of climate-relevant gases (e.g., N_2O , DMS, and HVOCs) and toxic DA, **are monitored using optical sensors equipped autonomous moored profiler and/or autonomous benthic vehicle.**”**

→ 5.2.4 Year four plan (lines 21–23, page 29):

“(6) Monitor possible side effects, such as **the decline in oxygen contents and the production of climate-relevant gases and toxic DA, **using optical sensors equipped autonomous moored profiler and/or autonomous benthic vehicle.**”**

11. Section 3.2: The authors make no mention of observations from natural fertilization studies.

- Thank you for pointing this out. We have added the information about ‘*the observations from natural fertilization*’ as follows:

→ 3.2 Considering environmental side effects (lines 31–33, page 18):

“The SO nOIF experiment conducted in 2011 year (i.e., Kerguelen Ocean and Plateau compared Study-2: KEOPS-2) (Table 1) showed that CH₄ concentrations were 4-fold higher in the naturally iron-fertilized patch than in the control area (Farías et al., 2015).”

→ 3.2 Considering environmental side effects (lines 12–13, page 19):

“However, the SO nOIF experiment (i.e., KEOPS-2) suggested that nOIF acts as both a sink and a source for N₂O (Farías et al., 2015).”

→ 3.2 Considering environmental side effects (lines 36–38, page 19):

“On the other hand, the SO nOIF experiment (KEOPS-1) conducted in 2005 year (Table 1) showed that DMS production was not markedly higher in the naturally fertilized area compared to the surrounding waters (Belviso et al., 2008).”

12. p. 20, lines 34-36: This statement is not correct. Keller et al, 2014 does find significant changes in ocean O₂ under climate change scenarios.

- We apologize for the confusion. We have revised this paragraph and added the information about decline in oxygen content as follows:

→ 3.2 Considering environmental side effects (lines 18–24, page 20):

“Early modeling studies suggest that anoxic conditions may develop after long-term, large-scale aOIF (Fuhrman and Capone, 1991; Sarmiento and Orr, 1991), whereas a recent study based on more sophisticated models showed sustained well-oxygenated conditions (O₂ ≈ 120 μM) even under simulated aOIF south of 30°S on a 100 year timescale from 2010 to 2110 (Oschlies et al., 2010). Keller et al. (2014) found that simulated SO large-scale aOIF south of 40°S from the year 2020 to 2100 under a high CO₂ emissions scenario (Meinshausen et al., 2011) may develop suboxia (O₂ <10 μM) in the year 2125. Clearly, the circumstances under which a substantial decline in oxygen inventory can be caused by large-scale aOIF need further study.”

→ Abstract (lines 36–39, page 1):

“(5) Monitoring of side effects on marine/ocean ecosystems, including production of climate-relevant gases (e.g., N₂O, dimethyl sulfide, and halogenated volatile organic compounds), **decline in oxygen inventory**, and development of toxic algae blooms, **with optical sensor equipped autonomous moored profilers and/or autonomous benthic vehicles.**”

→ 3.2 Considering environmental side effects (lines 9–13, page 21):

“The production of climate-relevant gases such as N₂O, DMS, and HVOCs, which is influenced by the remineralization of sinking particles that follows OIF-induced blooms, **the decline in oxygen inventory**, and the production of DA are particularly important to understand. These processes can directly and indirectly modify the effectiveness of carbon sequestration, with either positive or negative effects. Therefore, monitoring **declines in oxygen content** and production of climate-relevant gases and DA to evaluate the effectiveness of aOIF as a geoengineering approach is essential.”

→ 4 Future: Designing future aOIF experiments (lines 16–19, page 26):

“The emissions of climate-relevant gases, such as N₂O, DMS, and HVOCs, may directly contribute to warming or cooling effects, and **oxygen decrease and** toxic DA production may have a negative impact on marine/ocean ecosystems (Law, 2008; Silver et al., 2010; Trick et al., 2010; **Williamson et al., 2012**), resulting in significant offsets against the benefits of aOIF experiments.”

→ 4 Future: Designing future aOIF experiments (lines 33–35, page 26):

“(5) The side effects on marine/ocean ecosystems, including **decline in oxygen contents** and the production of climate-relevant gases (e.g., N₂O, DMS, and HVOCs) and toxic DA, are monitored using optical sensors equipped autonomous moored profiler and/or autonomous benthic vehicle.”

→ 5.2.2 Year two plan (lines 25–27, page 28):

“(1) Using the ice breaker RV *ARAON*, undertake a field investigation in/near the eastern Bransfield Basin to determine physical and biogeochemical parameters associated with both carbon sequestration and aOIF side effects (e.g., **decline in oxygen inventory** and production of N₂O, DMS, HVOCs, and DA), based on the first-year results.”

→ 6 Summary (lines 18–19, page 30):

“Likewise, the possible environmental negative side effects in response to iron addition, such as **decline in oxygen contents** and the production of climate-relevant gases and toxic DA, could not be fully evaluated due to the widely differing outcomes,”

13. p. 21, lines 17-27: I believe the topic of C sequestration is also largely influenced by model simulations, therefore, how additional experiments might inform ocean biogeochemical models might be an important aspect when presenting the scientific rationale (see for instance Losch et al., 2014, <http://dx.doi.org/10.1016/j.jmarsys.2013.09.003>).

● Thank you for your suggestion. However, this paragraph summarizes which environmental side effects should be essentially quantified to evaluate the effectiveness of aOIF as a means for reducing atmospheric CO₂ based on previous aOIF experiments, nOIF experiments, and modeling results. Improvement of the ocean biogeochemical models can be realized from future scientific additional aOIF experiment, like KIFES. Therefore, we think that adding the sentence about *'how additional experiments might inform ocean biogeochemical models might be an important aspect when presenting the scientific rationale'* to Section 5.2.5 KIFES section would be more reasonable. We added this part as follows:

➔ **5.2.5 Year five plan** (lines 33–35, page 29):

“(5) Evaluate effectiveness and environmental side effects of large-scale SO aOIF via more realistic simulations under various scenarios with ocean biogeochemical models using the integrated results of KIFES.”

14. p. 23 § 2: The issue of zooplankton grazing is mentioned only in passing in the next page (§ 1 on timing of experiment), however, zooplankton composition and stocks should also guide the choice of location (for instance areas were krill or salps dominate?) and scientific rationale for KIFES.

● Good point. We have added the mention about *'the issue of zooplankton grazing'* in Section 4 as follows:

➔ **4 Future: Designing future aOIF experiments** (lines 18–24, page 23):

“Previous aOIF experiments have shown that silicate concentration and mesozooplankton **stocks (i.e., copepods) are the crucial factors controlling** diatom **blooms (Boyd et al., 2000; Gervais et al., 2002; Coale et al., 2004; Tsuda et al., 2007; Smetacek et al., 2012).** Therefore, to obtain the greatest possible carbon export flux in response to iron addition, aOIF experiments should be designed in regions with high silicate concentrations **and low grazing pressure. It will be important to conduct initial surveys to measure the degree of grazing pressure in HNLC region with** high silicate concentrations such as in the subarctic NP (e.g., SEEDS-1 experiment) and the south of SO PF (e.g., SOFeX-S experiment) $>15 \mu\text{M}$ (Fig. 4c).”

15. p. 24, § 2: *Duration: The authors should consider the life span of eddies as well as the possibility of using autonomous observation platforms. In addition, risk assessment and monitoring as stipulated in the LC-LP.2 (2010) resolution should also be considered as a basis to determine the duration of the experiments. Further, carrying repeated fertilization might delay sinking events due to iron limitation of phytoplankton.*

- We provided the information about the life span of mesoscale eddies in ‘Where’ of Section 4 (lines 27-28, page 23).

- We have added the sentence about consideration about ‘the possibility of using autonomous observation platforms’ in Section 4 as follows:

→ 4 Future: Designing future aOIF experiments (lines 29–31, page 24):

“In addition, autonomous observation platforms are essential to monitor post-assessment of effectiveness, capacity, and risks of aOIF for at least 12 months after experiment termination.”

- For the risk assessment and monitoring, please refer **Reviewer #2’s Response 10**.

16. p. 25, lines 14-25: *The arguments for several iron infusions are not very robust given that despite a decrease in dissolved iron concentrations Fv/Fm remained high in all experiments before iron additions were repeated. Further, based on the EIFEX results, one could argue that a single iron addition to a final concentration of 1.5-2 nM might be enough to trigger both a bloom as well as lead to significant export (see also previous comment).*

- Thank you for your comments. As we had mentioned in Section 2.3.1 (lines 32–34, page 8), bottle incubation experiments showed that maximum phytoplankton growth rates in response to iron additions occurred when dissolved iron concentrations increased to the target concentrations, 1–2 nM (Fitzwater et al., 1996). However, IronEx-1, single iron addition experiment, showed that dissolved iron concentration rapidly decreased from 3.6 to 0.25 nM ~4 days after iron addition in the center of the fertilized patch due to subduction of the patch and sinking of colloidal aggregation and/or large particles containing iron, suggesting a limit to the level required for phytoplankton growth (please refer Section 2.3.1: lines 11–13, page 9). As we had mentioned in How of Section 4 (lines 10–15, page 25), SOIREEE also showed that first infusion elevated the dissolved iron concentration up to 2.7 nM, and then dissolved iron concentrations was rapidly decreased to 0.2–0.3 nM in ~60 hours after first addition, requiring re-infusion (Bowie et al., 2001). During IronEx-1, Fv/Fm ratio, which indicates iron limitation on phytoplankton growth, showed that increased Fv/Fm after first iron addition continuously decreased ~4 days after first iron addition, but during IronEx-2 increased Fv/Fm after first iron addition remained throughout experiment duration following additional iron infusions (Barber and Hiscock, 2006) (please refer Section 2.4.1: lines 21–26, page 10). During SOIREEE multiple additions of iron led to persistently high levels of both dissolved and particulate iron within the mixed layer, with a

rapid reduction at the end of the experiment, combined with an increase in the concentration of iron-binding ligands (Bowie et al., 2001) (please refer Section 4: lines 12–15, page 25). In both EIFEX and SOFeX-S, it was also found that multiple iron infusions allowed iron to persist in the mixed layer longer than its expected oxidation kinetics (Croot et al., 2008) (please refer Section 4: lines 15–18, page 25). Therefore, we think that the suggestion of multiple iron infusions would be more reasonable for aOIF experimentation.

17. p. 27: *I still find the argument for the location (Bransfield straight) somewhat tenuous as they could apply to other location as well. Further, the location seems to be situated in a sensitive area (CCAMLR subarea 48.1) with high krill stocks adding a whole new level of complexity in following both the evolution of the patch and impacts on higher trophic levels. I suggest the location should be re-evaluated based on considerations including food webs and hydrography (including what would be the fate of the patch and surface properties as well as exported organic matter once the eddy collapses).*

- Thank you for your suggestion. We believe that we had faithfully answered to Reviewers's comment on the location of Bransfield region with detailed explanations in **Response 13 in Author's response version 2**. Detailed background/information are found in '*specific proposal for KIFES*', '*description of the logic for conducting this in the Bransfield Strait*', '*what are the hypothesis*', and '*how the KIFES project improve on previous experiments*'.

- For aOIF experiment in the krill habitat, Smetacek and Naqvi (2008) suggested that the krill habitat may provide a suitable site for a future SO aOIF experiment, as they have a potential (Tovar-Sanchez et al. 2007) for sustaining biological pump after the experiment and in turn recovering the whale populations, leading to positive effect on marine ecosystem. However, there has been little information about the relationship between mesoscale eddies and krill habitat in previous aOIF experiments. Therefore, we think that at present it is inappropriate dealing with the subject in this review.

-Annotated Comments in the Manuscript:

❖ AC1: In ‘caution is required because artificially high levels of SF₆ injection may negatively impact the interpretation of low-level SF₆ signals dissolved in seawater via air-sea exchange.’,

* SF₆ has other advantages: can help provide estimates of vertical and horizontal mixing/diffusivity.

* meaning of this sentence unclear. The main issue with SF₆ is that concentrations decrease relatively rapidly (mixing and air-sea exchanged). Hence, it allows marking of the patch as a whole only once and for a limited period of time.

- We have modified this sentence and added sentences concerning ‘main issue with SF₆’ in Section 2.3.2 and Section 4 as follows:

→ 2.3.2 Tracing iron-fertilized patch (lines 29–34, page 9):

“Although these earlier experiments demonstrated that the injection of artificial SF₆ is a useful technique for following iron-fertilized patches, **SF₆ can only be used for limited period (~2 weeks) due to the loss at the surface through air-sea gas exchange (Law et al., 2006; Tsumune et al., 2009; Martin et al., 2013). Furthermore**, caution is required because artificially high levels of SF₆ injection may negatively impact the interpretation of low-level SF₆ signals dissolved in seawater via air-sea exchange **to estimate tracer-based water mass ages for understanding physical circulation (Fine, 2011).**”

→ 4 Future: Designing future aOIF experiments (lines 32–33, page 25):

“**However, tracing via SF₆ allows for only a limited period (~2 weeks) due to air-sea gas exchange (Law et al., 2006; Tsumune et al., 2009; Martin et al., 2013).**”

❖ AC2: In ‘(~50%)’,

* this value must be explained.

- We have added explanation as follows:

→ 2.4.1 Equatorial Pacific (lines 7–9, page 11):

“The biomass of meso-zooplankton (200–2,000 μm), such as copepods, grew simultaneously, substantially increasing the **community grazing effect of larger animals on phytoplankton standing stocks from 7.8% d⁻¹ outside patch to 11.4% d⁻¹ in the patch** (Coale et al., 1996).”

❖ **AC3: In ‘2.4.2 Southern Ocean’,**

** As the Southern Ocean is the most relevant here, I would shift this section to the end of section 2.4.*

- Thank you. However, based on Reviewers’s comments in second revision (please refer **Response 21 in Author’s response version 2**), we had organized the Section 2 and Figures 6–7 ‘*by region with HNLC regions*’. As we had presented the data by the regions (EP-SO-Subarctic NP) in Section 2.1 and Section 2.2, it is likely appropriate to sustain original version (EP-SO-Subarctic NP) in Section 2.4.2.

❖ **AC4: In ‘results suggested that diatoms were abundant in the two experiments.’,**

** How so?*

- We have revised this sentence in Section 2.4.2 as follows:

➔ **2.4.2 Southern Ocean (lines 20–22, page 12):**

“**Although both initially dominated by diatoms**, SOFeX-S had a somewhat greater ΔNO_3^- ($-3.5\ \mu\text{M}$) and $\Delta p\text{CO}_2$ ($-36\ \mu\text{atm}$) than EIFEX (ΔNO_3^- : $-1.6\ \mu\text{M}$ and $\Delta p\text{CO}_2$: $-30\ \mu\text{atm}$) (Coale et al., 2004; Hoffmann et al., 2006; Smetacek et al., 2012; Assmy et al., 2013).”

❖ **AC5: In ‘This augmentation was the largest among all the aOIF experiments (Tsuda et al., 2003). The dramatic chlorophyll-a increase observed during SEEDS-1 was partly attributed to the particular range of seawater temperature in the region, which was conducive to diatom growth (i.e., 8–13°C) as well as to the shallower MLD (~10 m), which provided a relatively longer surface water residence time for the additional iron (Figs. 6c and d) (Noiri et al., 2005; Takeda and Tsuda, 2005; Tsumune et al., 2005).’,**

** When comparing integrated values (over the mixed layer), other experiments reached similar high biomass. Given the broad range in mixed layer depths, comparing concentrations can give a wrong impression of the actual biomass increase.*

- We apologize for missing your comment in first revision (**Response 17 in Author’s response version 1**) when we re-organized our manuscript during second revision. We have re-added mention about ‘*the difference between surface chlorophyll-a concentrations*’ and ‘*integrated chlorophyll-a concentrations or/and primary production*’ in Section 2.5, which we had revised in first revision and have added ‘surface’ word in the sentence.

➔ **2.4.3 Subarctic North Pacific (lines 29–31, page 13):**

“The dramatic **surface** chlorophyll-a increase observed during SEEDS-1 was partly attributed to the particular range of seawater temperature in the region, which was conducive to diatom growth (i.e., 8–13°C) as well as to the shallower MLD (~10 m),”

➔ 2.5 Summary of the significant results from aOIF experiments (lines 7–14, page 15):

“**Among previous aOIF experiments**, the subarctic NP SEEDS-1 experiment, which was conducted under temperature conditions ideal for diatom growth ($\sim 8^{\circ}\text{C}$) and with shallow MLDs ($\sim 10\text{ m}$), produced the greatest changes in **surface phytoplankton biomass**. **However, influence of iron addition on the phytoplankton growth covers from surface to euphotic depth as added iron is mixed within the ML by physical processes (Coale et al., 1998). Although maximum surface chlorophyll-a concentration during SEEDS-1 ($\sim 22\text{ mg m}^{-3}$) was much higher than EIFEX ($\sim 3.2\text{ mg m}^{-3}$), the MLD-integrated chlorophyll-a concentrations were similar to $\sim 250\text{ mg m}^{-2}$ between two experiments. Therefore, to quantify the exact changes in phytoplankton biomass in response to iron addition, it would be eligible to consider the MLD-integrated PP for comparison.**”

❖ AC6: In ‘2.5 Summary of the significant results from a OIF experiments’,

* *These are not results!*

- Please refer **Reviewer #2’s Response 5**.

❖ AC7: In ‘changeover to a diatom-dominated community after iron addition is needed.’,

* *How about SOFEX-N then?*

- Please refer **Reviewer #2’s Response 6**.

❖ **AC8: In ‘addition were not dramatic compared to natural values.’,**

** I am not sure I understand the argument here: natural values are not necessarily low.*

- We had referred ‘Buesseler et al. (2005)’ and added it.

In ‘Buesseler et al. (2005)’,

‘Thus, the lack of a stronger inverse correlation between surface SF_6 and ^{234}Th , is evidence that during the 4-week occupation of the SOFeX bloom, surface ocean export was not particularly elevated, or at least was small relative to other observations of particle export associated with natural blooms from other regions.’

‘When converted to POC and $bSiO_2$ export rates, these fluxes were not large relative to natural blooms and were, in fact, smaller than those observed under natural conditions at this site during a different field season.’

→ 2.4.2 Southern Ocean (lines 39–40, page 12):

“During SOFeX-S, significantly enhanced POC fluxes below the **MLD** similar to those observed in natural blooms, were estimated from ^{234}Th measurements after iron enrichment (**Buesseler et al., 2005**).”

→ 2.5 Summary of the significant results from aOIF experiments (lines 21–22, page 15):

“However, the changes in export flux, after iron addition, were not dramatic compared to natural values (**Buesseler et al., 2005**).”

❖ **AC9: In ‘offset leading’,**

** I would change the wording, as offset is used in the literature as counteracting the effects OIFs, This is not the case here.*

- We have modified this wording as follows:

→ 3.2 Considering environmental side effects (lines 15–16, page 20):

“**Another important consideration is the extent to which** the effectiveness of **aOIF is cancelled out by its tendency to** lead to ocean ecosystem changes such as a decrease in dissolved oxygen and an increase in domoic acid (DA) levels.”

❖ **AC10: In ‘It is likely that several phytoplankton samples (e.g., *Pseudo-nitzschia* abundance: 1.3×10^6 cells L⁻¹ in IronEx-2 and 7.5×10^4 cells L⁻¹ in SOFeX-S) collected with a net tow were suitable to detect these changes.’,**

** unclear. please explain.*

● We apologize for the confusion. Unlike the results of Trick et al. (2010) that showed no DA change during EisenEx and SERIES, Silver et al. (2010) suggested that discernable change in DA during IronEx-2 and SOFeX-S could be found from phytoplankton samples taken with net tows (20- to 30-μm mesh phytoplankton nets) by allowing larger phytoplankton samples including *Pseudo-nitzschia* to concentrate. During IronEx-2 and SOFeX-S, DA concentrations were as high as 45 ng DA l⁻¹ and 220 ng DA l⁻¹ in the water, respectively, i.e., toxin levels high enough to damage marine communities in coastal waters (Scholin et al., 2000; Schnetzer et al., 2007). Based on these results, we concluded that it is necessary to quantify DA production in response to iron additions, with concentrated larger phytoplankton samples taken from a net tow. To explain clearly these contents, we have modified these sentences in Section 3.2.

➔ 3.2 Considering environmental side effects (line 35, page 20 – line 1, page 21):

“It is likely that **detection was possible because these samples were collected with net tows (20- to 30-μm mesh phytoplankton nets), which provided concentrated samples of larger phytoplankton including *Pseudo-nitzschia*** (e.g., *Pseudo-nitzschia* abundance: 1.3×10^6 cells L⁻¹ in IronEx-2 and 7.5×10^4 cells L⁻¹ in SOFeX-S). During IronEx-2 and SOFeX-S, high cell abundances of *Pseudo-nitzschia* (10^6 and 10^5 cells l⁻¹, respectively) combined with moderate DA **cell** quotas (0.05 and 1 pg DA cell⁻¹, respectively) produced toxin levels as high as 45 ng DA l⁻¹ and 220 ng DA l⁻¹ **in the water**, respectively, i.e., toxin levels high enough to damage marine communities in coastal waters (**Scholin et al., 2000; Schnetzer et al., 2007**).”

➔ 3.2 Considering environmental side effects (lines 4–5, page 21):

“As a result, it is necessary to **clarify/quantify** DA production in response to aOIF, with concentrated **larger** phytoplankton samples **collected using** net tows **(20- to 30-μm mesh phytoplankton net)**.”

❖ **AC11: In ‘toxin levels high enough to damage marine communities in coastal waters.’,**

** add references.*

● Done. Please refer **Reviewer #2’s AC10**.

❖ **AC12:** In ‘Therefore, it is necessary to quantify DA production in response to iron additions, with concentrated phytoplankton samples (i.e., large numbers of cells) using a net tow.’,

* *Where does this come from? It is not the conclusion I would make from the previous. Rather handle the samples in such a way that DA remains unchanged. Also what about DA in the water?*

- Please refer **Reviewer #2’s AC10**.

❖ **AC13:** In ‘Where: The first consideration for a successful aOIF experiment is the location. The dominance of diatoms in phytoplankton communities plays a major role in increasing the biological pump because diatom species can sink rapidly as aggregates or by forming resting spores (Tréguer et al., 1995).’,

* *see also more recent literature:*

Rembauville, Mathieu, et al. Export fluxes in a naturally iron-fertilized area of the Southern Ocean—Part 2: Importance of diatom resting spores and faecal pellets for export." Biogeosciences 12.11 (2015): 3171-3195. Rembauville, M., et al. Strong contribution of diatom resting spores to deep-sea carbon transfer in naturally iron-fertilized waters downstream of South Georgia." Deep Sea Research Part I: Oceanographic Research Papers 115 (2016): 22-35. Salter, Ian, et al. Estimating carbon, silica and diatom export from a naturally fertilised phytoplankton bloom in the Southern Ocean using PELAGRA: A novel drifting sediment trap." Deep Sea Research Part II: Topical Studies in Oceanography 54.18 (2007): 2233-2259.

- Thank you. We have added these references and added the sentence that mentions diatom resting spores.

➔ **4 Future: Designing future aOIF experiments (lines 14–18, page 23):**

“The dominance of diatoms in phytoplankton communities plays a major role in increasing the biological pump because diatom species can sink rapidly as aggregates or **by forming resting spores to efficiently bypass the intense grazing pressure of mesozooplankton (e.g., copepods, salps, and krill) and export carbon out of the winter ML** (Tréguer et al., 1995; Salter et al. 2007; Assmy et al., 2013; Rembauville et al., 2015; Rembauville et al., 2016).”

❖ **AC14:** In ‘while after January (i.e., during late summer and early autumn from February to March) it is mainly limited due to silicate availability.’,

* *please provide references*

- We have added references.

❖ AC15: In ‘(de Baar et al., 2005;’,
* *De Baar 2005 states that light is limiting*

- Thank you for pointing this out. We have modified this sentence.

➔ 4 Future: Designing future aOIF experiments (line 41, page 23 – line 3, page 24):

“In previous SO aOIF experiments conducted between spring and early autumn, PP was mainly limited by iron and/or silicate availability rather than light availability **(except when heavy clouds led to severe light limitation, only occurred for a few days during EisenEx) (Gervais et al., 2002; Bakker et al., 2005; Smetacek and Naqvi, 2008; Pelouquin et al., 2011b).**”

❖ AC16: In ‘(Coale et al., 2004; Martin et al., 2013),’,
* *I have not found convincing evidence for grazing control of diatom growth this in the articles cited.*

- We have removed these references and added more appropriate references **(Schultes et al., 2006; Smetacek and Naqvi, 2010)**, reporting grazing control on diatom growth.

In ‘Schultes et al. (2006)’:

‘Overall grazing impact of copepods in these studies reached 25% of phytoplankton standing stock removed per day with the major fraction being attributable to C. simillimus.’

In ‘Smetacek and Naqvi (2010)’:

‘The second fertilization had no noticeable effect on growth rates or biomass of phyto- or bacterioplankton. Apparently, the main reason why biomass did not build up to higher levels was due to heavy grazing of the large copepod population. Incubation experiments indicated that the copepods increased their feeding and faecal production rates inside the patch.’

❖ AC17: In ‘(Le Quéré et al., 2016).’,

* *This is a model simulation paper. Should not be referred here.*

- We have removed this paper and added more appropriate references as follows:

Hunt, B. P. V. and Hosie, G. W.: The seasonal succession of zooplankton in the Southern Ocean south of Australia, part II: The Sub-Antarctic to Polar Frontal Zones, *Deep-Sea Res. Pt. I*, 53, 1203-1223, 2006.

Rembauville, M., Blain, S., Armand, L., Quéguiner, B., and Salter, I.: Export fluxes in a naturally iron-fertilized area of the Southern Ocean – Part 2: Importance of diatom resting spores and faecal pellets for export, *Biogeosciences*, 12, 3171-3195, <https://doi.org/10.5194/bg-12-3171-2015>, 2015.

❖ AC18: In ‘(2) below the depth of the winter MLD to detect iron-induced export carbon fluxes into intermediate/deeper waters (Bidigare et al., 1999; Nodder et al., 2001; Boyd et al., 2004; Buesseler et al., 2004; Coale et al., 2004; Aono et al., 2005; Buesseler et al., 2005; Tsuda et al., 2007; Smetacek et al., 2012; Martin et al., 2013). Sinking-particle profiling systems mounted on autonomous floats, such as a transmissometer and UVP that measure and photograph sinking particles, could provide a record of the temporal and vertical evolution of iron-induced POC stocks through successive depth layers down to ~3,000-m depth for ~20 months after deployment, once calibrated using POC fluxes measured from sediment traps and/or a water-column based ²³⁴Th method (Bishop et al., 2004; Smetacek et al., 2012; Martin et al., 2013).’,

* *What about deep-sea (moored) traps and benthic oxygen measurements?*

- Thank you for comments. Please refer **Reviewer #2’s response 10**.

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Ocean Iron Fertilization Experiments: Past–Present–Future looking to a future Korean Iron Fertilization Experiment in the Southern Ocean (KIFES) Project

5 Joo-Eun Yoon¹, Kyu-Cheul Yoo², Alison M. Macdonald³, Ho-Il Yoon², Ki-Tae Park², Eun Jin Yang²,
Hyun-Cheol Kim², Jae Il Lee², Min Kyung Lee², Jinyoung Jung², Jisoo Park², Jiyoung Lee¹, Soyeon
Kim¹, Kitae Kim^{2*}, and Il-Nam Kim^{1*}

¹Department of Marine Science, Incheon National University, Incheon 22012, Republic of Korea

10 ²Korea Polar Research Institute, Incheon 21990, Republic of Korea

³WHOI, MS 21, 266 Woods Hole Rd., Woods Hole, MA 02543, USA

*Correspondence to: Il-Nam Kim (ilnamkim@inu.ac.kr) and Kitae Kim (ktkim@kopri.re.kr)

15 **Abstract.** Since the start of the industrial revolution, human activities have caused a rapid increase in atmospheric CO₂
concentrations, which have, in turn, had an impact on climate leading to global warming and ocean acidification. Various
approaches have been proposed to reduce atmospheric CO₂-concentrations. The 'Martin (or Iron) Hypothesis' suggests that
ocean iron fertilization (OIF) could be an effective method for stimulating oceanic carbon sequestration through the
biological pump in iron-limited, high-nutrient, low-chlorophyll (HNLC) regions. To test the Martin hypothesis, 13 artificial
20 OIF (aOIF) experiments have been performed since 1990 in ~~the Southern Ocean (seven experiments), in the subarctic Pacific~~
~~(three experiments), in the equatorial Pacific (two experiments), and in the subtropical Atlantic (one experiment)~~ HNLC
regions. These aOIF field experiments have demonstrated that primary production can be significantly enhanced by the
artificial addition of iron. However, except in the Southern Ocean European Iron Fertilization Experiment, the effectiveness
of aOIF of export production (i.e., –the amount of iron-induced carbon export flux below the winter mixed layer depth
25 export of carbon from surface waters into intermediate/deep waters) revealed by the aOIF experiments has been–
was unexpectedly low compared to that achieved by natural phytoplankton blooms, except in the Southern Ocean European Iron
Fertilization Experiment. These results, including possible side effects (e.g., changes in climate relevant gas emissions and
an increase in toxic phytoplankton species) have been debated amongst those who support and oppose aOIF experimentation,
but and many questions such as effectiveness of scientific aOIF, environmental side effects, and international aOIF law
30 frameworks remain. In the context of increasing global and political concerns associated with climate change, it is valuable
to examine the validity and usefulness of the aOIF experiments. To maximize the effectiveness of aOIF experiments under
international aOIF regulations in the future, we suggest a design that incorporates several conditions components. (1)
Experiments are conducted in the center of an eddy structure when grazing pressure is low and silicate levels are high (e.g.,
in the case of the Southern Ocean, at the south of polar front during the early summer). (2) Shipboard observations are made
35 duration extending over a minimum of ~40 days, with multiple iron injections (at least 2 (or three) iron infusions of ~2,000 kg at
least three times, with an interval of ~10–15 days, to fertilize a patch of 300 km² to and obtain a ~2 nM concentration). (3)
Tracing of the The iron fertilized patch is traced using both physical (e.g., a drifting buoy) and biogeochemical (e.g., sulfur
hexafluoride and photosynthetic quantum efficiency the Fv/Fm ratio, where Fm is the maximum chlorophyll fluorescence
yield and Fv is the difference between Fm and the minimum chlorophyll fluorescence yield) tracers. (4) Employment of A
40 neutrally buoyant sediment traps system and application of the water-column derived ²³⁴Thorium method are employed at

two depths (~~i.e., one within is just below~~ the *in situ* mixed layer ~~(i.e., 10–20 m below)~~ depth and ~~another below it at the depth of the depth of the~~ winter mixed layer depth ~~(i.e., ~450 m)~~), with autonomous profilers equipped with an underwater video profiler and a transmissometer, ~~to estimate accurately the carbon export flux.~~ (5) Monitoring of ~~The~~ side effects on marine/ocean ecosystems ~~are monitored~~, including ~~the~~ production of climate-relevant gases (e.g., N₂O, dimethyl sulfide, and halogenated volatile organic compounds), ~~the decline in oxygen inventory~~, and ~~an increase in the abundance of toxic phytoplankton species~~ ~~the development of toxic~~ algae blooms ~~(e.g., production of domoic acid)~~, ~~with are monitored using optical sensor equipped autonomous moored profilers and/or autonomous benthic vehicles.~~ ~~Finally~~ ~~Lastly~~, we introduce the scientific aOIF experimental design guidelines for a future Korean Iron Fertilization Experiment in the Southern Ocean.

Keywords: Ocean Iron Fertilization; High-Nutrient and Low-Chlorophyll regions; Biological Pump; Phytoplankton; Iron

1 Introduction

Since the start of the industrial revolution, human activities have caused a rapid increase in atmospheric carbon dioxide (CO₂, ~~awhich is major greenhouse gas~~) from ~280 ppm (pre-industrial revolution) to ~400 ppm (present day) (<http://www.esrl.noaa.gov/>), which has, in turn, led to global warming and ocean acidification, indicating that there is an urgent need to reduce global greenhouse gas emissions (IPCC, 2013) (Fig. 1). As the Anthropocene climate system has rapidly become more unpredictable, the scientific consensus is that the negative outcomes are a globally urgent issue that should be resolved in a timely manner for the sake of all life on Earth (IPCC, 1990, 1992, 1995, 2001, 2007, 2013). The various ideas/approaches that have been proposed to relieve/resolve the problem of global warming (Matthews, 1996; Lenton and Vaughan, 2009; Vaughan and Lenton, 2011; IPCC, 2014; Leung et al., 2014; Ming et al., 2014) largely fall into two categories: (1) reduction of atmospheric CO₂ by the enhancement of biological CO₂ uptake (including ocean fertilization) and/or the direct capture or storage of atmospheric CO₂ through chemically engineered processes, and (2) control of solar radiation by artificial aerosol injection into the atmosphere to augment cloud formation and cloud brightening to elevate albedo (Fig. 2). One of the most attractive methods among the proposed approaches is ocean fertilization (<https://web.whoi.edu/ocb-fert/>), which targets the drawdown of atmospheric CO₂ by nutrient addition (e.g., iron, nitrogen, or phosphorus compounds) to stimulate phytoplankton growth and, subsequently, carbon export to the deep ocean or sediments via the ocean biological pump (ACE CRC, 2015).

The ocean biological pump is frequently depicted as a single combined process, whereby organic matter produced by phytoplankton during photosynthesis in surface waters is quickly transported to intermediate and/or deep waters (Fig. 3a) (Volk and Hoffert, 1985; De La Rocha, 2007). Although the effectiveness of the biological pump is primarily controlled by the supply of macro-nutrients (i.e., nitrate, phosphate, and silicate) from the deep ocean into the mixed layer (ML) leading to new production (Sarmiento and Gruber, 2006), iron acts as an essential micro-nutrient to stimulate the uptake of macro-nutrients for phytoplankton growth (Fig. 3b) (Martin and Fitzwater, 1988; Martin, 1990; Morel and Price, 2003). In the subarctic North Pacific (NP), equatorial Pacific (EP), and Southern Ocean (SO), which are well known as high-nutrient and low-chlorophyll (HNLC) regions (Figs. 4a and b), phytoplankton cannot completely utilize the available macro-nutrients (particularly nitrate) for photosynthesis due to a lack of iron. As a consequence, primary production (PP) in these HNLC regions is relatively low, despite the high availability of macronutrients (in particular nitrate and phosphate) (Figs. 4a and b).

Analyses of trapped air bubbles in Arctic/Antarctic ice cores have revealed that atmospheric CO₂ (~180 ppm) during the Last Glacial Maximum (LGM; ~20,000 years ago) was much lower than during pre-industrial times (~280 ppm) (Neftel et al., 1982; Barnola et al., 1987; Petit et al., 1999). Over the last 25 years, several hypotheses have been proposed to explain the lowered atmospheric CO₂ level during the LGM (Broecker, 1982; McElroy, 1983; Falkowski, 1997; Broecker and Henderson, 1998; Sigman and Boyle, 2000). Dust inputs are generally regarded as a major natural iron source for ocean fertilization, and Martin (1990) hypothesized that during the LGM increased dust inputs relieved iron-limitation and, thereby, substantially enhanced the biological pump in HNLC regions, particularly in the SO (Fig. 3b). Since Martin's hypothesis was first published, there has been an enormous interest in ocean iron fertilization (OIF) because only a small amount of iron (C:Fe ratios = 100,000:1, Anderson and Morel, 1982) is needed to stimulate a strong phytoplankton response. Therefore, much of the investigative focus has centered on the artificial addition of iron to HNLC regions as a means of enhancing carbon fixation and subsequent export via the biological pump (ACE CRC, 2008).

To test Martin's hypothesis, ~~two-six~~ natural OIF (nOIF) and 13 artificial OIF (aOIF) experiments have been performed to date in the subtropical North Atlantic (NA), EP, subarctic NP, and SO (Blain et al., 2007; Boyd et al., 2007; Pollard et al., 2009; Strong et al., 2009; Smetacek et al., 2012; Martin et al., 2013; Blain et al., 2015) (Fig. 4a and Table 1). These OIF experiments demonstrated, particularly for the SO, that PP could be significantly increased after iron addition (de Baar et al.,

2005; Boyd et al., 2007). However, for aOIF to be considered as a useful geoengineering approach (IPCC, 2007), in the long run, the most critical issue is the ‘effectiveness of a-OIF’. That is, whether ~~thea significant portionsubstantial amounts~~ of ~~the~~ organic carbon produced by aOIF in the surface waters is exported below the winter mixed layer depth (MLD) ~~lead to a significant export~~ to intermediate/deep layers ~~andfor~~ long-term (~1,000 years) storage (Fig. 3c) (Lampitt et al., 2008). A high carbon export was observed in the nOIF experiments in the SO near the Kerguelen Plateau and Crozet Islands (Blain et al., 2007; Pollard et al., 2009). However, all aOIF experiments have shown unexpectedly low carbon exports compared to natural systems (de Baar et al., 2005; Boyd et al., 2007), except for the SO European Iron Fertilization Experiment, EIFEX (Smetacek et al., 2012). The results of these experiments, as well as the potential side effects (e.g., N₂O production and development of hypoxia) (Fuhrman and Capone, 1991), have been scientifically debated amongst those who support and oppose aOIF experimentation (Chisholm et al., 2001; Johnson and Karl, 2002; Lawrence, 2002; Buesseler and Boyd, 2003; Smetacek and Naqvi, 2008; Williamson et al., 2012).

In the context of increasing global (social-political-economic) concerns associated with rapid climate change, it is necessary to examine the validity and usefulness of aOIF experimentation as a climate change mitigation strategy. Therefore, the purpose of this paper is to: (1) provide a thorough overview of the aOIF experiments conducted over the last 25 years; (2) discuss aOIF-related important unanswered questions, including carbon export measurement methods, potential side effects, and international law; (3) suggest considerations for the design of future aOIF experiments to maximize the effectiveness of the technique and begin to answer open questions; and (4) introduce design guidelines for a future Korean Iron Fertilization Experiment in the Southern Ocean (KIFES) project.

2 Past: Overview of previous aOIF experiments

A total of 13 aOIF experiments have been conducted in the following areas: 12 experiments were conducted in the three main HNLC (i.e., nitrate >~10 µM) regions: two in the EP, three in the subarctic NP, and seven in the SO (Table 1, Figs. 4a and b). One experiment was conducted in the subtropical NA, known to be a low-nutrient and low-chlorophyll (LNLC) (i.e., nitrate <1 µM) region. These aOIF experiments have been conducted with various/multiple objectives/hypotheses to investigate the biogeochemical responses of ocean environments to artificial iron additions (Table 2). This overview of past aOIF experimentation begins in Section 2.1, with a presentation of the reasons why each experiment was performed and the main hypotheses (Table 2). The unique ocean conditions for the various experiments are described in Section 2.2. Iron addition and ~~the~~ tracing methods of iron are described in Section 2.3. The biogeochemical responses to the aOIF experiments are presented in Section 2.4, and finally the significant findings from these experiments are summarized in Section 2.5.

2.1 Objectives/hypotheses of previous aOIF experiments

2.1.1 Equatorial Pacific

Initially, Martin’s hypothesis was supported by the results of laboratory and shipboard iron-enrichment bottle experiments (Hudson and Morel, 1990; Brand, 1991; Sunda et al., 1991; DiTullio et al., 1993; Hutchins et al., 1993). However, the extrapolation of these results based on bottle incubations that exclude the lower trophic levels in a marine ecosystem to higher trophic community levels has been strongly criticized due to possible underestimates in grazing rates ~~forcing~~ and other bottle containment effects. To deal with these issues, *in situ* iron fertilization experiments at the whole-ecosystem level are required. Under the hypothesis that ~~artificial iron addition~~ aOIF would increase phytoplankton

productivity by relieving iron limitations on phytoplankton in HNLC regions, the first aOIF experiment, Iron Enrichment Experiment (IronEx-1), was conducted over 10 days in October 1993 in the EP where high light intensity ~~light~~ and ~~warm~~ temperatures would ~~promote~~assist rapid phytoplankton growth (Table 1 and Fig. 4a) (Martin et al., 1994; Coale et al., 1998).

However, the magnitude of the biogeochemical responses in IronEx-1 was not as large as expected (Martin et al., 1994). ~~Three-Four~~ hypotheses were advanced to explain the weak responses observed: (1) the possibility of ~~other~~ unforeseen micro-nutrient (e.g., zinc, cadmium, and manganese) or macro-nutrient (e.g., silicate) limitations, (2) the short residence time of bioavailable iron in the surface patch due to ~~an unstable water column structure colloidal aggregation and/or sinking of larger particles containing iron~~, (3) ~~insufficient light brought about by subduction of the patch~~, and (34) ~~the extremely~~ high grazing pressure ~~by~~of zooplankton (Martin et al., 1994; Cullen, 1995; Coale et al., 1996; Gordon et al., 1998). To test the ~~three-four~~ hypotheses, a second aOIF experiment, IronEx-2, was conducted in May 1995 (Coale et al., 1996). The IronEx-2 research cruise investigated the same area for a longer period (17 days), providing more time to collect information about the biogeochemical, physiological, and ecological responses to the aOIF-experiment.

2.1.2 Southern Ocean

The SO plays an important role in intermediate and deep-water formation, and has the ~~largest~~greatest potential ~~of any of the major ocean basins~~ for carbon sequestration associated with artificial iron addition (Martin, 1990; Sarmiento and Orr, 1991; Cooper et al., 1996; Marshall and Speer, 2012). It is known as the largest HNLC region in the World Ocean and models simulating aOIF have predicted that among all HNLC regions, the effect of OIF on carbon sequestration is greatest in the SO (Sarmiento and Orr, 1991; Aumont and Bopp, 2006). However, a simple extrapolation of the IronEx-2 results to the SO was not deemed appropriate because of the vastly different environmental conditions (Coale et al., 1996) ~~and~~, therefore, ~~based on the lessons from the EP experiments, several aOIF experiments were carried out in the SO this basin became the next region selected for an aOIF experiment~~ (Frost, 1996; Boyd et al., 2000; Smetacek, 2001; Coale et al., 2004; Harvey et al., 2010; Smetacek et al., 2012; Martin et al., 2013). ~~To test the roles of~~ With the hypothesis that iron and light availability ~~may act~~ as key factors ~~controlling that control~~ phytoplankton dynamics, community structure, and grazing in the SO, the Southern Ocean Iron Release Experiment (SOIREE) (Table 1 and Fig. 4a), ~~which was~~ the first *in situ* aOIF experiment performed in the SO, took place in February 1999 (13 days) in the Australasian-Pacific sector (Boyd et al., 2000).

The following year, a second aOIF experiment in the SO, EisenEx ('Eisen' means iron in German), was performed in November within an Antarctic Circumpolar Current eddy in the Atlantic sector (Smetacek, 2001). This region is considered to have a relatively high iron supply, which is supported by dust inputs and ~~possibly icebergs sea ice melt~~ (de Baar et al., 1995; Quéguiner et al., 1997; Smetacek et al., 2002). EisenEx was designed to test ~~how the hypothesis that~~ atmospheric dust, an important source of iron in ocean environments, might have led to a dramatic increase in ocean productivity during the LGM due to the relief of iron-limiting conditions for phytoplankton growth (Abelmann et al., 2006).

In addition to iron availability, the supply of silicate is also considered to be an important factor controlling PP in the SO. Silicate-requiring diatoms, which are large-sized phytoplankton, ~~play have~~ an important role in the biological pump and are responsible for ~75% of the annual PP in the SO (Tréguer et al., 1995). The silicate concentrations in the SO ~~show have~~ a decreasing northward gradient, in particular, on either side of the Antarctic Polar Front (PF), with low silicate concentrations (<5 µM) in the sub-Antarctic waters north of the PF (<61°S) and high silicate concentrations (>60 µM) to the south of the PF (Fig. 4c). Therefore, to address the ~~impact of potential for~~ iron and silicate on phytoplankton communities and export interactions to regulate the diatom bloom, two aOIF experiments were conducted during January–February 2002 in two distinct regions: the Southern Ocean iron experiment-north (SOFeX-N) and -south (SOFeX-S) of the PF (Table 1) (Coale et al., 2004; Hiscock and Millero, 2005). ~~In these two experiments, it was hypothesized that conditions that provided sufficient silicate and iron would lead to high diatom production, while sufficient iron alone would not lead to a~~

diatom bloom (Coale et al., 2004).

Two years later, the Surface Ocean Lower Atmosphere Study (SOLAS) Air–Sea Gas Exchange (SAGE) experiment was conducted during March–April 2004 (15 days) in sub-Antarctic waters, which are typically HNLC with are HNLC and low silicate concentrations waters (HNLC_{LSi}). The aim was to determine the response of phytoplankton dynamics to iron addition in an HNLC_{LSi} region (Fig. 4c) (Law et al., 2011). SAGE was designed with the assumption that the response of phytoplankton blooms to artificial iron addition aOIF could be detected by enhanced air-sea exchanges of climate-relevant gases (e.g., CO₂ and dimethyl sulfide (DMS)) (Harvey et al., 2010; Law et al., 2011).

These early aOIF experiments resulted in demonstrated clear increases in phytoplankton biomass and PP, but the impact on association with export production (i.e., exported carbon export from the surface waters into intermediate/deep water to below the winter MLD) was not evident obscure (Fig. 3c) (de Baar et al., 2005; Boyd et al., 2007). Therefore, to determine if aOIF could increase export production, the experiment EIFEX was carried out conducted during in the closed core of a cyclonic eddy February–March 2004 in a cyclonic eddy core near the PF during the austral summer of 2004 (Fig. 5). Because it was designed to investigate the termination of a bloom and resulting export production, EIFEX was a much longer experiment (39 days) than compared to earlier experiments studies (mean \pm SD = 22 \pm 10 days; SD represents standard deviation ~28 days or less) (Smetacek et al., 2012).

Of similar duration To trace the fate of an iron-stimulated phytoplankton bloom and deep carbon export, the Indo-German iron fertilization experiment (LOHAFEX; ‘Loha’ is iron in Hindi) was conducted during January–March 2009 (40 days), also in a PF cyclonic eddy in HNLC_{LSi} waters (Smetacek and Naqvi, 2010; Martin et al., 2013).

2.1.3 Subarctic North Pacific

By the 20th century, the subarctic NP was the only HNLC region in which an aOIF experiment had not been performed (Table 1) (de Baar et al., 2005; Boyd et al., 2007). The subarctic NP shows a strong longitudinal gradient in aeolian dust deposition (i.e., high dust deposition in the west, but low in the east) (Duce and Tindale, 1991; Tsuda et al., 2003; Takeda and Tsuda, 2005), which is different from the other two HNLC regions (i.e., EP and SO). To investigate the relationship between the phytoplankton biomass/community and dust deposition, the Subarctic Pacific iron Experiment for Ecosystem Dynamics Study-1 (SEEDS-1) was conducted in July–August 2001 (13 days) in the western subarctic gyre (Tsuda et al., 2003, 2005). In 2004, the experiment was repeated (SEEDS-2) in almost the same location and season. In the intervening year, 2002, the Subarctic Ecosystem Response to Iron Enrichment Study (SERIES) was performed in July–August 2002 (25 days) in the Gulf of Alaska (representing the eastern subarctic gyre ecosystem) to compare the response of phytoplankton in this area with that in the western subarctic (Boyd et al., 2004, 2005). The SEEDS-1/2 experiments focused on changes in phytoplankton composition, vertical carbon flux, and climate-relevant gas production stimulated by artificial iron addition (Tsuda et al., 2005, 2007). The main objective of SEEDS-2 and SERIES was to determine the most significant factor (i.e., nutrient supply and/or grazing foreign) controlling the iron induced phytoplankton bloom from its beginning to its end (Tsuda et al., 2003; Boyd et al., 2004).

2.1.4 Subtropical North Atlantic

Unlike HNLC regions, PP in LNLC regions, which are predominantly occupied by N₂ fixers, is generally co-limited by phosphate and iron (Mills et al., 2004) (Mills et al., 2004). To investigate the impact of iron and phosphate co-limitation on PP, the *in situ* phosphate and iron addition experiment (FeeP) was conducted by adding both phosphate and iron in a LNLC region of the subtropical NA during April–May 2004 (21 days) (Rees et al., 2007). The location of the subtropical NA experiment corresponded to a typical LNLC region (Figs. 4a and b, Tables 3 and 4) with low nutrients (nitrate: <0.01 μ M, phosphate: ~0.01 μ M, and iron: <0.4 nM) and chlorophyll-a (<0.1 mg m⁻³) conditions much lower than other experimental

sites (~~Rees et al., 2007~~). The FeeP experiment reported that pico-plankton (0.2–2.0 μm) abundances increased after iron and phosphate additions (Rees et al., 2007); however, no other details on the biogeochemical response to ~~iron addition~~ aOIF in FeeP have been reported. This experiment will, therefore not be discussed further.

2.2 Environmental conditions prior to iron addition

The initial environment (~1–7 days before iron addition) can affect the outcome of an aOIF experiment, and the experiments described above were conducted under a wide range of physical and biogeochemical conditions. ~~Below we consider~~~~We considered~~ the similarities and differences in these environments according to the physical and biogeochemical properties of the sites (~~Steinberg et al., 1998~~; Coale et al., 1998; Steinberg et al., 1998; Bakker et al., 2001; Boyd and Law, 2001; Gervais et al., 2002; Coale et al., 2004; Boyd et al., 2005; Takeda and Tsuda, 2005; Tsuda et al., 2007; Cisewski et al., 2008; Harvey et al., 2010; Cavagna et al., 2011) (Fig. 6, Tables 3 and 4).

2.2.1 Equatorial Pacific

The first two aOIF experiments, IronEx-1 and IronEx-2, which were both conducted in the EP, were performed in different seasons (i.e., IronEx-1: October, IronEx-2: May). However, the initial surface physical conditions were ~~rather~~ similar, with warm temperatures (~~mean \pm SD =~~24.1 \pm 1.2°C; ~~SD represents standard deviation~~), high surface photosynthetic available radiation values ($\sim 51.7 \pm 2.1 \text{ mol m}^{-2} \text{ d}^{-1}$), and shallow ~~mixed layer depths (MLDs)~~ ($27.5 \pm 2.5 \text{ m}$) (Figs. 6c and d) (Coale et al., 1996; Coale et al., 1998; Steinberg et al., 1998; de Baar et al., 2005).

The initial surface biogeochemical conditions were high nutrients (i.e., nitrate = $10.6 \pm 0.2 \mu\text{M}$, phosphate = $0.9 \pm 0.06 \mu\text{M}$, and silicate = $4.5 \pm 0.6 \mu\text{M}$) and low chlorophyll-a concentrations ($0.2 \pm 0.05 \text{ mg m}^{-3}$) (Tables 3 and 4). The pico-phytoplankton (~~0.2–2.0 μm~~) community, including *Synechococcus* and *Prochlorococcus*, was dominant (Martin et al., 1994; Coale et al., 1996; Cavender-Bares et al., 1999). Initial surface nutrient concentrations were relatively low compared with other ocean basin aOIF sites (Table 3 and Fig. 6e). Initial photosynthetic quantum efficiency (i.e., Fv/Fm ratio, where Fm is the maximum chlorophyll fluorescence yield and Fv is the difference between Fm and the minimum chlorophyll fluorescence yield) (Butler, 1978), which is widely used to determine the degree to which iron is the limiting nutrient for phytoplankton growth (the Fv/Fm ratio ranges from 0.2 to 0.65 where conditions are less iron limited as Fv/Fm approaches 0.65), was less than ~ 0.3 (Fig. 6g and Table 4), suggesting severe iron limitation (Behrenfeld et al., 1996; Barber and Hiscock, 2006; Aiken et al., 2008). ~~In aOIF experiments, the initial surface partial pressure of CO_2 ($p\text{CO}_2$) has been recorded using continuous shipboard measurement systems (Wanninkhof and Thoning, 1993; Steinberg et al., 1998; Bakker et al., 2001; Bakker et al., 2005; Hiscock and Millero, 2005; Takeda and Tsuda, 2005; Smetacek et al., 2005; Wong et al., 2006; Tsumune et al., 2009; Currie et al., 2011).~~ In the EP, initial surface partial pressure of CO_2 ($p\text{CO}_2$) values were $504.5 \pm 33.5 \mu\text{atm}$, which were much higher than those observed in the SO ($355.6 \pm 11.7 \mu\text{atm}$) or the subarctic NP ($370.0 \pm 16.3 \mu\text{atm}$) (Table 3) (Steinberg et al., 1998).

2.2.2 Southern Ocean

The initial physical conditions for the aOIF experiments in the SO (SOIREE, EisenEx, SOFeX-N/S, EIFEX, SAGE, and LOHAFEX) were very different from those found in the EP; MLDs were much deeper ($57.9 \pm 19.2 \text{ m}$) (Fig. 6c) and sea surface temperature (SST) was much lower ($4.7 \pm 3.4 ^\circ\text{C}$) (Fig. 6d). During SOFeX-N/S, which were conducted along the same line of longitude, on either side of the PF, there were distinct differences in SST: 5.0°C in SOFeX-N and -0.5°C in SOFeX-S (Coale et al., 2004). SAGE was the northernmost of the aOIF experiments in the SO (Table 1) and, therefore, had

the highest SST (11.5°C) (Fig. 6d) (Harvey et al., 2010).—

The ~~locations~~regions used for the aOIF experiments were selected following preliminary surveys to confirm ~~that the sites were subject to~~ HNLC conditions, i.e., based on satellite imagery, nutrient concentrations, and Fv/Fm. ~~nitrate concentration ($>10\ \mu\text{M}$) and chlorophyll a concentration ($<1\ \text{mg}\ \text{m}^{-3}$)~~. Initial nitrate concentrations ranged from 7.9 μM (SAGE) to 26.3 μM (SOFeX-S) (Fig. 6e and Table 3). Among the various aOIF HNLC experiment sites, the SO had the highest initial nitrate concentrations (~~$21.4 \pm 5.8\ \mu\text{M}$~~ $21.2 \pm 5.8\ \mu\text{M}$), while the EP had the lowest ($10.6 \pm 0.2\ \mu\text{M}$). Initial nitrate and phosphate concentrations at aOIF sites in the SO followed a latitudinal gradient, with higher values to the south of 50°S (nitrate: $24.6 \pm 1.6\ \mu\text{M}$ and phosphate: $1.6 \pm 0.2\ \mu\text{M}$) and lower values to the north (nitrate: ~~$17.1 \pm 6.7\ \mu\text{M}$~~ $16.6 \pm 6.2\ \mu\text{M}$ and phosphate: $1.1 \pm 0.4\ \mu\text{M}$) (Table 3, Figs. 4b and 6e). The ~~SO presented the~~ full range of initial silicate concentrations has been covered by the various SO for all aOIF experiments, with values ranging ~~widely~~ from $\sim 1.0\ \mu\text{M}$ (SAGE) in the most northernmost site to $\sim 60\ \mu\text{M}$ (SOFeX-S) in the most southernmost (Table 3, Figs. 4c and 6f). With the specific intent of investigating the co-limitation of iron and silicate ~~to diatom blooms~~, SOFeX-N, SAGE, and LOHAFEX were all conducted in HNLC_{Si} regions, with initial silicate concentrations less than $2.5\ \mu\text{M}$ (Figs. 4c and 6f) (Coale et al., 2004; Harvey et al., 2010; Martin et al., 2013; Ebersbach et al., 2014). Initial $p\text{CO}_2$ values ~~were~~ lowest in the SO ($355.6 \pm 11.7\ \mu\text{atm}$) ranging from $330\ \mu\text{atm}$ (SAGE) to $367\ \mu\text{atm}$ (SOFeX-N) (Table 3).

As in the EP, initial ~~SO~~Fv/Fm values were below ~ 0.33 (Table 4 and Fig. 6g), indicating ~~a~~ severe iron limitation. Prior to iron addition, initial chlorophyll-a concentrations ranged from ~ 0.15 to $0.70\ \text{mg}\ \text{m}^{-3}$. The maximum ~~SO initial~~ chlorophyll concentrations ~~occurred in~~ were found at EIFEX, which started with a community ~~which was~~ dominated by ~~a micro-phytoplankton ($20\text{--}200\ \mu\text{m}$) community such as~~ diatoms (Hoffmann et al., 2006; Assmy et al., 2013), while the lowest initial ~~minimum~~ chlorophyll concentrations ~~occurred in~~ were observed at SOFeX-N, with a community ~~which was~~ dominated by a nano-~~phytoplankton ($2.0\text{--}20\ \mu\text{m}$) community~~, such as prymnesiophytes, pelagophytes, and dinoflagellates (Coale et al., 2004).

2.2.3 Subarctic North Pacific

The subarctic NP aOIF experiments (i.e., SEEDS-1/-2 and SERIES) were performed in regions with high nitrate ($15.6 \pm 4.0\ \mu\text{M}$) and low chlorophyll-a concentrations ($0.7 \pm 0.2\ \text{mg}\ \text{m}^{-3}$) (Tables 3 and 4, Figs. 6e and h). Compared with the other aOIF experiments, these subarctic experiments had much higher initial silicate concentrations ($27.3 \pm 9.6\ \mu\text{M}$) (Table 3 and Fig. 6f) and shallower MLDs (Fig. 6c). Although SEEDS-1 and SEEDS-2 were conducted in almost the same location and season in the western basin (Tsuda et al., 2007), the MLD in SEEDS-1 (8.5 m) was shallower than in SEEDS-2 (28 m).

Unlike the latitudinal gradients seen in the aOIF experiments in the SO, there were longitudinal gradients in physical and biogeochemical properties in the subarctic NP experiments (Tables 3–4, Figs. 4b–c, and ~~Figs.~~ 6d–h). Initial SSTs in the ~~subarctic NP~~SO were lower in the western region (7.5°C in SEEDS-1 and 8.4°C in SEEDS-2) than in the eastern region (12.5°C in SERIES) (Fig. 6d). Initial nutrient concentrations were much higher in the west (nitrate: $18.5 \pm 0.1\ \mu\text{M}$ and silicate: $34.0 \pm 2.2\ \mu\text{M}$) compared to the east (nitrate: $10\ \mu\text{M}$ and silicate: $14\ \mu\text{M}$) (Table 3, Figs. 4b–c and 6e–f). There was also a longitudinal gradient in chlorophyll-a concentrations, with relatively high values in the west (SEEDS-1: $0.8\ \text{mg}\ \text{m}^{-3}$ and SEEDS-2: $0.8\ \text{mg}\ \text{m}^{-3}$) and low values in the east (SERIES: $0.4\ \text{mg}\ \text{m}^{-3}$) (Fig. 6h). Before the first SEEDS-1 iron infusion, micro-phytoplanktons ($20\text{--}200\ \mu\text{m}$), such as the pennate diatom “*Pseudo-nitzschia turgidula*”, were dominant, whereas the areas for SERIES and SEEDS-2 were exclusively occupied by pico- and nano-phytoplankton, such as *Synechococcus* and haptophytes (Tsuda et al., 2005; Boyd et al., 2005; Tsuda et al., 2005; Sato et al., 2009). Initial Fv/Fm ratios in the subarctic NP aOIF experiments were <0.3 , indicating a severe iron limitation (Fig. 6g).

2.2.4 Subtropical North Atlantic

One exception to the focus on HNLC study sites was the FeeP experiment, which was conducted in the subtropical NA, a typically LNLC region (Figs. 4a and b, Tables 3 and 4). To test the effects of the co-limitation of iron and phosphate on PP, FeeP was conducted under much lower initial nutrient (nitrate: $<0.01\ \mu\text{M}$, phosphate: $<0.01\ \mu\text{M}$, and iron: $<0.4\ \text{nM}$) and chlorophyll *a* ($<0.1\ \text{mg}\cdot\text{m}^{-3}$) conditions than any of the other experimental sites (Rees et al., 2007).

2.3 Iron addition and tracing methods

2.3.1 Iron addition

Iron(II) and sulfate aerosols are ubiquitous in the atmosphere and, therefore, iron-sulfate ($\text{FeSO}_4\cdot\text{H}_2\text{O}$), a common form of combined iron that enters the ocean environment via dust deposition, has been frequently regarded as a bioavailable iron source during glacial periods (Zhuang et al., 1992; Zhuang and Duce, 1993; Spolaor et al., 2013). Iron-sulfate is a common inexpensive agricultural fertilizer that is relatively soluble in acidified seawater (Coale et al., 1998). Therefore, all aOIF experiments have been conducted by releasing commercial iron-sulfate dissolved in acidified seawater into the propeller wash of a moving ship (Fig. 5), to ensure mixing with surface waters during iron additions.

In general, background dissolved iron concentrations in HNLC regions are $<0.4\text{--}2\ \text{nM}$ (Table 1). Iron-enrichment bottle incubation experiments performed in deck incubators using *in situ* ocean-seawater have indicated the maximum phytoplankton growth rates in response to iron additions of 1–2 nM (Fitzwater et al., 1996). In aOIF experiments performed in the ocean, targeted iron concentrations within the MLD have ranged between ~ 1 to 4 nM, depending on the site (Table 1 and Fig. 6b) (Martin et al., 1994; Coale et al., 1996; Boyd et al., 2000; Bowie et al., 2001; Tsuda et al., 2003; Coale et al., 2004; Nishioka et al., 2005; Law et al., 2006; Tsuda et al., 2007; Harvey et al., 2010; Smetacek et al., 2012; Martin et al., 2013). If injected iron is well dispersed throughout the mixed-layer ML within 24 hours by convective mixing (Martin and Chisholm, 1992; Martin and Chisholm, 1992), the amount of added iron required to raise the background iron concentration to the target level can be calculated using a volume estimate (i.e., iron-fertilized water patch area \times MLD) (Watson et al., 1991). To minimize uncertainty between the first iron addition and phytoplankton response, aOIF experiments have involved multiple-small iron injections to the surface waters in the study area at ~ 0.4 to $\sim 1.5\ \text{km}$ intervals over a 1–2-day period (Coale et al., 1998). The patch size fertilized by the first iron addition varied from $25\ \text{km}^2$ (e.g., FeeP; Iron(II) addition of 1840 kg) to $300\ \text{km}^2$ (e.g., LOHAFEX; Iron(II) addition of 2,000 kg), and by the end of these experiments had spread to a maximum $\sim 2500\ \text{km}^2$ (Coale et al., 2004; Boyd et al., 2007; Martin et al., 2013) (Table 1, Figs. 6a and b).

During the experiments, dissolved iron concentrations increased to the target $\sim 1.0\text{--}4.0\ \text{nM}$ (Table 1 and Fig. 6b), but decreased to background concentrations within days. The fast decrease in dissolved iron concentrations indicates that iron was horizontally dispersed and/or rapidly incorporated into particles. These processes occur more rapidly in warmer waters (ACE CRC, 2015). For example, the first aOIF experiment, IronEx-1, showed that the dissolved iron concentration rapidly decreased from 3.6 to 0.25 nM $\sim 2\text{--}4$ days after iron addition in the center of the fertilized patch, suggesting a limit to the level required for phytoplankton growth (Coale et al., 1998; Gordon et al., 1998). As a result, except for the single iron addition experiments of IronEx-I, SEEDS-1, and FeeP (Martin et al., 1994; Tsuda et al., 2003; Rees et al., 2007), most aOIF experiments have involved multiple iron additions at the patch center, to continuously derive the stimulation of phytoplankton during the experiments. These experiments included: (2 additions) EIFEX, SERIES, SEEDS-2, LOHAFEX (Boyd et al., 2005; Tsuda et al., 2007; Smetacek et al., 2012; Martin et al., 2013); (3 additions) IronEx-2, EisenEx, SOFeX-N (Coale et al., 1996; Gervais et al., 2002; Coale et al., 2004; Nishioka et al., 2005); and (4 additions) SOIRE, SOFeX-S,

SAGE (Boyd et al., 2000; Coale et al., 2004; Bakker et al., 2005; Harvey et al., 2010) (Table 1).

2.3.2 Tracing iron-fertilized patch

To trace the iron-fertilized patch, aOIF experiments have used a combination of physical and biogeochemical approaches. All the aOIF experiments except EIFEX have used sulfur hexafluoride (SF₆) as a chemical tracer (Table 1) (Martin et al., 1994; de Baar et al., 2005; [Smetacek et al., 2012](#)). The SF₆, which is not naturally found in oceanic waters, is a useful tracer for investigating physical mixing and advection-diffusion processes in the ocean environment due to its nontoxicity, biogeochemically inert characteristics, and low detection limit (Law et al., 1998). The injected SF₆ is continuously monitored using gas chromatography with an electron capture detector system (Law et al., 1998; Tsumune et al., 2005). Usually only one SF₆ injection is necessary because background levels are generally extremely low in the ocean (<1.2 fM; f: femto-, 10⁻¹⁵) (Law et al., 1998; Law et al., [2006](#)[2003](#); [Martin et al., 2013](#)[Boyd et al., 2004](#)); however, in the SAGE experiment, with its higher mixing and lateral dilution, there were ~~three~~ [3](#) injections (Harvey et al., 2010). Although these earlier experiments demonstrated that the injection of artificial SF₆ is a useful technique for following iron-fertilized patches, [SF₆ can only be used for limited period \(~2 weeks\) due to the loss at the surface through air-sea gas exchange \(Law et al., 2006; Tsumune et al., 2009; Martin et al., 2013\)](#). Furthermore, caution is required because artificially high levels of SF₆ injection may negatively impact the interpretation of low-level SF₆ signals dissolved in seawater via air-sea exchange [to estimate tracer-based water mass ages for understanding physical circulation \(Fine, 2011\)](#). These techniques have been widely used to estimate anthropogenic carbon invasion as well as to understand ocean circulation in various ocean environments, with SF₆ being an important time-dependent tracer that has a well-recorded atmospheric history ~~(Fine, 2011)~~. [Thus, C](#)continuous sampling systems, measuring biogeochemical parameters such as Fv/Fm, pCO₂, and chlorophyll fluorescence, have also been used as an alternative means of following iron-fertilized patches (Gervais et al., 2002; Boyd et al., 2005; Tsuda et al., 2007; Harvey et al., 2010; Smetacek et al., 2012). The Fv/Fm ratio displays a particularly rapid increase (within 24 hours) in response to ~~ana~~ [initial-first](#) iron addition (Kolber et al., 1994; Behrenfeld et al., 1996; Smetacek et al., 2012), suggesting that it is an easy and convenient tracer for following a fertilized patch.

In addition, surface-drifting buoys equipped with ~~Array for Real-time Geostrophic Oceanography (ArgoRGO) and or Global Positioning System (GPS positioning systems) sensors~~ have been successfully used to track the movement of fertilized patches along with biogeochemical tracers (Coale et al., 1998; Boyd and Law, 2001; Law et al., 2006; Martin et al., 2013). However, floats tend to ~~drift out of the~~ [deviate from the location of](#) fertilized patches under strong wind forcing (Watson et al., 1991; Law et al., 1998; Stanton et al., 1998). NASA airborne oceanographic ~~L~~idar and ocean-color satellites have also been employed to assess the large-scale effects of iron addition on surface chlorophyll in fertilized patches, as compared to surrounding regions (Martin et al., 1994; Westberry et al., 2013).

2.4 Biogeochemical responses

Biogeochemical responses to artificial iron addition, in particular, Fv/Fm ratio, chlorophyll-a, PP, nutrients, CO₂ variables, and carbon export fluxes), are given in Tables 3–5 and Figs. 7–8. The results are important, as they have been used as a basis to determine whether the aOIF is effective. Here we address the biogeochemical response in each of the ocean basins to the aOIF experiments to date.

2.4.1 Equatorial Pacific

The IronEx-1/~~--and--~~2 experiments, which were conducted in similar initial conditions (refer to Section 2.2.1),

presented quite different biogeochemical responses (Tables 3–4 and Fig. 7). In IronEx-1, there were small increases in the Fv/Fm ratio, chlorophyll-a concentration, ~~and PP, and pCO₂ concentrations,~~ but no significant changes in nutrients ~~and pCO₂ concentrations~~ (Martin et al., 1994). On the other hand, IronEx-2 found dramatic changes in biogeochemical responses, providing support for Martin's hypothesis (Coale et al., 1996). The extremely different results from the two experiments are likely to be associated with additional iron injections (IronEx-1: no extra addition; IronEx-2: 2 additional injections) and different experimental durations (IronEx-1: 10 days; IronEx-2: 17 days).

The Fv/Fm ratios provided further detail. In IronEx-1 and IronEx-2, Fv/Fm rapidly increased within ~24 hours of iron addition and reached a maximum of ~0.60 on the second day (Table 4) (Barber and Hiscock, 2006; Aiken et al., 2008). While the elevated IronEx-1 Fv/Fm ratios promptly disappeared, ~~indicating suggesting~~ rapid iron loss due to the subduction of the fertilized patch and/or adsorption onto colloidal particles (perhaps indicative of insufficient iron supply), increased IronEx-2 Fv/Fm ratios were maintained for eight days through multiple iron additions, suggesting that additional iron enrichments are likely to be a determining factor in successfully artificially increasing PP through OIF (Kolber et al., 1994; Behrenfeld et al., 1996).

During IronEx-1, chlorophyll-a concentrations increased significantly (3-fold) reaching a maximum value of 0.65 mg m⁻³ in the first four days following iron addition (Martin et al., 1994). In IronEx-2, surface chlorophyll-a increased <27-fold with a maximum of 4 mg m⁻³ after day 7 (Table 4 and Fig. 7c) (Coale et al., 1996). To quantify the changes in carbon fixation following iron addition, the depth-integrated PP (from the surface to the critical depth, euphotic depth, or MLD) was estimated in the iron-fertilized patches. The depth-integrated PP values increased significantly compared to the initial values. The IronEx-2 ΔPP (where ΔPP = PP_{post-fertilization (postf)} – PP_{pre-fertilization (pref)}) was the highest (~1800 mg C m⁻² d⁻¹) of all the aOIF experiments discussed here (Table 4 and Fig. 7e).

Changes in pCO₂ during IronEx-1 were less than expected (ΔpCO₂ = [pCO₂]_{postf} – [pCO₂]_{pref} = -13 μatm) (Martin et al., 1994). However, substantial drawdowns of pCO₂ (ΔpCO₂ = -73 μatm) and dissolved inorganic carbon (DIC) (ΔDIC = [DIC]_{postf} – [DIC]_{pref} = -27 μM) during IronEx-2 were derived through the increased PP. The increased PP during IronEx-2 was, in turn, accompanied by drawdowns of pCO₂ (ΔpCO₂ = [pCO₂]_{postf} – [pCO₂]_{pref} = -73 μatm) and dissolved inorganic carbon (DIC) (ΔDIC = [DIC]_{postf} – [DIC]_{pref} = -27 μM) (Table 3 and Fig. 7f) (Steinberg et al., 1998). As the bloom developed, a significant nitrate uptake (e.g., ΔNO₃⁻ = [NO₃⁻]_{postf} – [NO₃⁻]_{pref} = -4.0 μM) was observed (Table 3 and Fig. 7b) and silicate concentrations also gradually decreased from 5.1 to 1.1 μM (i.e., limiting diatom growth) over eight days (Coale et al., 1996; Boyd, 2002). The depletion of macro-nutrients in fertilized patches provides indirect evidence that phytoplankton growth in surface waters is was driven by iron fertilization aOIF (Boyd and Law, 2001).

Although no phytoplankton community change was observed in IronEx-1, after iron addition in IronEx-2 ~~there was a~~ shift from a pico-phytoplankton dominated community to a micro-phytoplankton dominated community was observed, resulting in a a diatom-dominated blooms (Behrenfeld et al., 1996; Coale et al., 1996; Cavender-Bares et al., 1999). Diatom biomass increased <70-fold over eight days early in the experiment, compared to a less than a 2-fold increase for the pico-phytoplankton (Landry et al., 2000). The biomass of meso-zooplankton (200–2,000 μm), such as copepods, ~~also increased~~ grew simultaneously, substantially increasing the community grazing effect of larger animals on phytoplankton standing stocks from 7.8% d⁻¹ outside patch to 11.4% d⁻¹ in the patch (~50%) (Coale et al., 1996). However, ~~the grazing force of the increased biomass did not prevent the development of a~~ was insufficient to suppress the diatom bloom over eight days early in the IronEx-2 experiment (Table 4) (Coale et al., 1996; Rollwagen Bollens and Landry, 2000). The iron-induced diatom bloom began to decline after day ~8 of the experiment. The decline was probably associated with the combined effects of both the elevated grazing pressure and the onset of nutrient depletion (i.e., limitation in silicate and/or iron) (Cavender-Bares

et al., 1999; Boyd, 2002).

To determine whether the biological pump (i.e., export production) is enhanced after iron addition, the export flux of particulate organic carbon (POC) ~~can be~~ was estimated using ~~individually or in combination, chemical tracers such as,~~ the natural radiotracer thorium-234 (^{234}Th ; half-life = 24.1 days) ~~and the stable carbon isotope of particulate organic matter ($\delta^{13}\text{C}_{\text{org}}$), sediment traps, transmissometers, and underwater video profilers (UVPs)~~ (Table 5) (Bidigare et al., 1999; Nodder et al., 2001; Boyd et al., 2004; Buesseler et al., 2004; Coale et al., 2004; Aono et al., 2005; Tsuda et al., 2007; Smetacek et al., 2012; Martin et al., 2013). The ^{234}Th radionuclide has a strong affinity for particles, and the extent of ^{234}Th removal in the water column is indicative of the export of POC associated with surface PP out of the ~~ML mixed layer~~ (Buesseler, 1998). IronEx-2 was the first aOIF experiment in which the POC flux ~~was estimated (Bidigare et al., 1999). The ^{234}Th deficiency~~ from the surface to 25 m was measured ~~in the iron fertilized patch to estimate iron stimulated export production in the surface layer~~ (Table 5). However, no ^{234}Th measurements were made in the unfertilized patch for comparison, and no measurements in the deep ocean were undertaken to demonstrate deep carbon export (Bidigare et al., 1999).

2.4.2 Southern Ocean

As in the EP IronEx-1/-2 experiments, there were initial rapid increases in the Fv/Fm ratio within 24 hours of iron addition, indicating that phytoplankton growth was mainly limited by iron availability. Maximum values of the Fv/Fm ratio ranged from 0.5 (SOFEX-N and LOHAFEX) to 0.65 (SOIREE and SOFEX-S) (Table 4 and Fig. 7a). However, the time taken to reach the maximum Fv/Fm ratio was usually longer than ~10 days, i.e., much slower than in IronEx-1/-2 (~2 days) (Boyd and Abraham, 2001; Gervais et al., 2002; Coale et al., 2004; Smetacek et al., 2005; Peloquin et al., 2011a; Martin et al., 2013). The slower response time in the SO compared to the EP might be attributed to the colder temperatures (~5°C vs. ~24°C) and/or the deeper MLDs (~60 m vs. ~30 m) (Figs. 6c and d), ~~which were indicative of active physical mixing~~ (Boyd and Abraham, 2001; Boyd, 2002).

The aOIF experiments in the SO recorded >2-fold increases in chlorophyll-a concentrations compared to initial levels (<0.7 mg m⁻³), and maximum values between 1.25 mg m⁻³ (LOHAFEX) and ~3.8 mg m⁻³ (SOFEX-S) were obtained after artificial iron additions (Table 4 and Fig. 7c). Satellite observations were used to investigate the changing spatial and temporal distribution of chlorophyll-a concentrations in response to iron fertilization in the fertilized patches compared to the surrounding waters; for example, SOFEX-N/S found elevated chlorophyll-a concentrations in fertilized patches after iron addition through satellite images (Fig. 7d) (Boyd et al., 2000; Coale et al., 2004; ~~Boyd et al., 2005;~~ Westberry et al., 2013).

~~For example, spatial changes in chlorophyll-a resulting from SOFEX-N/S iron addition were detected using Sea-viewing Wide Field-of-view Sensor (SeaWiFS) and MODerate-resolution Imaging Spectrometer (MODIS) Terra Level 2 chlorophyll a images. The chlorophyll a image on day 24 after iron addition in the SOFEX N showed a phytoplankton bloom distribution resembling a long thread with 10-fold higher concentrations (1.0 mg m⁻³) than the surrounding waters (0.1 mg m⁻³), while a chlorophyll a image on day 20 of SOFEX S suggested a somewhat broader bloom pattern (0.5 mg m⁻³), with concentrations elevated ~5 fold over the surrounding levels (~0.1 mg m⁻³) (Fig. 7d) (Westberry et al., 2013).~~

Following artificial iron enrichment in the SO, ΔPP ranged from 360 (SAGE) to ~1356 mg C m⁻² d⁻¹ (SOFEX-N) (Table 4 and Fig. 7e). During SOIREE, EisenEx, and SOFEX-N/-S, ~~the~~ PP increased continuously throughout the duration of the experiments (Boyd et al., 2000; Gall et al., 2001a; Gervais et al., 2002; Coale et al., 2004; Assmy et al., 2007). However, in EIFEX, SAGE, and LOHAFEX there was a significant increase in PP for ~10 (SAGE) ~~to~~ 20 (EIFEX) days in response to the iron addition, and decreasing trends after day ~12 (SAGE) – 25 (EIFEX). The decrease was due to various ~~processes influences~~, such as ~~high-export production~~ (e.g., EIFEX), lateral dilution with surrounding waters (e.g., SAGE),

and high grazing pressure and ~~active~~ bacterial respiration (e.g., LOHAFEX) (Boyd, 2002; Gervais et al., 2002; Buesseler et al., 2004; Coale et al., 2004; Peloquin et al., 2011a; Smetacek et al., 2012; Thiele et al., 2012; Assmy et al., 2013; Martin et al., 2013; Latasa et al., 2014).

Using both microscopes and high-performance liquid chromatography pigment analysis, changes in phytoplankton community affected by iron addition have also been investigated. ~~Most SO aOIF iron additions experiments~~ have resulted in blooms of ~~diatoms relatively large sized phytoplankton~~ (Boyd et al., 2007). During SOIREE and EisenEx, the dominant phytoplankton community shifted from pico- and nano-phytoplankton (e.g., pico-eukaryotes and prymnesiophytes) to micro-phytoplankton (i.e., diatoms) (Gall et al., 2001a; Gervais et al., 2002; Assmy et al., 2007). In SOFeX-S and EIFEX, diatoms were already the most abundant group prior to iron addition (Coale et al., 2004; Hoffmann et al., 2006; Assmy et al., 2013). The contribution of large diatoms became especially clear in EIFEX where ~97% of the phytoplankton bloom was attributed to ~~this group these species~~ (Smetacek et al., 2012; Assmy et al., 2013). However, no taxonomic shift toward diatom-dominated ~~phytoplankton~~ communities (<5% of total phytoplankton community) was observed ~~during~~ SAGE and LOHAFEX, which were conducted under silicate-limited conditions (Harvey et al., 2010; Peloquin et al., 2011a; Martin et al., 2013; Ebersbach et al., 2014). Although SOFeX-N was conducted under low silicate conditions (Fig. 6f), the diatom biomass increased remarkably making up ~44% of the total phytoplankton community (Coale et al., 2004). This result was partly influenced by the temporary relief of silicate limitation through lateral mixing of the iron-fertilized waters with surrounding waters, with relatively higher silicate concentrations (Coale et al., 2004).

Iron-mediated increases in PP resulted in a significant uptake in macronutrients and $p\text{CO}_2$ throughout the aOIF experiments in the SO (except for SAGE) (Table 3, Figs. 7b and f). ~~The ΔNO_3^- ranged from -3.5 μM (e.g., SOFeX-S) to -1.4 μM (e.g., SOFeX-N) -1 μM (e.g., EisenEx) and $\Delta p\text{CO}_2$ ranged from -38 μatm (e.g., SOIREE) to -7 μatm (e.g., LOHAFEX). Although both initially dominated by diatoms, SOFeX-S had a somewhat greater ΔNO_3^- (-3.5 μM) and $\Delta p\text{CO}_2$ (-36 μatm) than EIFEX (ΔNO_3^- : -1.6 μM and $\Delta p\text{CO}_2$: -30 μatm) (Coale et al., 2004; Hoffmann et al., 2006; Smetacek et al., 2012; Assmy et al., 2013) both results suggested that diatoms were abundant in the two experiments. However, the smaller silicate uptake ΔSi ($\Delta\text{Si} = [\text{Si}]_{\text{postf}} - [\text{Si}]_{\text{pref}} = -4 \mu\text{M}$) observed during SOFeX-S (-4 μM) (-4 μM , compared to EIFEX (-11 μM) was associated with a decrease in silicification (i.e., changes in the adjustment of frustule in frustule thickness toward thinner frustules) of the dominant diatom species, (i.e., *Fragilariopsis* sp.) (Twining et al., 2004). During EIFEX, the ratio of heavily silicified diatoms (e.g., *Thalassiothrix antarctica*) to total diatom biomass increased from 0.24 (day 0) to 0.46 (day 37) leading to the higher Si uptake larger ΔSi (i.e., more demand for silicate) (Hoffmann et al., 2006; Assmy et al., 2013). Interestingly, the biogeochemical responses in SAGE were totally different from those seen in other experiments, in particular, as increases of increases in ΔNO_3^- (+3.9 μM), $\Delta p\text{CO}_2$ (+8 μatm), and ΔDIC (+25 μM) (Table 3, Figs. 7b and f) were observed (Table 3, Figs. 7b and f). These contrasting results were thought to be the result of entrainment through vertical and horizontal physical mixing into the iron-fertilized patch of surrounding the waters, with higher nutrient and $p\text{CO}_2$ biogeochemical concentrations (Currie et al., 2011; Law et al., 2011).~~

SOIREE was the first aOIF experiment in the SO to estimate the downward carbon flux into deep waters (Fig. 3c). ~~A They used a comprehensive suite of methods was used: such as the deployment of a~~ drifting traps, ^{234}Th and ~~the stable carbon isotope of particulate organic matter ($\delta^{13}\text{C}_{\text{org}}$) $\delta^{13}\text{C}_{\text{org}}$~~ estimates derived from high-volume pump sampling, and a beam transmissometer (Nodder and Waite, 2001). However, no measurable change in carbon export was observed in response to iron-stimulated PP (Table 5 and Fig. 8b) (Charette and Buesseler, 2000; Nodder and Waite, 2001; Trull and Armand, 2001; Waite and Nodder, 2001). During EisenEx, an increased downward carbon flux estimated from ^{234}Th deficiency was observed in the iron-fertilized patch as the experiment progressed. However, there were no clear differences between in- and

outside-patch carbon fluxes (Buesseler et al., 2005). During SOFeX-S, significantly enhanced POC fluxes below ~~the the mixed-layerMLD similar to those observed in natural blooms, were after iron enrichment-estimatedobtained~~ from ^{234}Th measurements after iron enrichment~~observations~~ (Buesseler et al., 2005). However, the absolute magnitude of these flux increases ~~was similar to that observed in natural blooms (Buesseler et al., 2005). During~~Uniquely, SOFeX-N ~~autonomous profilersused only free-profiling robotic Lagrangian carbon explorers~~ equipped with transmissometers ~~recorded ato estimate the~~ downward carbon flux ~~without employing chemical tracers, and observed large POC flux events~~ between day ~27 and ~45 after the first iron addition (Bishop et al., 2004; Coale et al., 2004). However, it was unclear whether surface-fixed carbon was well and truly delivered ~~into intermediate/deep depthsbelow the winter MLD. During~~For SAGE and LOHAFEX experiments, which were conducted under silicate limited conditions (Table 3, Figs. 4c and 6f), ~~no significant enhancement of carbon export following was detectedthere was no detection of fertilization induced export by any method~~ (Table 5) (Peloquin et al., 2011a; Martin et al., 2013). This result was likely ~~to be due to the dominance ofassociated with the pico-plankton andplankton and grazingdominated community, which that~~ led to rapid recycling ~~of organic matter in the MLmixed layer and less downward carbon flux~~. In contrast to the other aOIF experiments, EIFEX, which was conducted within the core of an eddy, ~~showedprovided~~ clear evidence of carbon export ~~well below 500 m,-depth~~ stimulated by artificial iron addition (Jacquet et al., 2008; Smetacek et al., 2012). During EIFEX, the initial export flux, estimated from ^{234}Th in the upper 100 m of the fertilized patch, was $\sim 340 \text{ mg C m}^{-2} \text{ d}^{-1}$ (Table 5 and Fig. 8a) (Smetacek et al., 2012). This value remained constant for about 24 days after iron addition. Between day 28 and 32 a massive increase in carbon export flux (maximum of $\sim 1692 \text{ mg C m}^{-2} \text{ d}^{-1}$) was observed in the fertilized patch, while the initial value remained constant in the unfertilized patch (Table 5 and Fig. 8a). The profiling transmissometer with high-resolution coverage confirmed this result, showing an increase in exported POC below 200 m after day 24. At least half the iron-induced biomass sank (via the formation of aggregates of diatom species, in particular '*Chaetoceros dictyota*') to a depth of 1,000 m, with a tenfold higher sinking rate (500 m d^{-1}), compared to the initial conditions (Smetacek et al., 2012). Significant changes in export production were not found in any of the other aOIF experiments and, therefore, the impact of artificial iron addition on this component of the biological pump needs to be resolved in future ~~a~~OIF experiments (Boyd et al., 2004; Smetacek et al., 2012; Martin et al., 2013).

2.4.3 Subarctic North Pacific

The observed increase in the Fv/Fm ratio in response to ~~artificial iron additiona~~OIF in the subarctic NP suggests that the relief in iron limitation may have assisted phytoplankton growth (Table 4 and Fig. 7a). SEEDS-1/-2, which were conducted in the western basin, showed continuous increases in the Fv/Fm ratio, with a maximum value of ~ 0.4 approximately 10 days after the first iron addition (Tsuda et al., 2003; ~~Tsuda et al.~~, 2007). During SERIES, which was conducted in the eastern basin, the Fv/Fm ratio rapidly increased and reached a maximum value of 0.55 within 24 hours of the first iron addition (Boyd et al., 2005). However, the Fv/Fm ratio returned toward the initial value of < 0.3 as the dissolved iron concentrations decreased to background levels ($< 0.2 \text{ nM}$) after about day 10 (Tsuda et al., 2003; Boyd et al., 2005; Tsuda et al., 2007).

Increases in chlorophyll-a concentrations were detected in the subarctic NP aOIF experiments in both basins after about the fifth day (Tsuda et al., 2003; Boyd et al., 2004; Suzuki et al., 2009). These increases were especially apparent in SEEDS-1, where they reached a maximum value of 21.8 mg m^{-3} (27 times the initial value of 0.8 mg m^{-3}) (Table 4 and Fig. 7c). This augmentation was the largest among all the aOIF experiments (Tsuda et al., 2003). The dramatic ~~surface~~ chlorophyll-a increase observed during SEEDS-1 was partly attributed to the particular range of seawater temperature in the region, which was conducive to diatom growth (i.e., $8\text{--}13^\circ\text{C}$) as well as to the shallower MLD ($\sim 10 \text{ m}$), which provided a relatively longer surface water residence time for the additional iron (Figs. 6c and d) (Noiri et al., 2005; Takeda and Tsuda,

2005; Tsumune et al., 2005). During SERIES, chlorophyll-a concentrations increased substantially from the initial value of 0.35 to $\sim 5 \text{ mg m}^{-3}$ over 17 days, ~~and the second highest concentration recorded in~~ all aOIF experiments ~~was recorded~~ (Table 4 and Fig. 7c) (Boyd et al., 2004). However, on the 18th day there was a downturn in chlorophyll-a as silicate concentrations decreased to $< 2 \text{ } \mu\text{M}$ (Boyd et al., 2005). Although SEEDS-2 was conducted under similar initial conditions to SEEDS-1 (refer to Section 2.2.3), there was a minimal increase in chlorophyll-a (i.e., maximum value of less than 3 mg m^{-3}) (Fig. 7c). This smaller increase was thought to be the result of ~~strong extensive~~ copepod grazing (SEEDS-2 had almost five times more copepod biomass than SEEDS-1) (Table 4) (Tsuda et al., 2007). A similar ~~range spread~~ was seen in depth-integrated PP, which increased ~~by~~ 7-fold or more after ~~various iron addition~~ ~~enrichments~~ in the subarctic NP aOIF experiments (e.g., from 300–420 to 1,000–2,000 $\text{mg C m}^{-2} \text{ d}^{-1}$) (Table 4 and Fig. 7e).

Changes in the composition of phytoplankton groups were investigated in the subarctic NP aOIF experiments. In SEEDS-1 there was a shift from oceanic diatoms (e.g., *Pseudo-nitzschia turgidula*), with growth rates of $0.5\text{--}0.9 \text{ d}^{-1}$, to faster-growing neritic diatoms (e.g., *Chaetoceros debilis*, 1.8 d^{-1}) (Tsuda et al., 2005). The shift in the dominant phytoplankton species during ~~the SEEDS-1 experiment~~ was an important contributor to ~~the what became the greatest aOIF-induced recorded~~ increase in phytoplankton biomass ~~yet recorded~~. During SERIES, the phytoplankton community changed from *Synechococcus* and haptophytes to diatoms, and the highest SERIES chlorophyll-a concentration (day 17) was associated with a peak in diatom abundance (Boyd et al., 2005). However, during SEEDS-2, no significant iron-induced diatom bloom was observed. Instead, pico-phytoplankton (e.g., *Synechococcus* ~~phytoflagellates~~) (67% of the total community) dominated throughout the duration of the experiment due to the heavy grazing pressure on diatoms (Table 4) (Tsuda et al., 2007; [Sato et al., 2009](#)).

In the subarctic NP experiments, significant ~~changes decreases~~ in macro-nutrient ~~uptake~~ ~~uptake~~ (i.e., ΔNO_3^- and ΔSi), ΔDIC , and $\Delta p\text{CO}_2$ in response to ~~artificial iron addition~~ aOIF were observed (Table 3 and Figs. 7b and f). SEEDS-1, which ~~saw exhibited~~ the largest increases in chlorophyll-a concentrations, also had the largest $\Delta p\text{CO}_2$ ($-130 \text{ } \mu\text{atm}$) and ΔDIC ($-58 \text{ } \mu\text{M}$) (Table 3 and Fig. 7f). These changes led, in turn, to the largest ΔNO_3^- ($-15.8 \text{ } \mu\text{M}$) (Fig. 7b) and ΔSi ($-26.8 \text{ } \mu\text{M}$) (Table 3) (Tsuda et al., 2003). The second largest increase in the chlorophyll-a concentration was observed in SERIES, where drawdowns of $p\text{CO}_2$ ($-85 \text{ } \mu\text{atm}$), DIC ($-37 \text{ } \mu\text{M}$), nitrate ($-8.5 \text{ } \mu\text{M}$), and silicate ($-13.6 \text{ } \mu\text{M}$) were recorded. During SEEDS-2, the nitrate concentration decreased remarkably from $18.4 \text{ } \mu\text{M}$ to $12.7 \text{ } \mu\text{M}$ after day 5; however, there was no significant change in silicate concentrations, which would have been expected as a signal of an iron-induced diatom bloom (Tsuda et al., 2007; Suzuki et al., 2009).

Despite the formation of a massive iron-induced phytoplankton bloom during SEEDS-1, there was no large POC export flux during the observation period (Table 5) (Tsuda et al., 2003; Aono et al., 2005; Aramaki et al., 2009). During SERIES and SEEDS-2, which allowed comprehensive time-series measurements of the development and decline of the iron-stimulated bloom, POC fluxes estimated by the drifting traps in the fertilized patch displayed temporal variations (Boyd et al., 2004; Aramaki et al., 2009). ~~However, the~~ The results suggested that ~~subsequently, the drifting trap captured~~ only a small part of the decrease in ~~ML the mixed layer POC in MLD was subsequently captured by the drifting trap,~~ and POC flux losses were mainly governed by bacterial remineralization and mesozooplankton grazing (Boyd et al., 2004; Tsuda et al., 2007).

2.4.4 Subtropical North Atlantic

~~Not much is known about the biogeochemical responses to OIF in the subtropical NA. The FeeP experiment reported~~

that pico-plankton abundances increased after iron and phosphate additions (Rees et al., 2007); however, no other details of the biogeochemical response to iron addition in FeeP have been reported.

2.5 Summary of the significant results from aOIF experiments

~~Artificial—The one exception was the FeeP experiment, which was performed in the subtropical NA. The initial environmental conditions associated with the physical and biogeochemical properties were determined at these OIF sites over 1–7 days before iron addition to allow the responses to the aOIF to be evaluated and quantified. Preliminary surveys confirmed that all sites, except FeeP in the subtropical NA, were subject to iron-limited HNLC conditions, with typical levels of iron <0.4 nM, nitrate >10 μM, and chlorophyll a <1 mg m⁻³. The initial Fv/Fm ratios were <0.3, suggesting that phytoplankton growth was severely iron-limited. In SEEDS-1, SOFeX-S, and EIFEX, prior to the addition of iron, the micro-phytoplankton (e.g., diatom) community accounted for half of the population and this was thought to be beneficial to the enhancement of export production. In the other experiments, pico- and nano-phytoplankton (e.g., *Synechococcus* and haptophytes) initially dominated; they are associated with rapid recycling in the mixed layer through the microbial loop rather than export production (Michaels and Silver, 1988; Coale et al., 1998; Landry et al., 2000; Boyd and Law, 2001; Gervais et al., 2002; Coale et al., 2004; Boyd et al., 2005; Tsuda et al., 2005; Hoffmann et al., 2006; Tsuda et al., 2007; Harvey et al., 2010; Martin et al., 2013).~~

~~Iron sulfates dissolved in acidified seawater have been commonly used for artificial iron addition because they are both highly bioavailable and inexpensive. The mixture is generally released into the ship's wake over a period of 24 hours. The amount of added iron was determined so to reach a target dissolved iron concentration (at least >1 nM) by volume (defined as the MLD × patch area). To achieve this, a wide range of 225–2,000 kg was applied. Except in IronEx-1 and SEEDS-1, the experiments used multiple (2–4) iron additions to reinforce the increased iron levels. To trace the iron-fertilized patches, physical tracers (i.e., ARGO or other GPS-tracked drifting buoys) and/or chemical tracers such as SF₆ were used. In addition, biogeochemical parameters, such as the Fv/Fm ratio, macro-nutrients, and CO₂ variables, were used to detect responses through a comparison of before and after conditions (i.e., $\Delta = [\text{parameter}]_{\text{postf}} - [\text{parameter}]_{\text{pref}}$). In particular, it should be noted that the Fv/Fm ratios promptly increased from <0.3 to 0.56 (±0.08) in the two days following iron addition, indicating a relief in the iron limitation on phytoplankton growth. The subarctic NP SEEDS-1 experiment, which was conducted under temperature conditions ideal for diatom growth (~8°C) and with shallow MLDs (~10 m), produced the greatest changes in biogeochemical parameters.~~

~~The aOIF experiments have generally led to changes in the size of the phytoplankton community from pico and nano-phytoplankton to micro-phytoplankton. This effect was particularly noticeable as diatoms became the dominant species during IronEx-2, SOIRE, EisenEx, SEEDS-1, SOFeX-S, EIFEX, and SERIES₁, with micro-phytoplankton accounting for ~44% of total phytoplankton community in SOFeX-N. The shift to a diatom-dominated community appears to be related to the initial availability of silicate (i.e., initial silicate was >~3 μM in all the experiments just listed). Diatom-dominated blooms induced >4.5-fold increases in chlorophyll-a concentrations and accounted for >65% of the chlorophyll-a increase (Boyd et al., 2000; Gervais et al., 2002; Coale et al., 2004; Smetacek et al., 2012). The shift to a diatom-dominated community appears to be related to initial availability of silicate (i.e., initial silicate was >~5 μM in all the experiments just listed above). However, as silicate concentrations decreased to <2 μM due to removal by phytoplankton~~the elevated diatom abundance~~, the extent of diatom blooms rapidly declined. In SAGE and LOHAFEX, ~~started with~~had low initial levels of silicate (<2 μM); As a consequence, pico and nano-phytoplankton dominated their communities, and diatom growth was limited by the lack of available silicate. However during SOFeX-N, initial silicate limitation (<~3 μM) in the iron-fertilized waters was~~

temporarily relieved through lateral mixing with the surrounding waters that had relatively higher silicate concentrations (Coale et al., 2004), which contributed to a taxonomic shift toward diatom-dominated communities (from 16% to 44% of total phytoplankton community). These results suggest that to develop ~~large-a massive~~ phytoplankton bloom, a changeover to a diatom-dominated community after iron addition is needed. A necessary, but not sufficient condition, for such a change to occur is the availability of silicate. Silicate alone is not expected to be sufficient because diatom-dominated blooms ~~with distinct increases in the chlorophyll a concentration~~ were not observed in all experiments with high initial silicate concentrations. IronEx-1 and SEEDS-2 had ~~the~~ high initial silicate levels ($>4 \mu\text{M}$), ~~considered which were~~ conducive to the development of a diatom-dominated bloom, but ~~the blooms~~ ~~were~~as suppressed due to high grazing pressure. Taken together, the aOIF results suggest that both mesozooplankton grazing rates ~~as well as and the~~ initial silicate concentrations played a role in limiting the stimulation of diatom-dominated blooms after artificial iron enrichment.

In ~~some~~ experiments ~~with smaller increases (<3 times) in plankton biomass~~ (IronEx-1, SEEDS-2, SAGE, and LOHAFEX) there was little change in the carbon export flux. ~~Among previous aOIF experiments, the subarctic NP SEEDS-1 experiment, which was conducted under temperature conditions ideal for diatom growth ($\sim 8^\circ\text{C}$) and with shallow MLDs ($\sim 10 \text{ m}$), produced the greatest changes in surface phytoplankton biomass. However, influence of iron addition on the phytoplankton growth covers from surface to euphotic depth as added iron is mixed within the ML mixed layer by physical processes (Coale et al., 1998). Although maximum surface chlorophyll-a concentration during SEEDS-1 ($\sim 22 \text{ mg m}^{-3}$) was much higher than EIFEX ($\sim 3.2 \text{ mg m}^{-3}$), the MLD-integrated chlorophyll-a concentrations were similar to $\sim 250 \text{ mg m}^{-2}$ between two experiments. Therefore, to quantify the exact changes in phytoplankton biomass in response to iron addition, it would be eligible to consider the MLD-integrated PP for comparison. During IronEx-2, SOIREE, EisenEx, SEEDS-1, SOFeX-N/-S, EIFEX, and SERIES, a >2 -fold increase in PP within the ML mixed layer, with massive diatom-dominated blooms ~~were~~as observed. However, changes in the carbon export varied substantially and differed from experiment to experiment, while in others (IronEx 2, SOIREE, EisenEx, SEEDS 1, SOFeX S, EIFEX, and SERIES) there was a >2 fold increase in PP within the mixed layer, with massive diatom dominated blooms. However, even in the latter, changes in the carbon export flux differed from experiment to experiment. In SEEDS-1 and SOIREE there was little increase in export flux. However, it has been reported that changes in in the POC concentrations following an increase in PP can take three to four weeks (Buesseler et al., 2005), whereas these ~~These~~ two experiments were conducted over only about two weeks. ~~The short, which suggests that the~~ duration of each experiment ~~could have prevented the detection of~~ was too short to detect downward carbon export. In SERIES, there was a distinct increase in the ~~carbon-POC~~ export flux within the ~~ML mixed layer~~ ($\text{MLD} = 30 \text{ m}$), but there was no increase in the ~~carbon~~ export flux below ~~this MLD~~ because the ~~produced POC was rapidly remineralized due to elevated abundance of heterotrophic bacteria~~ was elevated after iron addition rapidly remineralized ~~POC respiration~~ within the ~~ML mixed layer~~ (Boyd et al., 2004). In SOFeX-S the export flux was enhanced at ~~100-m depth m~~, below the MLD (45 m). However, the changes in export flux, after iron addition, were not dramatic compared to natural values (Buesseler et al., 2005). It is possible that the duration of SOFeX-S was also insufficient (~ 4 weeks) (Table 2). EIFEX was the only aOIF experiment that produced significant carbon export to deeper layers (down to 3,000 m). This high flux was due to aggregate formation with fast sinking rates (Smetacek et al., 2012). EIFEX observed an entire cycle (i.e., development – decline – fate) of the iron-induced phytoplankton bloom during the 39 days of the experiment, which strongly suggests that a sufficient experimental duration is a prerequisite for detecting fully formed diatom aggregates (i.e., carbon export). It should also be noted that the rates of bacterial remineralization and grazing pressure on the diatoms were in the same range inside the fertilized patch as outside, which might have assisted the delivery of iron-induced POC from the ~~MLD to the intermediate/deep depths to deep layers~~ (Smetacek et al., 2012). These results suggest that to detect significant carbon exported ~~to below the winter MLD-deep waters~~ following an increase in PP, at least three conditions are necessary: (1) a shift to a diatom-dominated community, (2) low bacterial respiration and grazing pressure rates within the ~~ML mixed layer~~, and~~

(3) a sufficient experimental duration, enabling both immediate and delayed responses to iron addition to be observed.

3 Present: Unanswered aOIF questions - export flux, possible side effects, and international law

OIF has been proposed as a potential technique for rapidly and efficiently reducing atmospheric CO₂ levels at a relatively low cost (Buesseler and Boyd, 2003), but there is still much debate. Over the past 25 years, controlled aOIF experiments have shown that substantial increases in phytoplankton biomass can be stimulated in HNLC regions through iron addition, resulting in the drawdown of DIC and macronutrients (de Baar et al., 2005; Boyd et al., 2007; Smetacek et al., 2012; Martin et al., 2013). However, the impact on the net transfer of CO₂ from the atmosphere to ~~the below the winter MLD intermediate and/or deep ocean layers~~ through the 'biological pump' (Fig. 3c) is not yet fully understood or quantified and appears to vary with environmental conditions, export flux measurement techniques, and other unknown factors (Smetacek et al., 2012). ~~There have also been a wide range of the estimates of atmospheric CO₂ drawdown resulting from large-scale and long-term aOIF based on model simulations~~ (Joos et al., 1991; Peng and Broecker, 1991; Sarmiento and Orr, 1991; Kurz and Maier-Reimer, 1993; Gnanadesikan et al., 2003; Aumont and Bopp, 2006; Denman, 2008; Jin et al., 2008; Zahariev et al., 2008; Strong et al., 2009; Sarmiento et al., 2010). While it is generally agreed that OIF effectiveness needs to be determined through ~~both tracking and quantification~~ of export fluxes, there has been no discussion ~~of about~~ which export flux measurement techniques are the most effective. ~~In the meantime~~, concern has been expressed regarding possible environmental side effects in response to iron addition (Fuhrman and Capone, 1991). These side effects include the production of greenhouse gases (e.g., N₂O and CH₄) (Lawrence, 2002; Jin and Gruber, 2003; Liss et al., 2005; Law, 2008; Oschlies et al., 2010), the development of hypoxia/anoxia in the water column (Sarmiento and Orr, 1991; Oschlies et al., 2010; Keller et al., 2014), and toxic algal blooms (e.g., *Pseudo-nitzschia*) (Silver et al., 2010; Trick et al., 2010). ~~These unwanted~~ ~~Although they are accidental, these~~ side effects could lead to negative climate and ecosystem changes (Fuhrman and Capone, 1991; Sarmiento and Orr, 1991; Jin and Gruber, 2003; Schiermeier, 2003; Oschlies et al., 2010). ~~Model studies suggested that the unintended ecological and biogeochemical consequences in response to large-scale aOIF might cancel out the effectiveness of aOIF. For example, aOIF enhanced N₂O production may have offset (up to ~40%) the benefits of CO₂ sequestration in the EP (Sarmiento and Orr, 1991; Jin and Gruber, 2003; Oschlies et al., 2010; Hauck et al., 2016).~~ Core unanswered questions remain concerning the different carbon export flux results from different measurement techniques (Nodder and Waite, 2001; Aono et al., 2005; ~~Boyd et al., 2002~~), the possible side effects that could directly influence the aOIF effectiveness, and the legal framework that is in place to ~~regulate~~ ~~protect any imprudent~~ aOIF operations while simultaneously supporting further studies to increase our understanding of ~~the potential risks and benefits of aOIF about its potential risks and benefits~~ (Williamson et al., 2012). With the design of future aOIF experiments in mind, the following section discusses the ~~following these~~ core questions: (1) which of the methods are optimal for tracking and quantifying carbon export flux, (2) which of the possible side effects have negative impacts on aOIF effectiveness, and (3) what are the international aOIF experimentation laws and can they be ignored?

3.1 Export flux measurement methods

A traditional, direct method for estimating POC export fluxes in the water column is a sediment trap that collects sinking particles (Suess, 1980). Sediment traps are generally deployed at specific depths for days to years to produce estimates of total dried mass, POC, particulate inorganic carbon (PIC), particulate organic nitrogen (PON), particulate biogenic silica (~~BSi~~), $\delta^{13}\text{C}_{\text{org}}$, and ^{234}Th . A basic assumption for the use of a sediment trap is that it exclusively collects

settling particles, resulting from the gravitational sinking of organic matter produced in surface waters. However, although they are designed to ensure the well-defined collection/conservation of sinking particles, they have accuracy issues due to: 1) interference of the hydrodynamic flow across the trap (i.e., strong advective flow), 2) inclusion/invasion (accounting for 14–90% of the total POC collected) of metazoan zooplankton (e.g., copepods, amphipods, and euphausiids) capable of vertical migration (Karl and Knauer, 1989; ~~Buesseler, 1991~~~~Buesseler et al., 1991~~; Buesseler et al., 2007), and 3) loss of trapped particles by bacterial decay and/or dissolution during trap deployment and storage periods (Gardner et al., 1983; Knauer et al., 1984; Kähler and Bauerfeind, 2001). The application of sediment traps for the determination of the carbon export flux is relatively more biased in the ~~ML upper ocean (surface to ~1,000 m depth)~~~~mixed layer~~ where ocean currents are generally faster and zooplankton are much more active than deep water. These issues suggest that sediment traps alone may not accurately determine carbon export fluxes within the ML mixed layer.

Even when used at the same depth, traditional sediment traps, such as the surface-tethered drifting trap and bottom-moored trap, can greatly over- or under-estimate particulate ^{234}Th fluxes compared to water-column based estimates (~~Buesseler, 1991~~~~Buesseler et al., 1991~~). The water column-based total ^{234}Th deficiency method (the sum of dissolved and particulate activities) is less sensitive than sediment traps to the issues mentioned above, and provides better spatial and temporal resolution in flux estimates (~~Buesseler, 1998~~~~Buesseler et al., 1998~~). For these reasons, traditional sediment trap POC flux estimates have often been calibrated using the total ^{234}Th deficiency measured using a rosette bottle or high-volume pump samples (Coale and Bruland, 1985; Buesseler et al., 2006) as a reference. However, the water column-based ^{234}Th method is sensitive to the characterization of the POC to ^{234}Th ratio on sinking particles and/or the choice of ^{234}Th flux models difference between above and below MLD was greater than a factor of 2–10 (Buesseler et al., 2006). Therefore, sampling to estimate the POC to ^{234}Th ratio should be conducted both above and below MLD to accurately detect downward carbon export flux into intermediate/deep waters.

Several aOIF experiments have used both sediment traps and ^{234}Th deficiency to estimate the iron-induced POC export flux (Table 5). SOIREE reported distinct differences in POC fluxes estimated from drifting traps ($185 \text{ mg m}^{-2} \text{ d}^{-1}$) at a ~~110-m depth~~ over day 11–13 of the experiment and ^{234}Th ($\sim 87 \text{ mg m}^{-2} \text{ d}^{-1}$) at ~~100-m depth~~ (Charette and Buesseler, 2000; Nodder and Waite, 2001). While there was no measurable change in ^{234}Th -based POC fluxes during the 13 days of the SOIREE experiment (Fig. 8b), the traps suggested a 27% increase over the course of the experiment (from 146 to $185 \text{ mg m}^{-2} \text{ d}^{-1}$) (Table 5). It was later discovered that the sediment trap-based sampling biases caused this supposed increase was caused by sampling biases in trap (Nodder et al., 2001; Nodder and Waite, 2001). Likewise, in SEEDS-1 ^{234}Th -based POC fluxes at ~~50-m depth~~ over day 9–13 were estimated to be $423 \text{ mg m}^{-2} \text{ d}^{-1}$, but the drifting trap only recorded $141 \text{ mg m}^{-2} \text{ d}^{-1}$ at ~~40-m depth~~ over day 12–14, 3 times lower (Table 5) (Aono et al., 2005; Aramaki et al., 2009). This large discrepancy between the two methods might be caused by the under-sampling of POC into the drifting traps (~~Nodder and Waite, 2001~~; Aono et al., 2005).

To resolve the potential biases in traditional sediment traps, a neutrally buoyant (and freely drifting) sediment trap (NBST) was developed (Valdes and Price, 2000; Valdes and Buesseler, 2006; ~~Lampitt et al., 2008~~). Through preliminary experiments conducted in June and October 1997 at the Bermuda Atlantic Time-series Study site, Buesseler et al. (2000) showed that an NBST system could reduce the horizontal flow and invasion/inclusion of zooplankton into the trap samplers, and that NBST-based ^{234}Th fluxes were comparable with water-column based estimates. LOHAFEX has been the only aOIF experiment so far that has measured particle export using PELAGRA (Particle Export measurement using a LAGRAngian trap) sediment traps based on the NBST system deployed at two depths of 200 m and 450 m (below the winter MLD) (Martin et al., 2013). However, the PELAGRA sediment traps deployed below the mixed layer (at 200 m and 450 m) did not detect iron fertilizationaOIF-induced carbon export even though PP did increase within the ~~mixed layer~~ML. Water-column

based ^{234}Th measurements estimated the POC flux at a 100-m depth to be $\sim 94 \text{ mg m}^{-2} \text{ d}^{-1}$, whereas the PELAGRA sediment traps estimated the flux at 200 m and 450 m to be $\sim 12 \text{ mg m}^{-2} \text{ d}^{-1}$ (Table 5) (Martin et al., 2013). It should be noted that both sediment traps and water-column based ^{234}Th measurements have a limited ability to fully scan the vertical profile of POC fluxes and, therefore, these methods should ideally be complemented with additional techniques that can measure particle stocks at high depth resolution throughout the water column ~~are of limited use in determining the fate of iron-induced POC in the water column.~~

To resolve the full column more effectively, LOHAFEX employed an underwater video profiler (UVP) ~~UVP~~, which provided photographic evidence of sinking particles (particle size $\geq 100 \mu\text{m}$) from the surface down to $\sim 3,000$ -m depth, with $\sim 0.2 \text{ m}$ vertical resolution (Smetacek and Naqvi, 2010; ~~Smetacek et al., 2010;~~ Martin et al., 2013). Through an analysis of particle size distributions, the UVP also allowed particles to be classified into fecal pellets, aggregates, and live zooplankton. Total vertical particle volume profiles obtained from the UVP indicated the a maximum particle flux concentration at a 75-m depth ($\sim 0.3 \text{ mm}^3 \text{ L}^{-1}$), with a gradual decrease to 150 m ($\sim 0.15 \text{ mm}^3 \text{ L}^{-1}$). Interestingly, large particles (i.e., zooplankton) were copious between 75-m and 100-m depth, suggesting that there might be high grazing pressure. Heavy grazing which might explain the large discrepancy between the 100 m (water-column based ^{234}Th method) and 200 m / 450 m (PELAGRA sediment trap) POC flux estimates (i.e., rather than a sampling bias in sediment trap data) (Martin et al., 2013). To continuously monitor vertical changes in POC flux stocks following iron addition, EIFEX used a transmissometer, providing high vertical resolution (~ 24 data points per meter) and tracking of the iron-induced flux stocks down to $\sim 3,000 \text{ m}$, even though, unlike UVPs, transmissometers do not allow classification of particles (Smetacek et al., 2012). Improving on this method, SOFeX-N applied autonomous carbon flux explorers equipped with transmissometers, designed to float along with the currents. Three autonomous carbon flux explorers were deployed, two explored the ‘iron fertilized in-patch’ and one acted as a ‘control’ outside the patch. Carbon flux explorers could continuously monitor carbon flux in the field for up to 18 months beyond the initial deployment, which allowed SOFeX-N to observe ‘episodic raining’ in the iron-fertilized waters (Bishop et al., 2004), indicating a high carbon export flux long after artificial iron addition. Furthermore, recent studies also reported that use of optical spike signals in particulate backscattering and fluorescence, measured from autonomous platforms such as gliders and floats, can provides high-resolution observations of POC flux (Briggs et al., 2011; Dall’Omo and Mork, 2014).

The combination of multiple approaches is essential to the successful detection of the POC produced in response to iron addition and its fate. NBST systems (e.g., the PELAGRA sediment trap) are appropriate to should be deployed at two depths (i.e., below both the in situ MLD and the winter MLD) for to quantifying the aOIF-induced POC flux. mixed layer in the upper waters (< 400 m depth); This technique is improved, especially when accompanied by calibration using water-column based ^{234}Th . Particle profiling systems (e.g., a transmissometer and an UVP) can mounted on a CTD Rosette sampler provide continuous quantitative and qualitative information about sinking particles, with high vertical resolution and full coverage of the water column ($> 3,000$ -m depth). They are therefore useful for indirectly identifying deep carbon transport. Autonomous carbon flux explorers are an excellent alternative, allowing the for continuous observation of POC fluxes during and after an aOIF experiment.

3.2 Considering environmental side effects

The purpose of aOIF is to reduce the atmospheric CO_2 level by stimulating the sequestration of oceanic carbon through artificial iron additions in the HNLC regions, mitigating the global warming threat. Beyond the benefits of aOIF

experimentation, scientists have debated the unintended secondary consequences of aOIF, such as production of climate-relevant gases and ocean ecosystem changes. Therefore, it is important to consider the possible negative consequences of aOIF to evaluate whether the aOIF experiments are effective (i.e., net profit: positives > negatives).

To investigate changes in climate-relevant gas emissions produced by biological activities and/or photochemical reactions before and after iron additions, the production of CH₄, N₂O, DMS, and halogenated volatile organic compounds (HVOCs) ~~were~~^{was} measured during aOIF experiments (Liss et al., 2005), because their emission may lead to unintended consequences negating the desired effects of aOIF experiments on carbon sequestration. Among the climate-relevant gases, CH₄ has a ~20 times greater warming potential than CO₂ (IPCC, 1990). However, CH₄ has been considered to be relatively low risk because most of the CH₄ formed in the ocean is used as an energy source for microorganisms and is converted to CO₂ before reaching the sea surface (Smetacek and Naqvi, 2008; Williamson et al., 2012). The SO nOIF experiment conducted in 2011 year (i.e., Kerguelen Ocean and Plateau compared Study-2: KEOPS-2) (Table 1) showed that CH₄ concentrations were 4-fold higher in the naturally iron-fertilized patch than in the control area (FariásFariás et al., 2015). During the SOFeX-N experiment, measurements of dissolved CH₄ indicated concentrations were slightly elevated, i.e., by less than 1% (1.74 ppmv in fertilized patch and 1.72 ppmv outside fertilized patch) (Wingenter et al., 2004). Simulated SO large-scale aOIF has suggested that a 20% enhancement of CH₄ emissions would offset only <1% (~4 Tg C yr⁻¹) of the resulting carbon sequestration (Oschlies et al., 2010). Hence, additional CH₄ production from aOIF experiments is not likely to be ~~a serious problem~~^{significant}.

On the other hand, N₂O has a relatively long lifetime in the atmosphere (~110 years) and has a global warming potential about 300 times greater than CO₂ (Forster et al., 2007). The ocean is already a significant source of atmospheric N₂O (Nevison et al., 2003; Bange, 2006). Oceanic N₂O is mainly produced by bacterial remineralization. Therefore, increases in N₂O production after iron additions are expected and, in the long run, contribute to an increase rather than a decrease in the greenhouse effect (Fuhrman and Capone, 1991Bange, 2006). During the SOIREE experiment, a significant increase (~4%) in mean N₂O saturation in the pycnocline (65–80 m) of the fertilized patch (104.4 ± 2.4%), as compared to outside the fertilized patch (100.3 ± 1.7%), was associated with an increased phytoplankton biomass (Law and Ling, 2001). Measurements of N₂O saturation during SERIES also showed increases of 8% at 30–50 m, which were coincident with the accumulation of ammonium and nitrite attributable to increases in bacterial remineralization following increased POC levels (Boyd et al., 2004; Law, 2008). SOIREE-based model estimates suggested that potential N₂O production at timescales longer than six weeks would subsequently offset carbon reduction benefits resulting from the bacterial remineralization of additional carbon fixation by 6–12% (Law and Ling, 2001). This estimate is in line with the N₂O offset of 6–18% suggested by a modeling study (Jin and Gruber, 2003) and the 5–9% suggested by a more recent modeling study investigating the effectiveness of long-term and large-scale SO aOIF (Oschlies et al., 2010). However, the SO nOIF experiment (i.e., KEOPS-2) suggested that nOIF acted as both a sink and a source for N₂O (FariásFariás et al., 2015). Excess N₂O was not found after iron addition in EIFEX, where significant vertical export through the formation of rapidly sinking aggregates was found (Walter et al., 2005; Law, 2008). One explanation for the absence of N₂O accumulation below the EIFEX patch might be the limited bacterial remineralization due to the rapid export of organic matter well below the 500-~~m-depth~~ to the seafloor (Law, 2008Walter et al., 2005). Based on the results of previous studies, no consensus has yet been reached on the exact extent of additional N₂O production after iron additions. However, because there is the potential for excessive N₂O production that would not only impact the effectiveness of aOIF experiments but also positively contribute to global warming, further studies are required to reach a conclusion.

Unlike N₂O emissions, which have the potential to offset the effectiveness of aOIF, DMS, a potential precursor of

sulfate aerosols that cause cloud formation, may contribute to the homeostasis of the earth's climate by countering the warming due to increased CO₂ emissions (Charlson et al., 1987). DMS is produced by the enzymatic cleavage of planktonic dimethylsulfoniopropionate (DMSP). Microzooplankton grazing on nano-phytoplankton (e.g., haptophytes) is a key factor controlling oceanic DMS production (Dacey and Wakeham, 1986; Gall et al., 2001b; Park et al., 2014). The production of DMS in response to iron addition was measured during all aOIF experiments. In the EP and SO, DMS production increased, but in the subarctic NP, it remained constant or decreased (Lawrence, 2002; Boyd et al., 2007; Law, 2008). There were significant short-term increases in DMS production in IronEx-2 (from 2.5 to 4.2 nM), SOIREE (from 0.5 to 3.4 nM), EisenEx (from 1.9 to 3.1 nM), and SOFeX-N (7.7 nM in the fertilized patch and 1.6 nM outside the fertilized patch) (Turner et al., 1996; Turner et al., 2004; Wingenter et al., 2004; Liss et al., 2005; Wingenter et al., 2007). The maximum DMS production observed was a 6.8-fold increase after iron addition in SOIREE (Turner et al., 2004). During an early SOIREE experiment, the dominant phytoplankton species were haptophytes, and DMS production was increased by microzooplankton grazing on DMSP-rich haptophyte species (i.e., Prymnesiophyceae) (Gall et al., 2001b). Similarly, a 4.8-fold enhancement of DMS production was observed in SOFeX-N. Estimates derived by the extrapolation of SOFeX-N DMS production results suggested that fertilizing ~2% of the SO area over the course of a week would derive a 20% increase of the total SO DMS flux, enhance DMS production by 20%, which would lead to a 2°C decrease in air temperature over the SO (Wingenter et al., 2007). On the other hand, the SO nOIF experiment (KEOPS-1) conducted in 2005 year (Table 1) showed that DMS production was not markedly higher in the naturally fertilized area compared to the surrounding waters (Belviso et al., 2008). Twenty years simulated SO aOIF did not produce accumulation of DMS in surface waters (Bopp et al., 2008). Interestingly, there were no significant changes in DMS production after iron additions in the western subarctic NP SEEDS-1/-2 experiments, despite increases in PP (Turner et al., 1996; Takeda and Tsuda, 2005; Nagao et al., 2009). Furthermore, in the eastern subarctic NP, SERIES DMS production increased from 8.5–10.9 nM on day 1 to a maximum of 41.2 nM on day 10, but decreased to <0.03 nM by the end of the experiment due to an increase in bacterial abundance (Table 4) (Levasseur et al., 2006). It is therefore difficult to predict the iron-induced DMS response, because OIF itself is not the only source of DMS. Based on the results of previous aOIF experiments, DMS production was sensitive in the EP and SO, but was less sensitive in the subarctic NP (Law, 2008). These results indicate that further process and modeling studies for each region are required to determine the production and degradation of DMS, both following iron fertilization and in the natural environment.

HVOCs, such as CH₃Cl, CH₃Br, and CH₃I, are well known for their ability to destroy ozone in the lower stratosphere and marine boundary layer (Solomon et al., 1994), and were also measured during past aOIF experiments (Wingenter et al., 2004; Liss et al., 2005). However, no consistent results have been reported for HVOCs production (Liss et al., 2005). In SOFeX-N, the impact of iron addition on HVOCs was complicated, with CH₃Cl concentrations remaining unchanged, and CH₃Br concentrations increasing by 14% (6.5 pptv in the fertilized patch and 5.7 pptv outside the fertilized patch), while CH₃I concentrations decreased by 23% (4.9 pptv in fertilized patch and 6.4 pptv outside the fertilized patch) (Wingenter et al., 2004). In contrast, CH₃I concentrations increased ~2-fold during EisenEx (Liss et al., 2005). Such a complicated response suggests that, as for DMS, further study is needed to fully understand natural cycling of HVOCs and their responses to iron fertilization aOIF.

Secondly, Another important consideration is the extent to which the effectiveness of aOIF may also be cancelled out by its tendency to offset leading to changes in the ocean ecosystem following aOIF, changes such as a decrease in dissolved oxygen and an increase in domoic acid (DA) levels. The decomposition of iron addition-enhanced biomass may cause decreased oxygen concentrations in subsurface waters (Williamson et al., 2012), but Although mid-water oxygen depletion has not been reported from aOIF experiments to date (Williamson et al., 2012), Early modeling studies suggest that anoxic

conditions may develop after long-term, ~~and~~ large-scale aOIF (Fuhrman and Capone, 1991; Sarmiento and Orr, 1991). However, ~~more sophisticated and realistic models suggest that OIF produces whereas a recent study based on more sophisticated models showed sustained~~ well-oxygenated conditions ($O_2 \approx 12047 \mu M$) even under simulated aOIF south of $30^\circ S$ on a 100 year timescale from 2010 to 2110, ~~without the development of anoxic conditions, even under climate change scenarios, without the development of anoxic conditions, even under climate change scenarios~~ (Oschlies et al., 2010; Keller et al., 2014). Thus, hypoxia/anoxia development in response to iron additions is unlikely to be a primary concern. Keller et al. (2014) found ~~that simulated SO large-scale aOIF south of $40^\circ S$ from the year 2020 to 2100 under a high CO_2 emissions scenario (Meinshausen et al., 2011) may develop suboxia ($O_2 < 10 \mu M$) in the year 2125. Clearly, the circumstances under which a substantial decline in oxygen inventory can be caused by large-scale aOIF need further study.~~

The changes in phytoplankton community composition after iron addition discussed in Section 2.4 may also have unintended consequences. ~~For example, they such changes~~ could lead to potentially toxic species dominating plankton assemblages (Silver et al., 2010; Trick et al., 2010). Some aOIF experiments (e.g., IronEx-2, SOIREE, EisenEx, SOFeX-N/S, and SERIES) generated large blooms dominated by pennate diatoms belonging to the genus '*Pseudo-nitzschia*' (de Baar et al., 2005; ~~Silver et al., 2010; Trick et al., 2010~~). Some '*Pseudo-nitzschia*' species have the capacity to produce the neurotoxin DA that ~~canis known to~~ detrimentally affect marine ecosystems. However, no DA was found during EisenEx and SERIES, even though '*Pseudo-nitzschia*' were dominant (Gervais et al., 2002; Assmy et al., 2007; Marchetti et al., 2008; Assmy et al., 2007). ~~However, p~~Phytoplankton samples used to estimate DA production have ~~sometimes~~ been stored for a long time before the analysis, for example, 12 years in IronEx-2 and four years in SOFeX-S (Silver et al., 2010). Trick et al. (2010) argued that ~~storage might have affected the DA content in the samples, which led to phytoplankton samples stored for a long time would have degraded, leading to~~ an under-estimation in DA ~~concentrations~~production. This implies that accurate information about changes in DA production in response to iron addition might not be available. ~~However~~ Nevertheless, the IronEx-2, and SOFeX-S experiments found discernable changes in DA production; ~~were found even if the original DA might have degraded in IronEx-2; and SOFeX-S experiments~~ (Silver et al., 2010). It is likely that ~~detection was possible because these several phytoplankton samples were (e.g., *Pseudo-nitzschia* abundance: 1.3×10^6 cells L^{-1} in IronEx-2 and 7.5×10^4 cells L^{-1} in SOFeX-S)~~ collected with a-net tows (20- to 30- μm mesh phytoplankton nets), which ~~were suitable to detect these changes provided concentrated samples of larger phytoplankton including *Pseudo-nitzschia* (e.g., *Pseudo-nitzschia* abundance: 1.3×10^6 cells L^{-1} in IronEx-2 and 7.5×10^4 cells L^{-1} in SOFeX-S).~~ During IronEx-2 and SOFeX-S, high cell abundances of '*Pseudo-nitzschia*' (10^6 and 10^5 cells L^{-1} , respectively) combined with moderate DA cell quotas (0.05 and 1 pg DA cell $^{-1}$, respectively) produced toxin levels as high as 45 ng DA L^{-1} in IronEx-2 and 220 ng DA L^{-1} in SOFeX-S in the water, respectively, i.e., toxin levels high enough to damage marine communities in coastal waters (Scholin et al., 2000; Schnetzer et al., 2007). Trick et al. (2010) suggested that large-scale iron fertilizations OIF may induce DA accumulation with developing toxic *Pseudo-nitzschia* blooms. However, large uncertainties remain as Trick et al. (2010) simply extrapolated DA concentration based on bottle incubation experiments with HNLC surface waters to the DA production expected from large-scale iron fertilization OIF. ~~Therefore~~ As a result, it is necessary to clarify/quantify DA production in response to iron additions aOIF, with concentrated larger phytoplankton samples (i.e., large numbers of cells) using collected using a-net tows (20- to 30- μm mesh phytoplankton net). This, ~~once~~ Here again, existing research indicates that ~~such the~~ processes involved need to be better understood in the natural environment before the ramifications of aOIF can be fully ~~appreciated~~ understood.

Whether aOIF is a viable carbon removal strategy is still under debate (Boyd et al., 2007; Smetacek and Naqvi, 2008). The production of climate-relevant gases such as N_2O , DMS, and HVOCs, which is influenced by the remineralization of sinking particles that follows OIF-induced blooms, the decline in oxygen inventory, and the production of DA are particularly important to understand. These processes can directly and indirectly modify the effectiveness of carbon

sequestration, with ~~either positive or negative the effects being either positive or negative~~. Therefore, monitoring ~~of the declines in oxygen content and~~ production of climate-relevant gases and DA to evaluate the effectiveness of aOIF as a geoengineering approach is essential. ~~The processes isdiscussed here represents only a few of many possiblethe current state of knowledge concerning aOIF~~ side effects. The direct and indirect environmental consequences ~~of aOIF~~ remain largely unresolved due to the inconsistent and highly uncertain outcomes of the experiments conducted so far, as well as our poor understanding of the processes involved under both nOIF and aOIF conditions (Chisholm et al., 2001; Johnson and Karl, 2002; Williamson et al., 2012). Therefore, considering the increasing evidence for the necessity to keep warming at or below 1.5°C (Rogelj et al., 2015), there continues to be a need to quantitatively determine the effectiveness of aOIF as a long-term means for reducing atmospheric CO₂ through the quantification of aOIF side effects.

3.3 Regulation of aOIF: International law of the sea as it applies to aOIF

To prevent pollution of the sea from human activities, the international Convention on the Prevention of Marine Pollution by Dumping of Wastes and Other Matter (London Convention, 1972) was amended in 1972. In 1996, contracting parties to the London Convention adopted the Protocol to the London Convention (London Protocol, 1996). This places legal restrictions on the dumping of wastes and other matter that may cause hazard, harm, and damage in the ocean and/or interfere with the marine environment. However, the London Convention & Protocol (LC/LP) did not establish specific laws to protect the ocean environment against the side-effects of fertilization activities. In 2007, several commercial companies (e.g., GreenSea Venture [<http://www.greenseaventure.com>] and Climos [<http://www.climos.com>]) promoted large-scale (10,000 km²) commercial aOIF as a climate mitigation strategy and as a means to gain carbon credits (Chisholm et al., 2001; Buesseler and Boyd, 2003; Freestone and Rayfuse, 2008). Meanwhile, assessments of the effectiveness of aOIF have been limited to small fertilized patches (25–300 km²) (Fig. 6a) due to the time and expense of comparing fertilized and unfertilized areas (ACE CRC, 2008). As discussed earlier, these small-scale ~~area~~ experiments have left many unanswered scientific questions regarding both the effectiveness and the potential impacts of aOIF (Lawrence, 2002; Buesseler and Boyd, 2003). In the same year, noting the potential risks and benefits, the LC/LP scientific group released a statement on large-scale ocean fertilization and recommended that ocean fertilization activities be evaluated carefully to ensure that such operations were not contrary to the aims of the LC/LP.

At the 2008 LC/LP meeting, the contracting parties adopted Resolution LC-LP.1 (2008) on the regulation of ocean fertilization. This resolution prohibited ocean fertilization activities until such time that specific guidance could be developed to justify legitimate scientific research. There was an exception for ‘small-scale scientific research studies within coastal waters’ to permit the development of proposals that would lead to an assessment framework for scientific ocean fertilization research (Resolution LC-LP.1, 2008). In the meantime, there was a call to develop an assessment framework for ocean fertilization experiments to assess, accurately, scientific research proposals (Resolution LC-LP.1, 2008). In 2010, LC/LP parties developed Resolution LC-LP.2 (2010), adopting an “Assessment Framework for Scientific Research Involving Ocean Fertilization” to be used to assess, on a case-by-case basis, whether any proposed ocean fertilization activity constitutes legitimate scientific research falling within the aims and scope of Resolution LC-LP.1 (2008) (Fig. 9) (Resolution LC-LP.2, 2010). This framework demands preliminary scientific research prior to any aOIF experimentation. There must be a transparent/reasonable scientific rationale/purpose to the experiment and a risk analysis must be undertaken using parameters such as problem formulation, site selection, exposure and effect assessment, and risk characterization and management. Monitoring is also required as an integral component of all approved (i.e., legitimate) scientific aOIF research activity to assess ecological impacts and to review actual vs. intended geo-engineering benefits (ACE CRC, 2015). In October 2013,

the LC/LP parties adopted amendments that categorize aOIF as marine geo-engineering, thereby prohibiting operational aOIF activities, but enabling aOIF scientific research that meets the permit conditions through the environmental assessment framework (Resolution LP.4 (8), 2013). This means that large-scale (i.e., >300 km² based on previous aOIF experiments; exact areal sizes are not determined in the LC/LP) and/or commercial aOIF (e.g., ‘the 2012 Haida Gwaii Iron Dump’ off the west coast of Canada) are currently banned by international regulations. Under LC/LP, commercial aOIF efforts cannot proceed because of the large uncertainties related to large-scale aOIF.

4 Future: Designing future aOIF experiments

Scientific aOIF research has focused on improving our understanding of the effectiveness, capacity, and risks of OIF as an atmospheric CO₂ removal strategy both in the future and the past (in particular glacial periods). Although the first aOIF experiments took place more than twenty years ago, the legal and economic aspects of such a strategy in terms of the international laws of the sea and carbon offset markets are not yet clear (ACE CRC, 2015). Nonetheless, previous small-scale aOIF experiments have demonstrated a considerable potential for easily and effectively reducing atmospheric CO₂ levels. Accordingly, physical/biogeochemical/ecological models and nOIF experiments (long-term) have been conducted in an effort to overcome some of the limitations of short-term aOIF experiments (e.g., spatial and temporal scales) and to predict the effectiveness of long-term and large-scale fertilization (Aumont and Bopp, 2006; Blain et al., 2007; Denman, 2008; Pollard et al., 2009; Sarmiento et al., 2010). For example, earlier global biogeochemical models have indicated that massive fertilization could draw down atmospheric CO₂ by as much as 107 µatm in 100 years (Joos et al., 1991; Peng and Broecker, 1991; Sarmiento and Orr, 1991; Kurz and Maier-Reimer, 1993). Recent global models, with more realistic ecosystem and biogeochemical cycles predict values closer to a 33 µatm drawdown in atmospheric CO₂ (Aumont and Bopp, 2006). These results suggest that the amount of carbon sequestration resulting from aOIF represents only a modest offset, i.e., a contribution of ~10% over the range of IPCC future emission scenarios (IPCC, 2000; Aumont and Bopp, 2006; Denman, 2008; Zahariev et al., 2008). The nOIF experiments have also produced much higher carbon sequestration rates than the small-scale aOIF experiments (Morris and Charette, 2013). Furthermore, the results from nOIF experiments do not support the potential negative impacts proposed for OIF experiments, even at larger scales (Belviso et al., 2008). However, these nOIF results do not guarantee that aOIF as a geoengineering approach is able to achieve the high effectiveness associated with carbon sequestration and enables a simple scaling-up as a prediction tool, because the nOIF experiments differ from the aOIF experiments in the mode of iron supply. In particular, nOIF is a continuous and slow process and its iron source is based on the upwelling of iron-rich subsurface waters to the surface layer, whereas aOIF is intended to be episodic, with massive short-term iron additions (Blain et al., 2007). In addition, in nOIF it is difficult to accurately identify iron sources due to the complexity of the system, whereas in aOIF there is quantitative and qualitative information about iron additions and sources (Blain et al., 2008). Contrary to the results of aOIF experiments in the SO (e.g., SOIREE and SOFeX-N), no increase in DMS emissions was found in SO nOIF experiments in the SO (i.e., ~~Kerguelen Ocean and Plateau compared Study-KEOPS-1~~) (Belviso et al., 2008), suggesting that it might be difficult to identify the potential long-term negative effects of aOIF ~~through the by study ofing the~~ naturally fertilized systems, at least in the SO. Therefore, it is important to continue undertaking small-scale studies to obtain a better understanding of natural processes in the SO as well as to assess the associated risks, and so lay the groundwork for evaluating the potential effectiveness and impacts of large-scale aOIF as a geoengineering solution to anthropogenic climate change. It is therefore of paramount importance that future aOIF experiments continue to focus on the effectiveness and capacity of aOIF as a means of reducing atmospheric CO₂, but they should also carefully consider the location (i.e., ‘where’), timing (i.e., ‘when’), and duration (i.e., ‘how long’), as well as modes of iron addition (i.e., ‘how’), tracing methods/parameters measurements/protocols (i.e., ‘what’), and side effects on

marine/ocean ecosystems (i.e., ‘what concerns’). They ~~is should will~~ build on the results of previous aOIF experiments ~~to~~ develop our understanding of the magnitude and sources of uncertainties, and provide confidence in our ability to reproduce results.

Where: The first consideration for a successful aOIF experiment is the location. The dominance of diatoms in phytoplankton communities plays a major role in increasing the biological pump because diatom species can sink rapidly as aggregates or by forming resting spores ~~to efficiently bypass the intense grazing pressure of mesozooplankton (e.g., copepods, salps, and krill) and export carbon out of the winter ML mixed layer~~ (Tréguer et al., 1995; Salter et al. 2007; Assmy et al., 2013; Rembauville et al., 2015; Rembauville et al., 2016). Previous aOIF experiments have shown that silicate concentration ~~and mesozooplankton stocks (i.e., copepods) is are~~ the crucial factors ~~inducing-controlling~~ diatom blooms (Boyd et al., 2000; Gervais et al., 2002; Coale et al., 2004; Tsuda et al., 2007; Smetacek et al., 2012). Therefore, to obtain the greatest possible carbon export flux in response to iron addition, aOIF experiments should be designed in regions with high silicate concentrations ~~and low grazing pressure. It will be important to conduct initial surveys onto measure the degree of grazing pressure in HNLC region with high silicate concentrations,~~ such as in the subarctic NP (e.g., SEEDS-1 experiment) and the south of ~~SO PF-of the SO~~ (e.g., SOFeX-S experiment) $>15 \mu\text{M}$ (Fig. 4c). In selecting sites for ~~iron fertilization aOIF,~~ it is also important to distinguish the iron-fertilized patch from the surrounding unfertilized waters to ~~easily and efficiently~~ observe; ~~easily and efficiently,~~ iron-induced changes (Coale et al., 1996). Ocean eddies provide an excellent setting for aOIF experimentation because they tend to naturally isolate interior waters from the surrounding waters. Mesoscale eddies range from 25–~~to~~ 250 km in diameter and maintain their characteristics for 10–100 days after formation (Morrow and Le Traon, 2012; Faghmous et al., 2015). Eddy centers, ~~in which fertilization is performed,~~ tend to be subject to relatively slow current speeds, with low shear and high vertical coherence, providing ideal conditions for tracing the same water ~~column~~ from the surface to ~~the deep layers below the winter MLD, as well as while simultaneously~~ minimizing lateral stirring and advection (Smetacek et al, 2012). ~~Therefore, f~~Finding an appropriate eddy setting in a study area should be a high priority consideration when ~~condue~~ designing an aOIF experiment (Smetacek and Naqvi, 2008).

When: The second consideration for a successful aOIF experiment is timing, which includes when an experiment starts. PP in ocean environment is generally limited by nutrient availability and/or by light availability, often referred to as a single- or co-limitation. PP in the SO, a representative HNLC region, is subject to co-limitation by micro/macro-nutrients (i.e., iron and/or silicate) and light availability (Mitchell et al., 1991). ~~In-To~~ the south ~~of the SO PF-of SO,~~ phytoplankton blooms usually occur during ~~the~~ early summer (i.e., from late December to early January) due to ~~an the~~ increasing in the nutrient flux from ~~the~~ subsurface waters ~~induced by winter mixing to the surface waters by the shoaling of MLD,~~ along with the favorable light conditions provided by a shoaling of ML ~~the receipt of sufficient solar radiation~~ (Moore and Abbott, 2002). Prior to December, phytoplankton growth is mainly limited due to light availability (Mitchell et al., 1991; Veth et al., 1997; Abbott et al., 2000), while after January (i.e., during late summer and early autumn from February to March) it is mainly limited due to iron and silicate availability (Abbott et al., 2000; Mengelt et al., 2001; Nelson et al., 2001). In previous SO aOIF experiments ~~in the SO that have been~~ conducted between spring and early autumn, PP was mainly limited by iron and/or silicate availability rather than light availability (except when heavy clouds led to severe light limitation, only occurred for a few days during EisenEx) (Gervais et al., 2002; Bakker et al., 2005; de Baar et al., 2005; Smetacek and Naqvi, 2008; Peloquin et al., 2011b). ~~T~~In addition, the grazing pressure of mesozooplankton ~~(i.e., copepods)~~ on large diatoms was also a major limiting factor in diatom production (~~Coale et al., 2004; Schultes et al., 2006; Smetacek and Naqvi, 2010; Martin et al., 2013,~~ and was generally higher during late summer and early autumn (February to March) (~~Hunt and Hosie, 2006; Rembauville et al., 2015; Le Quéré et al., 2016.~~ Considering the key factors (i.e., micro/macro nutrient availability, light availability, and grazing pressure) controlling PP in the SO, the most appropriate timing for the start of an SO aOIF

experiment ~~to start in the SO~~ is likely to be the early summertime (i.e., late December to early January).

How long: The third consideration for a successful aOIF experiment is the duration. Although the periods that phytoplankton blooms have been maintained by aOIF have lasted from ~10 to ~40 days (Kolber et al., 1994; Martin et al., 1994; Coale et al., 1996; Boyd et al., 2000; [Boyd et al., 2004](#); [Coale et al., 2004](#); Tsuda et al., 2005; ~~Coale et al., 2004~~; ~~Boyd et al., 2004~~; Smetacek et al., 2012), it has also been suggested that most aOIF experiments did not cover the full response times from onset to termination (Boyd et al., 2005). For example, SOIREE and SEEDS-1 had relatively short observation periods (13 days) and saw increasing trends in PP throughout the experiments (Fig. 10a), suggesting that the observation period should have been extended. Furthermore, after the end of SOIREE, ocean color satellite images showed continued high chlorophyll-a concentrations ($>1 \text{ mg m}^{-3}$) in the iron fertilized patch, which was visible as a long ribbon shape that extended some 150 km for >40 days (~6 weeks) after the ~~initial-first~~ iron addition (Fig. 10b) (Abraham et al., 2000; Westberry et al., 2013). This indicates that short experimental durations may not be sufficient for detecting the full influence of ~~artificial iron additions~~ aOIF on PP and ecosystem (Figs. 8b and 10) (Boyd et al., 2000; Tsuda et al., 2003; de Baar et al., 2005). SOFeX-S also resulted in relatively low export production despite the high PP due to the experimental duration being insufficient to cover the termination of the phytoplankton bloom. However, SERIES, SEEDS-2, EIFEX, and LOHAFEX did fully monitor all phases of the phytoplankton bloom from onset to termination. EIFEX, the third-longest aOIF experiment, at 39 days, was the only one that observed iron-induced deep export production between day 28 and 32 (Table 5 and Fig. 8a) (Smetacek et al., 2012; Assmy et al., 2013). Furthermore, long-term observations covering the later stage of bloom development during nOIF experiments resulted in much higher ~~Fe:C:Fe~~ export efficiencies compared to the short-term aOIF (Blain et al., 2007; Pollard et al., 2009). Based on previous aOIF experiments, it would, therefore, be important to detect the full phase of a phytoplankton bloom to determine accurately the amount of iron-induced POC exported out of the ~~winter ML mixed layer~~. The observation period is, therefore, an important ly considering factor ~~to consider with regard to-in~~ budget and effectiveness estimates. It is suggested that the experimental duration should be a minimum of ~40 days based on the SOIREE experiment, which produced the longest iron-induced bloom (i.e., ~~the longevity of~~ >40 days). In addition, autonomous observation platforms are essentially required to monitor post-assessment of effectiveness, capacity, and risks of aOIF for at least 12 months after the experiment termination.

How: The fourth consideration for a successful aOIF experiment lies in the strategy/approach of adding and maintaining dissolved iron within the ~~ML upper mixed layer~~ to produce a phytoplankton bloom. First, the chemical form for iron addition should be acidified iron-sulfate, which is less expensive and more bioavailable than other iron compounds. The amount of iron-sulfate required is calculated according to the target concentration of the dissolved iron and volume ($\text{MLD} \times \text{patch size}$). Based on bottle incubation experiments, target iron concentrations of ~2–4 nM are recommended to stimulate maximum phytoplankton growth due to the rapid losses of added iron by horizontal advection/diffusion and oxidation to poorly bioavailable iron(III) (Coale et al., 1996; Coale et al., 1998; Bowie et al., 2001). For patch size, a biogeochemical model study showed that a fertilized patch size of 156 km^2 maintained an iron concentration above 0.3 nM for 56 days, while a longer period of 194 days required a fertilized patch size of $160,000 \text{ km}^2$ (Xiu and Chai, 2010). ~~This is because, compared to larger iron fertilized patches, a smaller patch size tended to lose iron more rapidly due to dilution effects with unfertilized water. As a consequence of expansion and dilution,~~ previous aOIF experiments also produced similar results to this model study. The lateral dilution rate ($<0.25 \text{ d}^{-1}$) during SAGE, which had the smallest fertilized patch size (36 km^2) of the SO experiments, was two times higher than the rates ($<0.11 \text{ d}^{-1}$) in the SO experiments with a larger fertilized patch size (e.g., EIFEX fertilized with a patch size of 167 km^2 and SOFeX-~~N/S~~ fertilized with a patch size of 225 km^2) (Coale et al., 2004; Harvey et al., 2010; Law et al., 2011; Smetacek et al., 2012). Therefore, it would be more appropriate to fertilize a large area (e.g., LOHAFEX had the largest aOIF experiment at 300 km^2), which would reduce the dilution effect with unfertilized

waters (Xiu and Chai, 2010). Based on a ~2 nM iron concentration for a patch size of 300 km² and MLD of ~60 m, it would need ~2,000 kg of iron(II) to be applied in a fertilization experiment. Iron should be released into the wake of a ship, with the release track describing an expanding spiral (or square) in the eddy center, with a regular interval of ~1 km throughout the patch, because it is easier to locate a fertilized patch than a point release (Watson et al., 1991). In addition, it should be completed within ~24 hours because of the time-dependent phytoplankton response within the iron-fertilized patch. Previous aOIF experiments have shown that multiple iron additions (≥ 2 infusions) are needed to maintain the dissolved iron concentration required to derive maximum phytoplankton growth within the fertilized patch. For example, in SOIREE it was found that ~~four~~ 4 additions of iron at intervals of about three days led to persistently high levels of both dissolved and particulate iron within the ~~ML mixed layer~~, with a rapid reduction at the end of the experiment, combined with an increase in the concentration of iron-binding ligands (Bowie et al., 2001). In both EIFEX and SOFeX-S, it was also found that multiple iron(II) infusions (in particular, ~~two~~ 2 infusions with intervals of 13 days in EIFEX and ~~four~~ 4 infusions with intervals of four days in SOFeX-S) allowed iron to persist in the ~~ML mixed layer~~ longer than its expected oxidation kinetics. The relatively low oxidation rates were related to a combination of photochemical production, slow oxidation and, possibly, organic complexation (Croot et al., 2008). Blain et al. (2007) explained that the higher carbon sequestration effectiveness of nOIF experiments compared to aOIF experiments partly resulted from the slow and continuous iron addition that occurs in the natural environment. Large amounts of iron addition at one time can lead to a substantial loss of artificially added iron. Therefore, for an experimental duration of >~40 days, a minimum of ~~two~~ 2 (or three) iron infusions at intervals of ~10–15 days would be required to prevent the iron limitation on phytoplankton growth, based on the ~~EisenEx and~~ EIFEX experiments (Nishioka et al., 2005; Smetacek et al., 2012).

What: The fifth consideration for a successful aOIF experiment is the effective tracing of the fertilized patch, including the detection of carbon sequestration (Buesseler and Boyd, 2003). All previous aOIF experiments used physical tracers, in particular GPS and ~~ARGO~~ Argos equipped drifting buoys, to follow the iron fertilized patch. ~~The release of GPS and Argos equipped drifting buoys at the center of the patch after the iron infusions would provide a visual map showing the tracked positions of the fertilized patch, because Aa~~ drifting buoy is a natural and passive system moving along with the currents. ~~However, but~~ it can be escaped from the fertilized patch due to the action of strong winds (Tsumune et al., 2005). ~~Therefore, the release of GPS and ARGO Argos equipped drifting buoys at the center of the patch after the iron infusions would provide a visual map showing the tracked positions of the fertilized patch.~~ An inert chemical tracer, such as SF₆, would also be an excellent option for following the fertilized patch after iron addition. Previous aOIF experiments have shown that the SF₆ measurements based on underway sampling systems can be used to accurately determine ~~accurately~~ time-dependent vertical and lateral transport of iron-fertilized patches. ~~However, tracing via SF₆ allows for only a limited period (~2 weeks) due to air-sea gas exchange (Law et al., 2006; Tsumune et al., 2009; Martin et al., 2013). Thus, M~~ many subsequent aOIF experiments have also used tracing methods based on the observation of biogeochemical parameters (such as the Fv/Fm ratio, chlorophyll fluorescence, and underway pCO₂) before and after iron addition (Martin et al., 1994; Coale et al., 1996; Boyd et al., 2000; ~~Coale et al., 2004;~~ Boyd et al., 2004; ~~Coale et al., 2004;~~ Tsuda et al., 2005; Smetacek et al., 2012). The Fv/Fm ratio can be easily and promptly used as an indicator to track the fertilized patch due to its rapid response to iron addition. Direct measurements of carbon export fluxes to determine the effectiveness of aOIF should be conducted by deploying an NBST at two depths: (1) ~~within the mixed layer just below the in situ MLD mixed layer~~ to detect increases in iron-induced POC in the surface layer along with the calibration of ~~the a~~ water-column based ²³⁴Th method, and (2) ~~below at the depth of the winter MLD mixed layer MLD~~ to detect iron-induced ~~export~~ carbon ~~export~~ fluxes ~~into intermediate/deeper waters below winter MLD~~ (Bidigare et al., 1999; Nodder et al., 2001; Boyd et al., 2004; Buesseler et al., 2004; Coale et al., 2004; Aono et al., 2005; Buesseler et al., 2005; Tsuda et al., 2007; Smetacek et al., 2012; Martin et al., 2013). Sinking-particle profiling systems ~~(e.g., transmissometers mounted on autonomous floats and gliders), such as a transmissometer and UVP~~ that

measure ~~and photograph~~ sinking particles; could provide a record of the temporal and vertical evolution of iron-induced POC stocks through successive depth layers down to ~3,000 ~~-m depth~~ for ~20 months after deployment, once calibrated using POC fluxes measured from sediment traps and/or ~~the a-~~water-column based ^{234}Th method (Bishop et al., 2004; Smetacek et al., 2012; ~~Martin et al., 2013~~). Repeat casts with UVPs mounted on the rosette could also serve a similar purpose providing a
5 photographic history of the water column (Martin et al., 2013). Future ~~a~~OIF experiments would benefit from these technological advances, enabling a more efficient tracing of the carbon export flux and particle size and composition at higher vertical and temporal resolution than has been possible in the past. Hence, the application of an NBST system and water-column based ^{234}Th method to direct flux estimates, combined with autonomous sinking-particle profilers of a transmissometer and an UVP, will enable the quantitative and qualitative evaluation of the effectiveness of aOIF and direct
10 observation of iron-induced carbon export fluxes after artificial iron additions.

What concerns: The sixth consideration for a successful aOIF experiment is the monitoring of possible side effects. The LC/LP parties recently adopted Resolution LC-LP.2 (2010), which includes the “Assessment Framework for Scientific Research Involving Ocean Fertilization”. This considers possible side effects on marine/ocean ecosystems after artificial iron additions, such as the production of climate-relevant gases and negative ecosystem changes, which are vital to assess when
15 proposing an aOIF experiment. The emissions of climate-relevant gases, such as N_2O , DMS, and HVOCs, may directly contribute to warming or cooling effects, and oxygen decrease and toxic DA production may have a negative impact on marine/ocean ecosystems (Law, 2008; Silver et al., 2010; Trick et al., 2010; Williamson et al., 2012), resulting in significant offsets against the benefits of aOIF experiments. However, there is little quantitative and qualitative information regarding possible side effects following the previous aOIF experiments. Therefore, the future monitoring of these potential side effects
20 is a prerequisite to evaluate accurately the effectiveness of an aOIF experiment in the future. The possible side effects after an aOIF experiment can be continuously monitored from optical sensors equipped autonomous moored profiler and/or autonomous benthic vehicle (e.g., crawler, which is capable to perform a long-term benthic oxygen measurements for ~12 months) (Dunne et al., 2002; Purser et al., 2013; Wenzhöfer et al., 2016).

In summary, to maximize the effectiveness of aOIF experiments in the future, we suggest a design that incorporates
25 several conditions. (1) Experiments are conducted in the center of an eddy structure when grazing pressure is low and silicate levels are high (e.g., in the case of SO, at the south of PF during the early summer). (2) Shipboard observations are made during a minimum of ~40 days, with multiple iron injections (iron infusions of ~2,000 kg at least ~~two2 (or three3)~~ times, with an interval of ~10–15 days, to fertilize a patch of 300 km^2 to obtain a ~2 nM concentration). (3) The iron-fertilized patch is traced using both physical (e.g., a drifting buoy) and biogeochemical (e.g., SF_6 and the Fv/Fm ratio) tracers. (4)
30 NBST system and water-column derived ^{234}Th method are employed at two depths (~~i.e., one is within just below the in situ the MLDmixed layer (i.e., 10–20 m below) and—another is at the depth of the winter MLDmixed layer (i.e., ~450 m)below it~~), with autonomous profilers equipped with an UVP and a transmissometers to estimate accurately the carbon export flux. (5) The side effects on marine/ocean ecosystems, ~~are monitored,~~ including decline in oxygen contents and the production of climate-relevant gases (e.g., N_2O , DMS, and HVOCs) and toxic DA, ~~are monitored- using optical sensors equipped~~
35 autonomous moored profiler and/or autonomous benthic vehicle.

5. Design of the Korean Iron Fertilization Experiment in the Southern Ocean (KIFES)

5.1 Background - Bransfield Basin

A science-oriented aOIF project, KIFES (Fig. 11), was launched in 2016 with research funding from the Korean Ministry of Oceans and Fisheries. This project was largely managed by the Korea Polar Research Institute (KOPRI) with domestic collaborators (i.e., Incheon National University, Inha University, Pusan National University, Hanyang University, and Yeonsei University) and strengthened by international collaborators (i.e., Alfred-Wegener-Institut (AWI), Institute of Geological and Nuclear Sciences, ~~Massachusetts Institute of Technology~~ Woods Hole Oceanographic Institution (~~MIT~~ WHOI), University of Otago, University of California at Irvine, McMaster University, University of South Florida, Royal Netherlands Institute for Sea Research, and Dalhousie University). KIFES had four main aims. (1) To conduct the first scientific aOIF experiment complying with the “Assessment Framework for Scientific Research Involving Ocean Fertilization” after the framework was accepted from the LC/LP in 2010. (2) To evaluate the effectiveness of scientific aOIF in terms of atmospheric carbon sequestration (i.e., to identify/quantify significant increases in iron-induced carbon export fluxes ~~into intermediate/deeper waters below the winter MLD~~) in the SO. (3) To determine the environmental conditions that would maximize the effectiveness of aOIF. (4) To quantitatively and qualitatively monitor short- and long-term possible side effects derived from previous aOIF experiments.

A location near the eastern Bransfield Basin was considered for the site of KIFES based on the following three criteria: (1) the possibility of diatom blooms, (2) the proximity to meso-scale eddies, and (3) the availability of historical oceanographic data. The development of a diatom bloom is the first prerequisite to maximize the effectiveness of an aOIF experiment. The paleoclimate team at KOPRI had found geological evidence of massive amounts of organic carbon buried in the sediments, especially in the diatomaceous ooze layer, in the eastern Bransfield Basin on the Antarctic Peninsula (Yoo et al., 2016). The well-preserved diatomaceous ooze layer (Bahk et al., 2003; Kang et al., 2003; Bak et al., 2015) indicates high accumulation rates of fast sinking diatoms, suggesting the existence of a strong ‘biological pump (i.e., significant export production)’ in this basin. In addition, this basin has a high silicate concentration of $\sim 30 \mu\text{M}$ (Fig. 4c), which is a fundamental condition to produce a massive diatom bloom. Despite the favorable environmental conditions, the Fv/Fm ratio in/near the eastern Bransfield Basin ($< \sim 0.43$) (Park et al., 2013) was lower than the maximum value of 0.65 measured during the aOIF experiments in the SO (e.g., SOIREE and SOFeX-S), suggesting an iron limitation on diatom growth. Therefore, we hypothesized that the input of bioavailable iron enabling an increase in productivity and export would lead to a massive enhancement of the diatom flux in this basin. Accordingly, we expected that an aOIF in the diatom-dominated region with high sinking rates near the eastern Bransfield Basin would be more effective for carbon export, as compared to the previous aOIF experiments conducted in the SO. This hypothesis, based on sedimentary evidence, was not considered in the site selection for previous experiments. A second important factor was the presence of stable eddies in/near the eastern Bransfield Basin (Kahru et al., 2007; Sangrà et al., 2011), providing coherent structures that made it possible to track effectively the iron-induced carbon export fluxes (Smetacek et al., 2012). For example, Thompson et al. (2009) showed that a large standing eddy ($\sim 40 \text{ km}$ in diameter) was centered at $\sim 62^\circ\text{S}$ and 54°W and remained for ~ 30 days using historical drifters released during the period 1989–2005, and 40 drifters released in February 2007 as part of the Antarctic Drifter Experiment: Links to Isobaths and Ecosystems project. Satellite sea-level height images have indicated meso-scale eddies with long lifespans in/near the eastern Bransfield Basin (<https://www.aviso.altimetry.fr/>). Additionally, several historical oceanographic datasets are available for this basin (Grelowski et al., 1986; Helbling et al., 1995; Figueiras et al., 1999; Kang et al., 2001; Varela et al., 2002; Khim et al., 2005). The historical datasets provide general oceanographic characteristics about sites for an aOIF experiment as well as basic information helpful for designing the experiment, including a preliminary hydrographic survey. Unfortunately, KIFES has lost its source of funding. Nevertheless, optimism prevails that alternative funding will be found at a future date and the following section is intended, therefore, to provide a basic set of design guidelines, with the expectation that an opportunity to move forward with KIFES will occur in the near future.

5.2 A plan for the future: KIFES

The KIFES design entails a five-year project plan modeled on the ‘EIFEX’ program that found deep carbon by conducting an aOIF experiment in the center of an eddy structure. The KIFES project would include a preliminary environmental survey both outside and inside the center of an eddy structure formed in/near the eastern Bransfield Basin, a scientific aOIF experiment, and an assessment of the full KIFES project. In this section, we introduce the major goals, objectives, and main tasks of KIFES.

5.2.1 Year one plan

Goals: (1) Data collection with regard to oceanographic conditions in/near the eastern Bransfield Basin, including both eddy development and distribution. (2) Establishment of the study aims, hypothesis, and site for the KIFES experiment.

Objective: To understand the physical and biogeochemical oceanography of relevance to the eastern Bransfield Basin as an aOIF site through an analysis of earlier datasets and a review of published papers.

Main tasks: (1) Review databases of physical and biogeochemical parameters from previous surveys conducted in/near the eastern Bransfield Basin. (2) Review the eastern Bransfield Basin oceanographic conditions using data analysis and references. (3) Establish the study aims, hypothesis, and site in/near the eastern Bransfield Basin for an aOIF experiment, based on the results obtained from tasks (1) and (2). (4) Design an oceanographic cruise map for the first preliminary survey in/near the eastern Bransfield Basin. (5) Analyze eddy development and distribution using satellite data in/near the eastern Bransfield Basin. (6) Prepare scientific instruments for ocean physical and biogeochemical monitoring. (7) Establish an international collaborative aOIF network. (8) Submit KIFES field program proposal for the ‘Initial Assessment’ to determine that KIFES falls within the remit of ocean fertilization and should be evaluated in the LC/LP assessment framework based on the results from tasks (1) ~~and (24)~~.

5.2.2 Year two plan

Goal: First preliminary hydrographic survey to provide a foundational understanding of oceanographic conditions in/near the eastern Bransfield Basin.

Objectives: (1) To obtain information about oceanographic conditions from *in situ* measurements in/near the eastern Bransfield Basin. (2) To provide background information before the KIFES experiment.

Main tasks: (1) Using the ice breaker RV *ARAON*, undertake a field investigation in/near the eastern Bransfield Basin to determine physical and biogeochemical parameters associated with both carbon sequestration and aOIF side effects (e.g., decline in oxygen inventory and production of N₂O, DMS, HVOCs, and DA), based on the first-year results. (2) Prepare an ‘Environmental Assessment’ for the LC/LP assessment framework based on the first-year results and a preliminary hydrographic survey.

5.2.3 Year three plan

Goals: (1) Preliminary hydrographic survey outside/inside the center of an eddy structure prior to the KIFES experiment. (2) Approval of KIFES from LC/LP.

Objectives: (1) To compare oceanographic conditions inside and outside the center of an eddy structure formed in/near the eastern Bransfield Basin prior to the KIFES experiment. (2) To obtain a permission on the basis that the proposed KIFES is legitimate scientific research from the LC/LP.

Main tasks: (1) Using the ice breaker RV *ARAON*, detect an eddy formed in/near the eastern Bransfield Basin using observations from acoustic Doppler current profilers (ADCPs) and satellites. (2) Conduct intensive physical and biogeochemical field investigations both inside and outside the center of an eddy structure. (3) Assess the physical and biogeochemical properties outside vs. inside the center of an eddy structure prior to KIFES. (4) Establish a final design for KIFES. (5) Submit the research results for 'Environmental Assessment' stage of the LC/LP assessment framework and obtain approval for the KIFES experiment via the 'Decision Making' process from the LC/LP.

5.2.4 Year four plan

Goal: Conduction of the KIFES scientific aOIF experiment in the center of an eddy structure during the early summertime (Fig. 11).

Objective: To conduct a scientific aOIF experiment in the center of an eddy structure formed near/in the eastern Bransfield Basin.

Main tasks: (1) Using the ice breaker RV *ARAON*, detect an eddy formed in/near the eastern Bransfield Basin using observations from ADCPs and satellites, and investigate the initial environmental conditions for ~4 days before KIFES. (2) Execute the KIFES field campaign during a >~40-day period with the eddy structure. (3) At least ~~two~~² (or ~~three~~³) iron additions at intervals of ~15 days, with each iron injection being ~2,000 kg following a spiral ship track, with a regular interval of ~1 km to create a patch size of 300 km² (target dissolved iron concentration of ~2 nM). (4) Trace the fertilized patch with deployments of GPS and ~~ARGO~~^{Argos} equipped drifting buoys, biogeochemical tracers (SF₆ and Fv/Fm ratio) employing underway-sampling systems, and gliders equipped with biogeochemical sensors. (5) Measure iron-induced carbon export fluxes for the regions both inside and outside the center of an eddy structure using NBST systems at two depths (~~i.e., one is just below the in situ the MLDmixed layer (i.e., 10-20 m below) within the mixed layer and another is at the depth of the winter MLDmixed layer (i.e., ~450 m) below it~~) along with the calibration of water-column based ²³⁴Th measurements and autonomous profilers equipped with ~~a~~^a transmissometer and ~~an~~^{an} UVP. (6) Monitor possible side effects, such as ~~the decline in oxygen contents and~~ the production of climate-relevant gases and toxic DA, ~~using optical sensors equipped autonomous moored profiler and/or autonomous benthic vehicle~~. (7) Monitor continued responses after KIFES termination using satellite observations and autonomous profilers. (8) Assess the effectiveness of carbon sequestration and environmental (ocean and atmosphere) side effects for KIFES and prepare the KIFES assessment for the 'Results of Monitoring' stage of the LC/LP assessment framework.

5.2.5 Year five plan

Goal: Integrated assessment of the KIFES project.

Objective: To evaluate whether small-scale scientific aOIF experimentation can be an effective tool for detecting the effectiveness of artificially iron-induced export production and determining any negative impacts on climate change.

Main tasks: (1) Submit the KIFES assessment report. (2) Submit scientific results to international journals. (3) Collect feedback regarding the KIFES project from international scientific/oceanographic communities. (4) Produce a final aOIF

experimental summary (including main tasks (1)–(3)). (5) Evaluate effectiveness and environmental side effects of large-scale SO aOIF via more realistic simulations under various scenarios with ocean biogeochemical models using the integrated results of KIFES. (6) Submit a final report of the KIFES assessment to the LC/LP.

5.3 Final Remark

None of the KIFES scientists have commercial interests (i.e., carbon credits) related to aOIF experiments. The interests of KIFES participants all lie in the detailed investigation of the biogeochemical effects of scientific ~~artificial iron addition~~aOIF in the SO and in aOIF as a possible geo-engineering method to mitigate the climate change effects we will face in the future. We envisage a future where the KIFES, or similar projects, can be resumed, enabling a more robust assessment of the potential of aOIF as a geo-engineering solution to help reduce atmospheric CO₂ concentrations. A continuation of the KIFES project would provide fundamental information and guidelines for future scientific aOIF experiments in HNLC regions, as well as improving our understanding of SO biogeochemistry. The risks and side effects of aOIF should be thoroughly investigated to calm international concerns. Finally, we emphasize that international cooperation is essential for a project as organizationally and scientifically complex as KIFES, and that we seek to improve our knowledge and provide a positive outlook for the Earth's future.

6 Summary

To test Martin's hypothesis, a total of 13 scientific aOIF experiments have been conducted in HNLC regions during the last 25 years. These aOIF experiments have resulted in increases of PP and drawdowns of macro-nutrients and DIC. In most experiments, the phytoplankton group has tended to shift from small-sized to large-sized plankton cells (mostly diatom-dominated). However, their effectiveness in enhancing export production has not been confirmed, except for EIFEX. Likewise, the possible environmental negative side effects in response to iron addition, such as decline in oxygen contents and the production of climate-relevant gases and toxic DA, could not be fully evaluated due to the widely differing outcomes, with large uncertainties depending on aOIF experimental conditions and settings. In particular, the monitoring of N₂O, DMS, and HVOCs is essential to determine the effectiveness of aOIF as a geoengineering approach, because these potential trace gas emissions can directly and indirectly modify the carbon reduction benefits resulting from aOIF. Furthermore, oxygen decline and toxic DA production may cause serious damage to marine/ocean ecosystems. Therefore, the validation and suitability of aOIF for the mitigation of rapidly increasing atmospheric CO₂ levels is a subject of vigorous debate. At present, large-scale and/or commercial aOIF is prohibited by international regulation, while small-scale aOIF experimentation for scientific purposes is permitted. To maximize the effectiveness of aOIF, future aOIF experiments should be conducted by carefully considering the major factors including the methods for iron addition, tracking methods, measurement parameters, location, timing, and experimental duration, under international aOIF regulations. Finally, we envisage a future where the KIFES project, or a similar alternative, becomes a reality so that we may determine whether aOIF is a promising geo-engineering solution.

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Table 1. Summary of ocean iron fertilization (OIF) experiments; time, location, research vessel, added iron(~~II-III~~) (values in brackets correspond to the number of days from the first iron addition, e.g., the first iron addition becomes (0)), initial iron concentrations, ~~target~~~~after~~ iron ~~addition~~ concentrations (iron concentrations after iron addition), tracer, initial patch size, experiment duration, and regional characteristics (HNLC: high-nutrient and ~~low-low~~-chlorophyll).

Experiment	Time	Location	Research vessel	Added iron(II-III) (kg) (day)	Initial iron (nM)	After-Target iron_ addition (nM)	Tracer	Patch size (km ²)	Duration (days)	Regional characteristics
1 IronEx-1	Oct 1993	Equatorial Pacific 5° S, 90° W	RV <i>Columbus Iselin</i>	①450 (0)	0.06	3.60	SF ₆	64	10	HNLC
2 IronEx-2	May 1995	Equatorial Pacific 3.5° S, 104° W	RV <i>Melville</i>	①225 (0) ②112 (3) ③112 (7)	0.02	2.00 1.00 1.00	SF ₆	72	17	HNLC
3 SOIREE	Feb 1999	Southern Ocean- Australasian-Pacific sector 61° S, 140° E	RV <i>Astrolab</i>	①768 (0) ②312 (3) ③312 (5) ④353 (7)	0.08	3.80 2.60 2.60 2.50	SF ₆	50	13	HNLC
4 EisenEx	Nov 2000	Southern Ocean- Atlantic sector 48° S, 21° E	RV <i>Polarstern</i>	①780 (0) ②780 (7) ③780 (16)	0.06	2.00	SF ₆	50	23	HNLC
5 SOFeX-N	Jan–Feb 2002	Southern Ocean- Pacific sector 56.23° S, 172° W	RV <i>Revelle</i> RV <i>Melville</i>	①631 (0) ②631 (4) ③450 (29)		1.20 1.20 1.50	SF ₆	225	40	^a HNLCLSi ^a
6 SOFeX-S	Jan–Feb 2002	Southern Ocean- Pacific sector 66.45° S, 171.8° W	RV <i>Revelle</i> RV <i>Melville</i> RV <i>Polar star</i>	①315 (0) ②315 (5) ③315 (8) ④315 (12)		0.70 0.70 0.70 0.70	SF ₆	225	28	HNLC
7 EIFEX	Feb–Mar 2004	Southern Ocean- Atlantic sector 50° S, 2° E	RV <i>Polarstern</i>	①1410 (0) ②1410 (13)	0.08–0.20 0.20	1.50 0.34		167	39	HNLC

To be continued

Experiment		Time	Location	Research vessel	Added iron (II) (H) (kg) (day)	Initial iron (nM)	Target Aft er iron _addition (nM)	Tracer	Patch size (km ²)	Duration (days)	Regional characteristics
8	SAGE	Mar–Apr 2004	Southern Ocean- Southeast of New Zealand 46.5° S 172.5° E	RV <i>Tangaroa</i>	①265 (0) ②265 (6) ③265 (9) ④265 (12)	0.09	3.03 1.59 0.55 1.01	SF ₆	36	15	^a HNLC ^{Si} ^a
9	LOHAFEX	Jan–Mar 2009	Southern Ocean- Atlantic sector 48° S, 15° W	RV <i>Polarstern</i>	①2000 (0) ②2000 (18)		2.00	SF ₆	300	40	^a HNLC ^{Si} ^a
10	SEEDS-1	Jul–Aug 2001	Subarctic North Pacific- Western basin 48.5° N, 165° E	RV <i>Kaiyo-Maru</i>	①350 (0)	0.05	2.90	SF ₆	80	13	HNLC
11	SERIES	Jul-Aug 2002	Subarctic North Pacific- Eastern basin 50.14° N, 144.75° W	RV <i>John P. Tully</i> RV <i>El Puma</i> RV <i>Kaiyo-Maru</i>	①245 (0) ②245 (6)	<0.10	≥12.00 0.60	SF ₆	77	25	HNLC
12	SEEDS-2	Jul–Aug 2004	Subarctic North Pacific- Western basin 48° N, 166° E	RV <i>Hakuho-Maru</i> RV <i>Kilo-Moana</i>	①332 (0) ②159 (6)	0.17	1.38	SF ₆	64	26	HNLC
13	FeeP	Apr–May 2004	Subtropical North Atlantic- North-east Atlantic 27.5° N 22.5° W	RV <i>Charles Darwin</i> RV <i>Poseidon</i>	①1840 (0)	0.20–0.40	3.00	SF ₆	25	21	LNLC ^b
I	<u>Polar</u> <u>Front^c</u> <u>EX^e</u>	<u>Oct–Nov</u> <u>1992Nov–</u> <u>2004–</u> <u>Jan 2005</u>	<u>Southern Ocean–</u> <u>Atlantic Sector</u> <u>48° S, 6° W</u> <u>Southern Ocean–</u> <u>Crozet Plateau</u> <u>44° S, 50° E</u>	<u>RV <i>Polarstern</i></u> <u>RV <i>Discovery</i></u>	-		<u>0.55</u>		-	-	<u>HNLC</u>

<u>II</u>	<u>PlumeEx^c</u>	<u>Nov 1993</u>	<u>Equatorial Pacific</u> <u>2° S, 89° W</u>	<u>RV Columbus Iselin</u>	<u>0.05</u>	<u>0.2</u>				<u>HNLC</u>
<u>H</u>	<u>KEOPS-1^e</u>	<u>Jan–Feb 2005</u>	<u>Southern Ocean–</u> <u>Kerguelen Plateau</u> <u>50° S, 73° E</u>	<u>RV Marion Dufresne</u>	<u>0.09</u>	<u>0.35</u>	<u>-</u>	<u>-</u>	<u>-</u>	<u>HNLC</u>

To be continued

<u>-</u>	<u>Experiment</u>	<u>Time</u>	<u>Location</u>	<u>Research vessel</u>	<u>Added iron(II)</u> <u>(kg) (day)</u>	<u>Initial iron</u> <u>(nM)</u>	<u>Target</u> <u>iron (nM)</u>	<u>Tracer</u>	<u>Patch size</u> <u>(km²)</u>	<u>Duration</u> <u>(days)</u>	<u>Regional</u> <u>characteristics</u>
<u>III</u>	<u>CROZEX^c</u>	<u>Nov 2004–</u> <u>Jan 2005</u>	<u>Southern Ocean–</u> <u>Crozet Plateau</u> <u>44° S, 50° E</u>	<u>RV Discovery</u>	<u>-</u>		<u>0.55</u>		<u>-</u>	<u>-</u>	<u>HNLC</u>
<u>IV</u>	<u>KEOPS-1^c</u>	<u>Jan–Feb 2005</u>	<u>Southern Ocean–</u> <u>Kerguelen Plateau</u> <u>50° S, 73° E</u>	<u>RV Marion Dufresne</u>	<u>-</u>	<u>0.09</u>	<u>0.35</u>		<u>-</u>	<u>-</u>	<u>HNLC</u>
<u>V</u>	<u>DynaLiFe^c</u>	<u>Jan–Feb 2009</u>	<u>Southern Ocean–</u> <u>Pacific sector</u> <u>74° S, 105° W</u>	<u>RV Nathaniel B.</u> <u>Palmer</u>		<u>0.2</u>	<u>0.4</u>				<u>HNLC</u>
<u>VI</u>	<u>KEOPS-2^c</u>	<u>Oct–Nov 2011</u>	<u>Southern Ocean–</u> <u>Kerguelen Plateau</u> <u>50.63° S, 72.08° E</u>	<u>RV Marion Dufresne</u>	<u>-</u>				<u>-</u>	<u>-</u>	<u>HNLC</u>

^aHigh–Nutrient Low–Low–Chlorophyll and Low–Low–Silicate (HNLCLS) region; ^bLow–Nutrient Low–Chlorophyll (LNLC) region; ^cNatural OIF experiments (PlumeEx: Natural iron enrichment experiment near Galapagos Islands; CROZEX: CROZet natural iron bloom and EXport experiment; Kerguelen Ocean and Plateau compared Study-1/2; KEOPS-1/2: Kerguelen Ocean and Plateau compared Study-1/2; DynaLiFe: Dynamic Light on Iron Limitation program; DynaLiFe).

Sources are Martin et al. (1994); de Baar et al. (1995); Coale et al. (1996); Coale et al. (1998); Gordon et al. (1998); Boyd et al. (2000); Boyd and Law (2001); Gervais et al. (2002); Tsuda et al. (2003); Boyd et al. (2004); Coale et al. (2004); Bakker et al. (2005); Boyd et al. (2005); de Baar et al. (2005); Nishioka et al. (2005); Hoffmann et al. (2006); Law et al. (2006); Blain et al. (2007); Boyd et al. (2007); Rees et al. (2007); Tsuda et al. (2007); Pollard et al. (2009); Strong et al. (2009); Harvey et al. (2010); Gerringa et al. (2012); Smetacek et al. (2012); and Martin et al. (2013); and Blain et al. (2015).

Table 2. Summary of artificial ocean iron fertilization (aOIF) experiments; objectives, significant results, and limitations.

	Experiment	Objectives	Significant results	Limitations
1	IronEx-1	<ul style="list-style-type: none"> To test the hypothesis that artificial iron addition will increase phytoplankton productivity by relieving the iron limitation of phytoplankton in high-nutrient low chlorophyll regions 	<ul style="list-style-type: none"> Small increases in the pCO₂ concentrations, Fv/Fm ratio, chlorophyll-a concentration, and primary production (PP) Insignificant changes in nutrients and pCO₂ concentrations 	<ul style="list-style-type: none"> Single iron addition Insufficient experimental periods to observe the full phases of biogeochemical responses from the onset to termination after iron additions Micro/macro-nutrient limitations
2	IronEx-2	<ul style="list-style-type: none"> To test three-four hypotheses that were advanced to explain the weak biogeochemical response observed during IronEx-1 	<ul style="list-style-type: none"> Dramatic changes in biogeochemical responses; close to—support for Martin’s hypothesis Taxonomic shift toward diatom-dominated phytoplankton communities 	<ul style="list-style-type: none"> No export flux measurements in the deep—ocean Insufficient experimental duration
3	SOIREE	<ul style="list-style-type: none"> To test the iron hypothesis in the Southern Ocean 	<ul style="list-style-type: none"> Diatom-dominated bloom No measurable change in carbon export 	<ul style="list-style-type: none"> Insufficient experimental duration
4	EisenEx	<ul style="list-style-type: none"> To test the hypothesis that atmospheric dust inputs might have led to a dramatic increase in ocean productivity during the Last Glacial Maximum due to the relief of iron-limited conditions for phytoplankton growth 	<ul style="list-style-type: none"> Diatom-dominated bloom No clear differences in carbon flux between in-patch and outside-patch 	<ul style="list-style-type: none"> Light limitation by storms Insufficient experimental duration
5	SOFeX-N	<ul style="list-style-type: none"> To address the potential for iron and silicate interactions to regulate the diatom bloom 	<ul style="list-style-type: none"> Remarkable increase in diatom biomass Observation of large export flux event with transmissometers 	<ul style="list-style-type: none"> Entrainment of dissolved silicate into the fertilized patch by physical mixing No direct measurement of export fluxes with sediment traps
6	SOFeX-S	<ul style="list-style-type: none"> To address the potential for iron and silicate interactions to regulate the diatom bloom 	<ul style="list-style-type: none"> Significantly enhanced export fluxes out of the mixed—l— Layer (ML) depth (MLD), but similar to those for natural blooms 	<ul style="list-style-type: none"> Insufficient experimental duration
7	EIFEX	<ul style="list-style-type: none"> To confirm that aOIF experiments can increase export production 	<ul style="list-style-type: none"> Observation of all the phases of the phytoplankton bloom from onset to termination Significant carbon export to deeper layers (down to 3,000 m) due to the formation of aggregates with rapid sinking rates • The occurrence of rapidly sinking large aggregates— 	

To be continued

Experiment		Objective	Significant results	Limitations
8	SAGE	<ul style="list-style-type: none"> To determine the response of phytoplankton dynamics to iron addition in high--nutrient low--chlorophyll and low silicate (HNLCLSi) regions To test the assumption that the response of phytoplankton blooms to artificial iron addition can be detected by the enhanced air-sea exchanges of climate-relevant gases 	<ul style="list-style-type: none"> No shift to a diatom-dominated community No detection of fertilization-induced export 	<ul style="list-style-type: none"> High dilution rate by small patch size
9	LOHAFEX	<ul style="list-style-type: none"> To trace the fate of iron-stimulated phytoplankton blooms and deep carbon export in HNLCLSi regions 	<ul style="list-style-type: none"> Observation of all the phases of the phytoplankton bloom from onset to termination No shift to a diatom-dominated community No detection of fertilization-induced export High grazing pressure and active bacterial respiration 	
10	SEEDS-1	<ul style="list-style-type: none"> To investigate the relationship between phytoplankton biomass/community and dust deposition in the subarctic North Pacific (NP) To investigate changes in phytoplankton composition and vertical carbon flux 	<ul style="list-style-type: none"> A shift from oceanic diatoms to fast-growing neritic ones The largest changes in biogeochemical parameters of all aOIF experiments No detection of large POC-carbon export flux 	<ul style="list-style-type: none"> Single iron addition Insufficient experimental duration
11	SERIES	<ul style="list-style-type: none"> To compare the response of phytoplankton in eastern subarctic with that in the western subarctic ecosystem To investigate the most significant factor that controls the beginning to the ending of the phytoplankton bloom induced by iron addition 	<ul style="list-style-type: none"> Observation of all phases of the phytoplankton bloom from onset to termination No significant increases in export fluxes below the mixed-layer depthMLDML High bacterial remineralization and mesozooplankton grazing pressure 	
12	SEEDS-2	<ul style="list-style-type: none"> To investigate the most significant factor that controls the beginning to the ending of the phytoplankton bloom induced by iron addition 	<ul style="list-style-type: none"> Observation of all phases of the phytoplankton bloom from onset to termination No shift to a diatom-dominated community No significant increases in export fluxes Extensive copepod grazing 	
13	FeeP	<ul style="list-style-type: none"> To investigate the impact of iron and phosphate co-limitation on PP 	<ul style="list-style-type: none"> Increases in pico-phytoplankton abundances 	

Sources are Martin et al. (1994); Coale et al. (1996); Coale et al. (1998); Bidigare et al. (1999); Boyd et al. (2000); Charette and Buesseler (2000); Gervais et al. (2002); Tsuda et al.

(2003); Boyd et al. (2004); Coale et al. (2004); Bakker et al. (2005); Boyd et al. (2005); de Baar et al. (2005); Hiscock and Millero (2005); Nishioka et al. (2005); Tsuda et al. (2005); Tsumune et al. (2005); Boyd et al. (2007); Rees et al. (2007); Tsuda et al. (2007); Harvey et al. (2010); Law et al. (2011); Smetacek et al. (2012); and Martin et al. (2013).

Table 3. Initial conditions and changes (Δ values) in chemical parameters during the artificial ocean iron fertilization (aOIF) experiments

	Experiment	Initial NO ₃ ⁻ (μ M)	Δ NO ₃ ⁻ (μ M)	Initial PO ₄ ³⁻ (μ M)	Δ PO ₄ ³⁻ (μ M)	Initial Si (μ M)	Δ Si (μ M)	Initial <i>p</i> CO ₂ (μ atm)	Δ <i>p</i> CO ₂ (μ atm)	Initial DIC (μ M)	Δ DIC (μ M)
1	IronEx-1	10.8	-0.70	0.92	-0.02	3.90	-0.02	471	-13.0	2044 ^a	^{**} -6.00 ^a
2	IronEx-2	10.4	-4.00	0.80	-0.25	5.10	-4.00	538	-73.0	2051 ^a	^{**} -27.0 ^a
3	SOIREE	25.0	-2.90	1.50	-0.24	10.0	-2.90	349	- (38.0–3 2.0)	<u>2137</u>	- (18.0–1 5.0)
4	EisenEx	<u>223.5</u> <u>0</u>	-1.0 <u>60</u>	1.60	-0.10 <u>6</u>	<u>104.0</u> <u>2</u>	<u>0</u>	~360	(20.0–1 8.0)	<u>2131</u>	- (15.0–1 2.0)
5	SOFEX-N	21.9	-1.40	1.40	-0.09	2.50	-1.10	367	-26.0	2109	-14.0
6	SOFEX-S	26.3	-3.50	1.87	-0.21	62.8	-4.00	365	-36.0	2176	-21.0
7	EIFEX	25.0	-1.60	1.80	[~] 0.30 ^b	19.0	-11.0	360	-30.0	2135	-13.5
8	SAGE	7.90– 10.5	1.30–3. 90	0.62– 0.85		0.83– 0.97		330	8.00	2057	25.0
9	LOHAFEX	20.0	-2.50	1.20– 1.30	[~] 0.15 ^c	0.60– 1.60		[*] ~358 ^d	- (15.0–7 .00)		
10	SEEDS-1	18.5	-15.8			31.8	-26.8	390	-130		-58.0
11	SERIES	10.0– 12.0	- (8.50–6 .50)	1.00	-0.50	14.0– 16.0	- (13.6–1 1.6)	350	-85.0	2030	-37.0
12	SEEDS-2	18.4	-5.70			36.1		370	-6.00		
13	FeeP	<0.01		~0.01							~-1.00

^aDissolved inorganic carbon (DIC) values in IronEx-1/-2 indicate normalized DIC (normalized DIC = DIC \times 35/Salinity).

^b Δ PO₄³⁻ in EIFEX was digitized from Figure 3 of Smetacek et al. (2012). ^c Δ PO₄³⁻ in LOHAFEX was digitized from Figure 5.1 of Smetacek and Naqvi (2010). ^d Δ pCO₂ in LOHAFEX was digitized from Figure 6.1 of Smetacek and Naqvi (2010).

Sources are Martin et al. (1994); Steinberg et al. (1998); Boyd et al. (2000); Bakker et al. (2001); Frew et al. (2001); Bakker et al. (2005); Boyd et al. (2005); Bozec et al. (2005); Hiscock and Millero (2005); Smetacek et al. (2005); Takeda and Tsuda (2005); Tsuda et al. (2005); Marchetti et al. (2006); Wong et al. (2006); Boyd et al. (2007); Tsuda et al. (2007); Tsumune et al. (2009); Harvey et al. (2010); Smetacek and Naqvi (2010); Berg et al. (2011); Currie et al. (2011); Law et al. (2011); Smetacek et al. (2012); Assmy et al. (2013); Ebersbach et al. (2014); and Latasa et al. (2014). ^{*} Δ PO₄ in EIFEX was digitized from Figure 3 of Smetacek et al. (2012); [~] Δ PO₄ in LOHAFEX was digitized from Figure 5.1 of Smetacek and Naqvi (2010). ^{*} Δ pCO₂ in LOHAFEX was digitized from Figure 6.1 of Smetacek and Naqvi (2010). ^{**}Dissolved inorganic carbon (DIC) values in IronEx-1/-2 indicate normalized DIC (normalized DIC = DIC \times 35/Salinity).

Sources are Martin et al. (1994); Steinberg et al. (1998); Boyd et al. (2000); Bakker et al. (2001); Frew et al. (2001); Gervais et al. (2002); Bakker et al. (2005); Boyd et al. (2005); Bozec et al. (2005); Hiscock and Millero (2005); Smetacek et al. (2005); Takeda and Tsuda (2005); Tsuda et al. (2005); Marchetti et al. (2006); Wong et al. (2006); Boyd et al. (2007); Tsuda et al. (2007); Tsumune et al. (2009); Harvey et al. (2010); Smetacek and Naqvi (2010); Berg et al. (2011); Currie et al. (2011); Law et al. (2011); Smetacek et al. (2012); Assmy et al. (2013); Ebersbach et al. (2014); and Latasa et al. (2014).

Table 4. Initial values of biological parameters and the values after fertilization. Note that maximum values were attained after fertilization.

Experiment	Initial Fv/Fm	After Fv/Fm	Initial Chlorophyll-a (mg m ⁻³)	After Chlorophyll-a (mg m ⁻³)	Initial PP (mg C m ⁻² d ⁻¹)	After PP (mg C m ⁻² d ⁻¹)	Initial Mesozooplankton biomass (mg C m ⁻³)	After Mesozooplankton biomass (mg C m ⁻³)	Initial Heterotrophic b Bacteria abundance (× 10 ⁵ cells ml ⁻¹)	After Heterotrophic b Bacteria abundance (× 10 ⁵ cells ml ⁻¹)
1 IronEx-1	~0.30	0.63	0.24	0.65	^a 300–450 ^a	^a 805–1330 ^a				
2 IronEx-2	0.25	^a ~0.57 ^b	0.15–0.20	4.00	^{**} ~630 ^c	^{**} ~2430 ^c	^d ~6 ^d 3.8 (0–55 m)	^d ~14 ^d 6.6 (0–55 m)	<u>6.5</u> 9.5	<u>10.8</u>
3 SOIREE	0.22	0.65	0.25	2.00	^{***} ~120 ^e	^{***} ~1300 ^e	^f 22.8 ^f (0–65 m)	^f 30.1 ^f (0–65 m)	<u>1.7</u> 3.7	<u>3.9</u>
4 EisenEx	0.30	0.56	0.50	2.50	130–220	790			<u>2.04</u> 4	<u>6.2</u>
5 SOFeX-N	0.20	0.5	^g ~0.15 ^g	^g ~2.60 ^g	^h ~144 ^h	^h ~1500 ^h			<u>4</u> 4	<u>10.9</u>
6 SOFeX-S	0.25	0.65	^g ~0.30 ^g	^g ~3.80 ^g	^h ~216 ^h	^h ~972 ^h			<u>3.34</u> 4	<u>5.3</u>
7 EIFEX	ⁱⁱ ~0.28 ⁱ	ⁱⁱ ~0.6 ⁱ	0.70	3.16	~750	1500				
8 SAGE	0.27	0.61	0.63	1.33	540	900				
9 LOHAFEX	~0.33	0.50	0.50	1.25	<960	1560				
10 SEEDS-1	ⁱⁱ ~0.19 ^j	ⁱⁱ ~0.42 ^j	0.80–0.90	21.8	420	1670	^f 6.8 ^f (0–20 m)	^f 7.5 ^f (0–20 m)	<u>3.2</u> 2.5	<u>8.1</u>
11 SERIES	0.24	0.55	0.35	~5.00	300	>2000	^f 7.3 ^f (0–30 m)		5.5	12
12 SEEDS-2	0.29	ⁱⁱ ~0.43 ^k	0.80	2.48	390	>1000	^f 18.9 ^f (0–20 m)	^f 38 ^f (0–20 m)		
13 FeeP			0.06	0.07						

^aPrimary productivity (PP) in IronEx-1 was estimated by multiplying PP (mg C m⁻³ d⁻¹) with the mixed layer depth (initial: 30 m and after: 35 m). ^bFv/Fm in IronEx-2 was digitized from the Figure 3 of Behrenfeld et al. (1996). ^cPP in IronEx-2 was digitized from the Figure 2 of Boyd (2002). ^dMesozooplankton biomass in IronEx-2 was digitized from the Figure 1 of Rollwagen Bollens and Landry (2000); Values in brackets correspond to the sampling layer. ^ePP in SOIREE was digitized from the Figure 3 of Gall et al. (2001a). ^fMesozooplankton biomass indicates copepod biomass; Values in brackets correspond to the sampling layer; After mesozooplankton biomass is the mean value averaged for the experimental period after iron addition. ^gChlorophyll-a concentrations in SOFeX-N/-S were digitized from the supplementary Figure 5 of Coale et al. (2004). ^hPP values in SOFeX-N/-S were digitized from the

Figure 4 of Coale et al. (2004). ^{¶¶¶}Fv/Fm in EIFEX was digitized from the Figure 2 of Berg et al. (2011); ^{¶¶¶}Fv/Fm in SEEDS-1 was digitized from the Figure 2 of Tsuda et al. (2003); ^{¶¶¶¶}Fv/Fm in SEEDS-2 was digitized from the Figure 6 of Tsuda et al. (2007).

[‡]Chlorophyll a concentrations in SOFeX N/S were digitized from the supplementary Figure 5 of Coale et al. (2004). ^{*}Primary productivity (PP) in IronEx 1 was estimated by multiplying PP (mg C m⁻³ d⁻¹) with the mixed layer depth (initial: 30 m and after: 35 m); ^{**}PP in IronEx 2 was digitized from the Figure 2 of Boyd (2002); ^{***}PP in SOIREE was digitized from the Figure 3 of Gall et al. (2001a); [§]PP values in SOFeX N/S were digitized from the Figure 4 of Coale et al. (2004). [‡]Mesozooplankton biomass indicates copepod biomass; Values in brackets correspond to the sampling layer; After mesozooplankton biomass is the mean value averaged for the experimental period after iron addition.

Sources are Kolber et al. (1994); Behrenfeld et al. (1996); Coale et al. (1996); Steinberg et al. (1998); Boyd et al. (2000); Rollwagen Bollens and Landry (2000); ~~Rollwagen and Landry (2000)~~; Boyd and Law (2001); Cochlan (2001); Gall et al.; (2001a); Hall and Safi (2001); Zeldis (2001); ~~Zeldis et al. (2001)~~; Boyd (2002); Gervais et al. (2002); Tsuda et al. (2003); Arrieta et al. (2004); Boyd et al. (2004); Coale et al. (2004); Oliver et al. (2004); Boyd et al. (2005); de Baar et al. (2005); Suzuki et al. (2005); Takeda and Tsuda (2005); Tsuda et al. (2005); Levasseur et al. (2006); Boyd et al. (2007); Tsuda et al. (2007); Kudo et al. (2009); Tsuda et al. (2009); Harvey et al. (2010); Berg et al. (2011); Currie et al. (2011); Peloquin et al. (2011a); Smetacek et al. (2012); Thiele et al. (2012); ~~and~~ Martin et al. (2013); and Latasa et al. (2014).

Table 5. Initial values of the export flux and the values after fertilization ($\text{mg C m}^{-2} \text{ d}^{-1}$), the corresponding depth inside and outside the fertilized patch for artificial ocean iron fertilization (aOIF) experiments, and measurement method. Values in brackets correspond to the day of measurement after fertilization.

Experiment	In-patch Initial (day)	In-patch After (day)	Outside-patch Initial (day)	Outside-patch After (day)	Depth (m)	Method
1 IronEx-1						
2 IronEx-2	84 (0)	600 (7-14 10)			25	Water-column ^{234}Th
3 SOIREE		~87			100	Water-column ^{234}Th
		185 (11–13)	146 (0–2)	78 (11–13)	110	Drifting trap
		74 (11–13)	73 (0–2)	38 (11–13)	310	Drifting trap
4 EisenEx						
5 SOFeX-N						
6 SOFeX-S	36 (5)	112 (27)	48 (6)	49 (26)	50	Water-column ^{234}Th
	19 (5)	142 (27)	38 (6)	56 (26)	100	Water-column ^{234}Th
7 EIFEX	*~340 (0) ^a	*~1692 (32) ^a	*~396 (0) ^a	*~516 (32) ^a	100	Water-column ^{234}Th
8 SAGE						
9 LOHAFEX	**~ 60-62 (0) ^b	**~94 (25) ^b	**~ 787 (4) ^b	**~ 9754 (2334) ^b	100	Water-column ^{234}Th
	~6 (0–2) ^c	~5 (13–15) ^c		~29 (26–27) ^c	200	Neutrally buoyant sediment trap
		~12 (28–37) ^c		~11 (24–29) ^c	450	Neutrally buoyant sediment trap
10 SEEDS-1	234 (1–3)	141 (12–14)	148 (1–6)	154 (10–14)	40	Drifting trap
	100 (0–2)	423 (9–13)			50	*Water-column ^{234}Th ^d
	68 (1–3)	85 (12–14)	61 (1–6)	91 (10–14)	100	Drifting trap
	121 (0–2)	460 (2–9)			200	Water-column ^{234}Th
11 SERIES	*~ 12038 (3) ^e	480 (24)	192 (3)	139 (15)	50	Drifting trap
	*~48 (3) ^e	*~192 (24) ^e			100	Drifting trap
12 SEEDS-2	290 (1–4)	580 (19–22)	300 (1–8)	509 (18–31)	40	Drifting trap
	316 (1–4)	337 (19–22)	213 (1–8)	204 (18–31)	100	Drifting trap
13 FeeP						

5 ^a*Export flux in EIFEX was digitized from the supplementary Figure 5.1 of Smetacek et al. (2012); ^b**Export flux in LOHAFEX was digitized from the Figure 4 of Martin et al. (2013); ^c*Export flux in LOHAFEX was digitized from the Figure 6 of Martin et al. (2013). ^d*Export flux in SEEDS-1 was determined from the suspended particles; ^e*Export flux in SERIES was digitized from the Figure 2 of Boyd et al. (2004).

Sources are Bidigare et al. (1999); Charette and Buesseler (2000); Nodder and Waite (2001); Boyd et al. (2004); Aono et al. (2005); Buesseler et al. (2005); Aramaki et al. (2009); Smetacek et al. (2012); and Martin et al. (2013).

Figure Captions

Fig. 1. Diagram showing the monthly atmospheric CO₂ concentrations (ppm) (blue) measured at the Mauna Loa Observatory, Hawaii (<http://www.esrl.noaa.gov/gmd/ccgg/trends/data.html>), global monthly land-surface air and sea surface temperature anomalies (°C) (red) (<http://data.giss.nasa.gov/gistemp/>), and pH (green) measured at ~~Station ALOHA~~ ALOHA station in the central ~~North~~ Pacific (http://hahana.soest.hawaii.edu/hot/products/HOT_surface_CO2.txt). The data values represent moving average values for 12 months and shading indicates the standard deviation for 12 months.

Fig. 2. Schematic representation of several proposed climate-engineering methods (modified from Matthews (1996)).

Fig. 3. The iron hypothesis, as suggested by Martin (1990). (a) Effectiveness of the biological pump under normal conditions, (b) Effectiveness of the biological pump following iron enrichment (modified from Sarmiento and Gruber (2006)), and (c) Schematic diagram of the decrease in the downward flux of organic carbon as a function of depth in the water column (modified from Lampitt et al. (2008)). OM is organic matter and DIC is dissolved inorganic carbon.

Fig. 4. Global annual distribution of surface (a) Chlorophyll concentrations (mg m⁻³), (b) Nitrate concentrations (μM), and (c) Silicate concentrations (μM). The chlorophyll-a concentration distribution was obtained from the Aqua MODIS chlorophyll-a composite from July 2002 to February 2016 (<https://oceancolor.gsfc.nasa.gov/cgi/l3>), nitrate and silicate were obtained from the World Ocean Atlas 2013 dataset (<https://odv.awi.de/en/data/ocean/world-ocean-atlas-2013>) and plotted using Ocean Data View (Schlitzer, 2017). The white circles indicate the locations of 13 artificial ocean iron fertilization (aOIF) experiments and the black triangles indicate the locations of ~~two~~ six natural OIF (nOIF) experiments. Note that the numbers indicate the order of the aOIF experiments and the Roman-numerals indicate the order of the nOIF experiments (see Table 1).

Fig. 5. Photographs of the iron addition procedure. Panels a-f taken during the European Iron Fertilization Experiment (EIFEX), Surface Ocean Lower Atmosphere Study (SOLAS) Air-Sea Gas Exchange (SAGE), and Indo-German iron fertilization experiment (LOHAFEX): (a) Iron-(II) sulfate bags. (b) The funnel used to pour iron and hydrochloric acid. (c) Tank system used for mixing Iron(II) sulfate, hydrochloric acid, and seawater (Smetacek, 2015). (d) Preparation for release: the deck of RV *Tangaroa* with the iron tanks on the left and the SF₆ tracer tanks on the right (Photo: Matt Walkington) (<https://www.niwa.co.nz/coasts-and-oceans/research-projects/sage>). (e) Outlet pipe connected to the tank system. (f) Pumping iron into the prop wash during EIFEX (Smetacek, 2015).

Fig. 6. (a) Maximum (bar with dotted line) and initial (bar with solid line) patch size (km²) during artificial ocean iron fertilization (aOIF) experiments. (b) ~~Total (bar with dotted line) and initial First target iron concentrations (bar with solid line) iron(II)-added (nMkg)~~. (c) Maximum (bar with dotted line) and minimum (bar with solid line) ~~mixed layer depth (MLD mixed layer depth) (MLD, m)~~ during aOIF experiments. (d) Initial sea surface temperature (SST, °C). (e) Initial nitrate concentrations (μM). (f) Initial silicate concentrations (μM). (g) Initial Fv/Fm ratios. (h) Initial chlorophyll-a concentrations (mg m⁻³). Note that the numbers on the X axis indicate the order of aOIF experiments as given in Fig. 4 and Table 1 and are grouped according to ocean basins; Equatorial Pacific (EP) (yellow bar), Southern Ocean (SO) (blue bar), subarctic North Pacific (NP) (red bar), and subtropical North Atlantic (NA) (green bar). Sources are Kolber et al. (1994); Martin et al. (1994); Behrenfeld et al. (1996); Coale et al. (1996); Coale et al. (1998); Steinberg et al. (1998); Boyd et al. (2000); Boyd and Law (2001); Gall et al. (2001b); Gervais et al. (2002); Law et al. (2003); Tsuda et al. (2003); Coale et al. (2004); Turner et al. (2004); Bakker et al. (2005); Boyd et al. (2005); Bozec et al. (2005); de Baar et al. (2005); Hiscock and Millero (2005); Takeda and Tsuda (2005); Tsuda et al. (2005); Tsumune et al. (2005); Law et al. (2006); Marchetti et al. (2006); Boyd et al. (2007); Rees et al. (2007); Tsuda et al. (2007); Suzuki et al. (2009); Tsumune et al. (2009); Harvey et al. (2010); Smetacek and Naqvi (2010); Berg et al. (2011); Hadfield (2011); Law et al. (2011); Peloquin et al. (2011a); Smetacek et al. (2012); Thiele et al. (2012); Martin et al. (2013); Ebersbach et al. (2014); and Latasa et al. (2014).

Fig. 7. (a) Maximum (bar with dotted line) and initial (bar with solid line) Fv/Fm ratios during artificial ocean iron fertilization (aOIF) experiments. (b) Changes in nitrate concentrations ($\Delta\text{NO}_3^- = [\text{NO}_3^-]_{\text{post-fertilization (postf)}} - [\text{NO}_3^-]_{\text{pre-fertilization (pref)}}$; μM). (c) Maximum (bar with dotted line) and initial (bar with solid line) chlorophyll-a concentrations (mg m⁻³). (d) Distributions of chlorophyll-a concentrations (mg m⁻³) on day 24 after iron addition in the Southern Ocean iron experiment-north (SOFEX-N) from MODIS Terra Level-2 daily image and on day 20 in the SOFeX-south (SOFEX-S) from SeaWiFS Level-2 daily image (white dotted box indicates phytoplankton bloom during aOIF experiments). (e) Changes in primary productivity (PP) ($\Delta\text{PP} = [\text{PP}]_{\text{postf}} - [\text{PP}]_{\text{pref}}$; mg C m⁻² d⁻¹). (f) Changes in partial pressure of CO₂ ($p\text{CO}_2$) ($\Delta p\text{CO}_2 = [p\text{CO}_2]_{\text{postf}} - [p\text{CO}_2]_{\text{pref}}$; μatm). The color bar indicates changes in dissolved inorganic carbon (DIC) ($\Delta\text{DIC} = [\text{DIC}]_{\text{postf}} - [\text{DIC}]_{\text{pref}}$; μM). Note that the PP (mg C m⁻² d⁻¹) of aOIF experiment number 1 (IronEx-1) was estimated by multiplying the

PP (mg C m⁻³ d⁻¹) with the mixed layer depth (initial: 30 m and after: 35 m). The numbers on the X axis indicate the order of aOIF experiments as given in Fig. 4 and Table 1 and are grouped according to ocean basins; Equatorial Pacific (EP) (yellow bar), Southern Ocean (SO) (blue bar), subarctic North Pacific (NP) (red bar), and subtropical North Atlantic (NA) (green bar). Sources are Kolber et al. (1994); Martin et al. (1994); Behrenfeld et al. (1996); Coale et al. (1996); Steinberg et al. (1998); Boyd et al. (2000); Boyd and Law (2001); Frew et al. (2001); Gall et al. (2001a); Boyd (2002); Gervais et al. (2002); Tsuda et al. (2003); Coale et al. (2004); Boyd et al. (2004); Bakker et al. (2005); Boyd et al. (2005); [Bozec et al. \(2005\)](#); de Baar et al. (2005); Hiscock and Millero (2005); Smetacek et al. (2005); Takeda and Tsuda (2005); Tsuda et al. (2005); Wong et al. (2006); Boyd et al. (2007); Tsuda et al. (2007); Kudo et al. (2009); Tsumune et al. (2009); Harvey et al. (2010); Smetacek and Naqvi (2010); Berg et al. (2011); Currie et al. (2011); Law et al. (2011); Peloquin et al. (2011a); Smetacek et al. (2012); Thiele et al. (2012); Assmy et al. (2013); Martin et al. (2013); ~~and~~ Ebersbach et al. (2014); [and Latasa et al. \(2014\)](#).

Fig. 8. (a) Time-series of particulate organic carbon (POC) fluxes estimated from the water-column based ²³⁴Th method (mg C m⁻² d⁻¹) of the upper 100-m layer inside (red bar) and outside the fertilized patch (blue bar) during the European Iron Fertilization Experiment (EIFEX) (modified from Smetacek et al. (2012)). (b) Time-series of vertically integrated ²³⁴Th (dpm l⁻¹) inside (red circles) and outside the fertilized patch (blue diamonds) relative to the parent ²³⁸U (dpm l⁻¹; dotted black line) during the Southern Ocean Iron Release Experiment (SOIREE) (modified from Nodder et al. (2001)).

Fig. 9. Assessment framework for scientific research involving ocean fertilization (OF) (modified from Resolution LC-LP.2, (2010)).

Fig. 10. (a) Time-series of mixed layer depth-integrated chlorophyll-a concentrations (mg m⁻²) during the Southern Ocean Iron Release Experiment (SOIREE) (brown line), Subarctic Pacific iron Experiment for Ecosystem Dynamics Study-1 (SEEDS-1) (coral line), Subarctic Ecosystem Response to Iron Enrichment Study (SERIES) (cyan line), SEEDS-2 (blue line), and European Iron Fertilization Experiment (EIFEX) (teal line). (b) The distributions of chlorophyll-a concentrations (mg m⁻³) on day 5 and day ~~45-42~~ during SOIREE from SeaWiFS Level-2 daily images. Sources are [Gall et al. \(2001b\)](#); ~~Boyd and Abraham (2001)~~; Tsuda et al. (2007); and Assmy et al. (2013).

Fig. 11. Schematic diagram of the Korean Iron Fertilization Experiment in the Southern Ocean (KIFES) representing the experiment target site (eddy structure) and survey methods (underway sampling systems, multiple sediment traps, sub-bottom profilers, sediment coring systems, and satellite observations).

Fig. 1

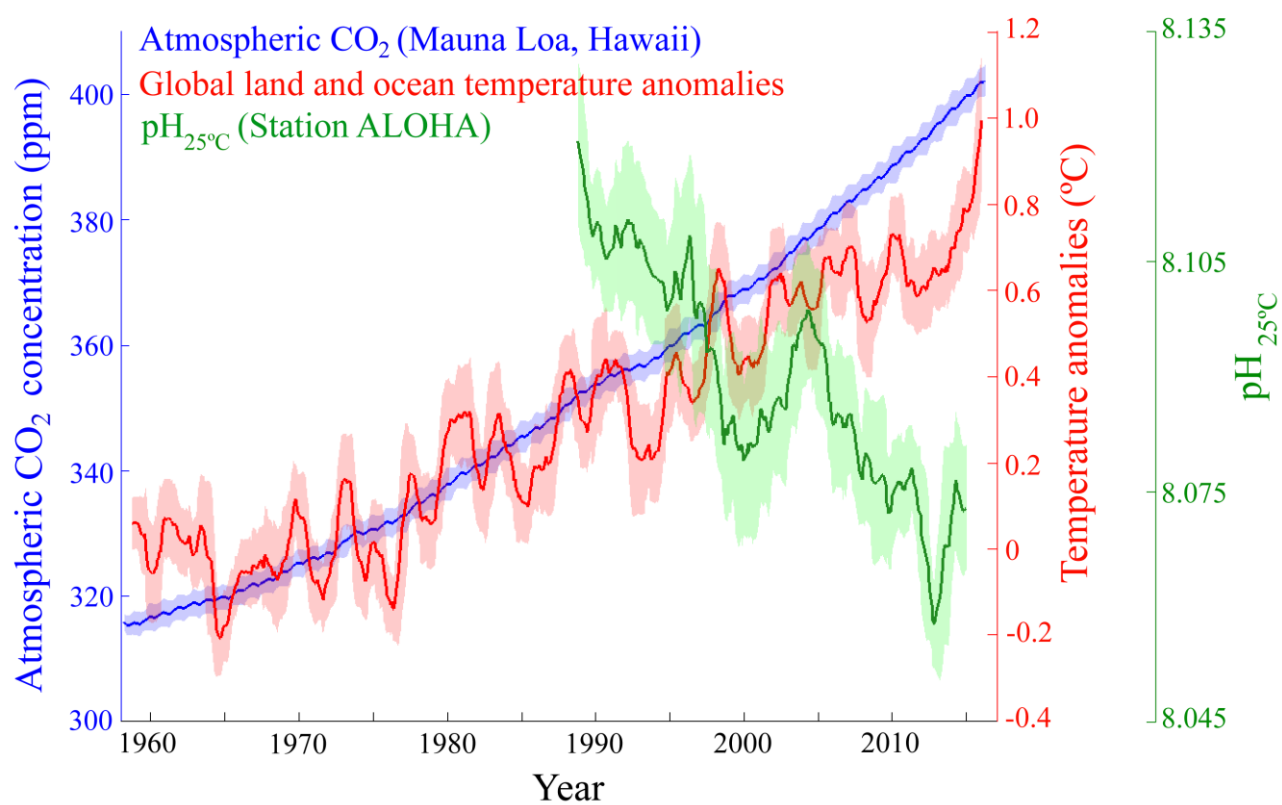


Fig. 2

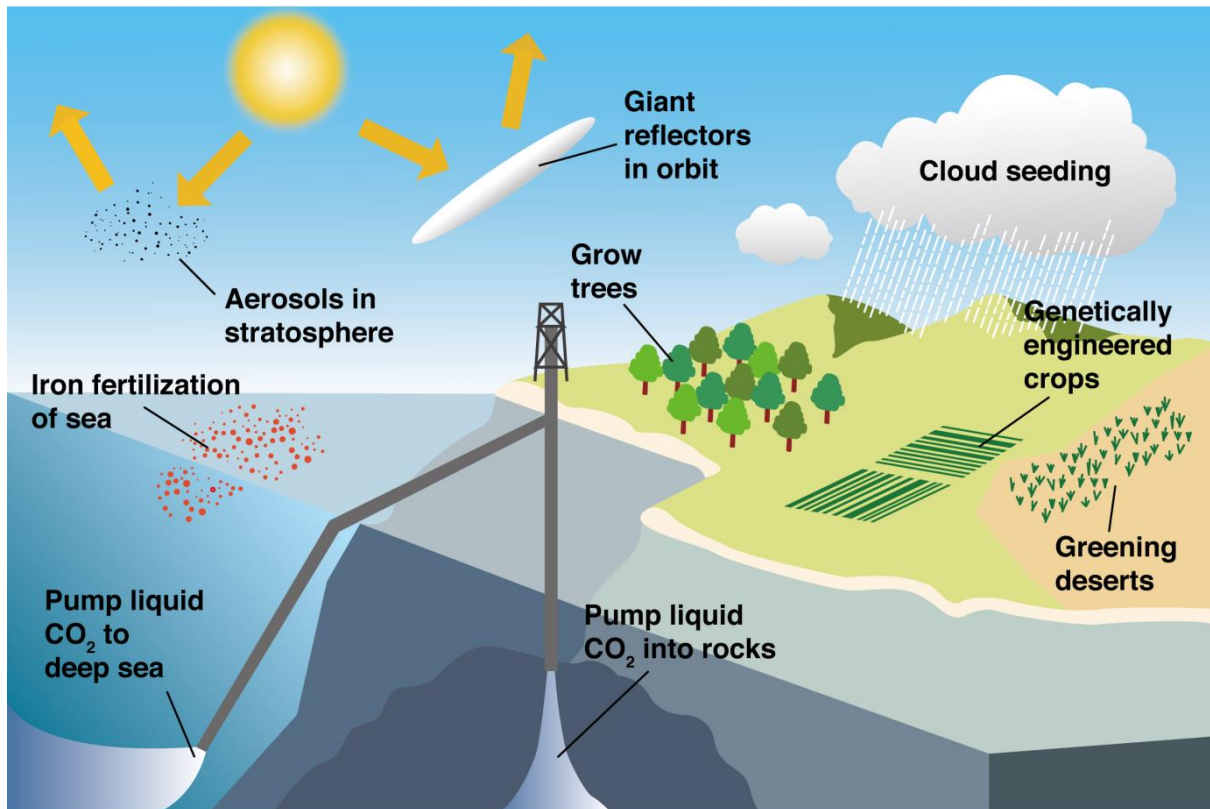


Fig. 3

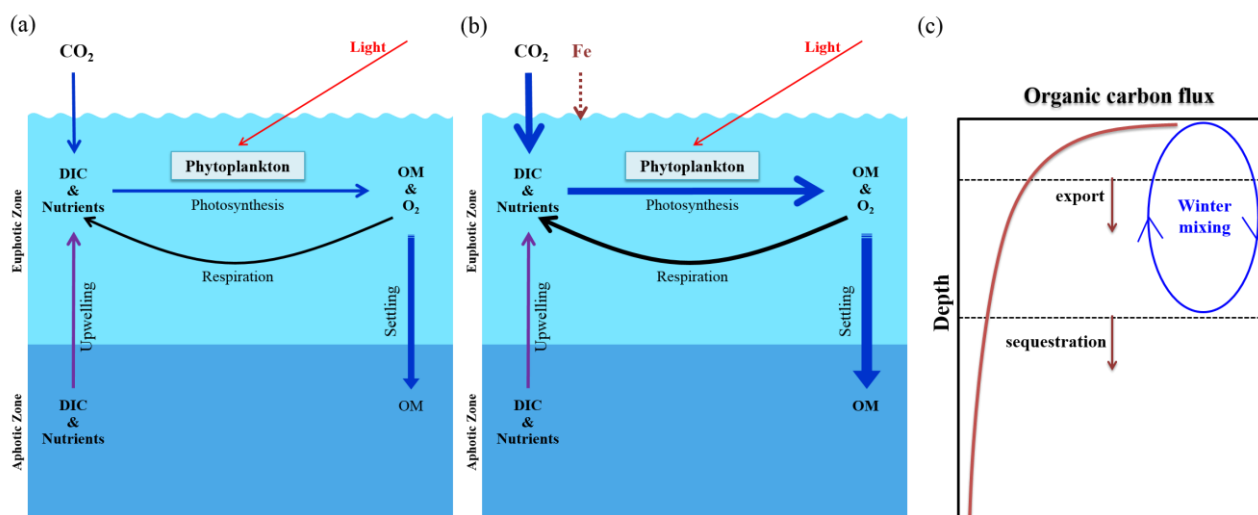
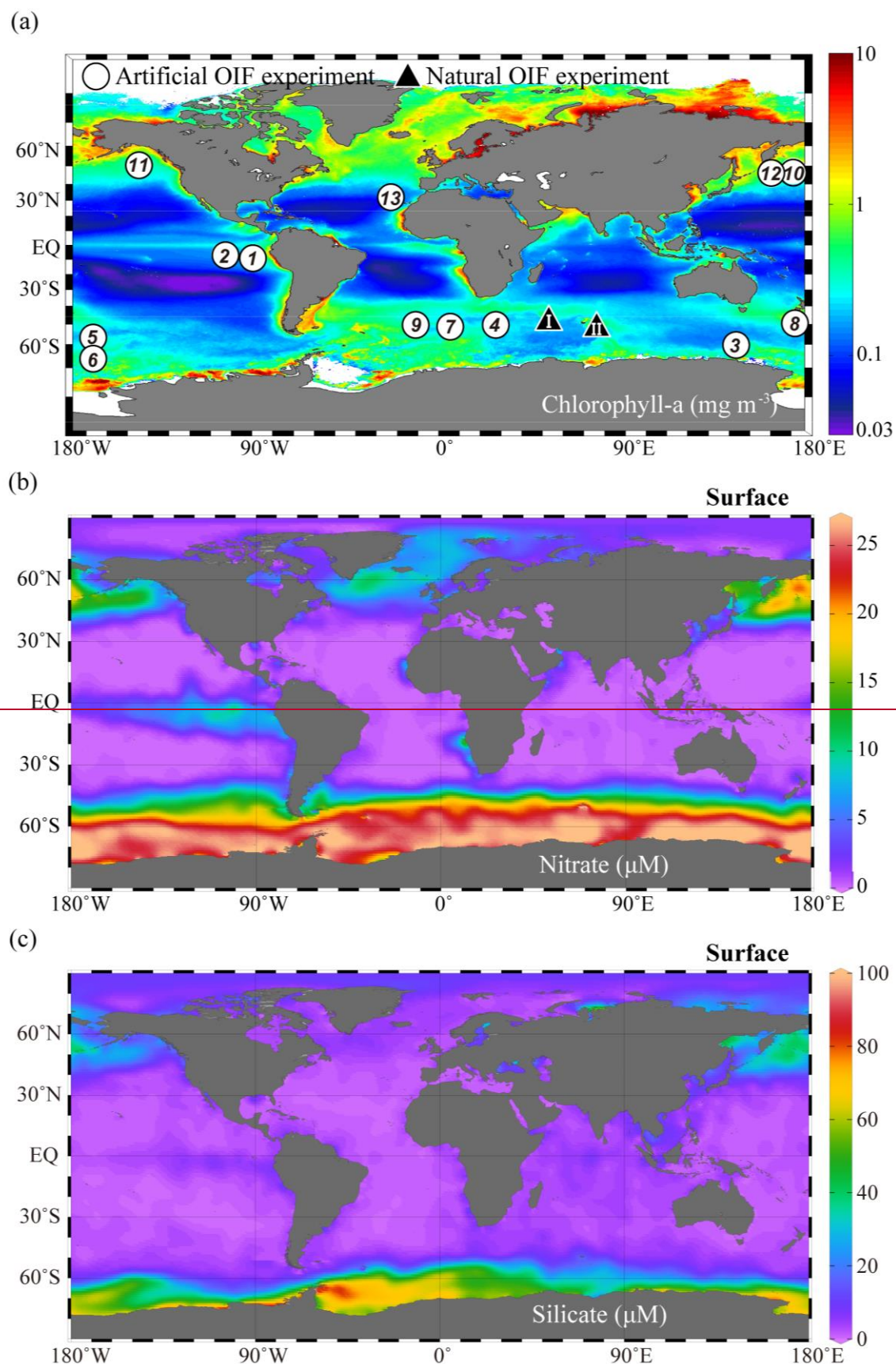
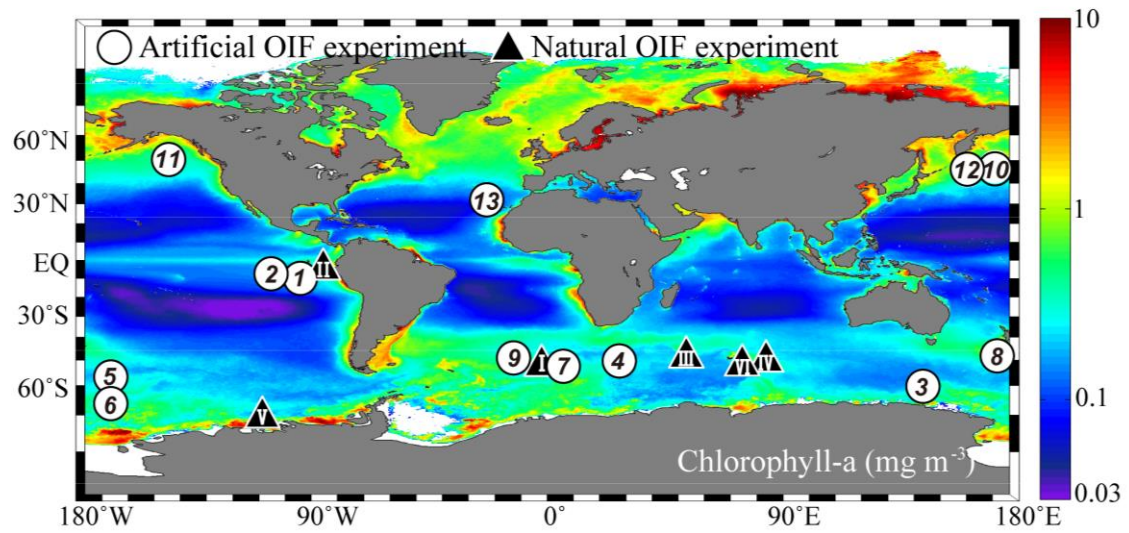


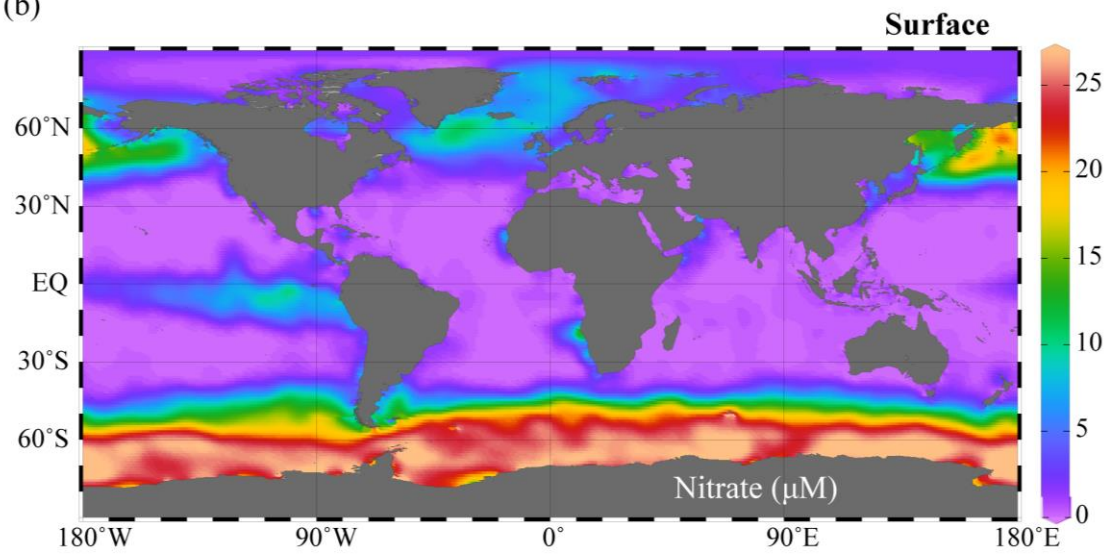
Fig. 4



(a)



(b)



(c)

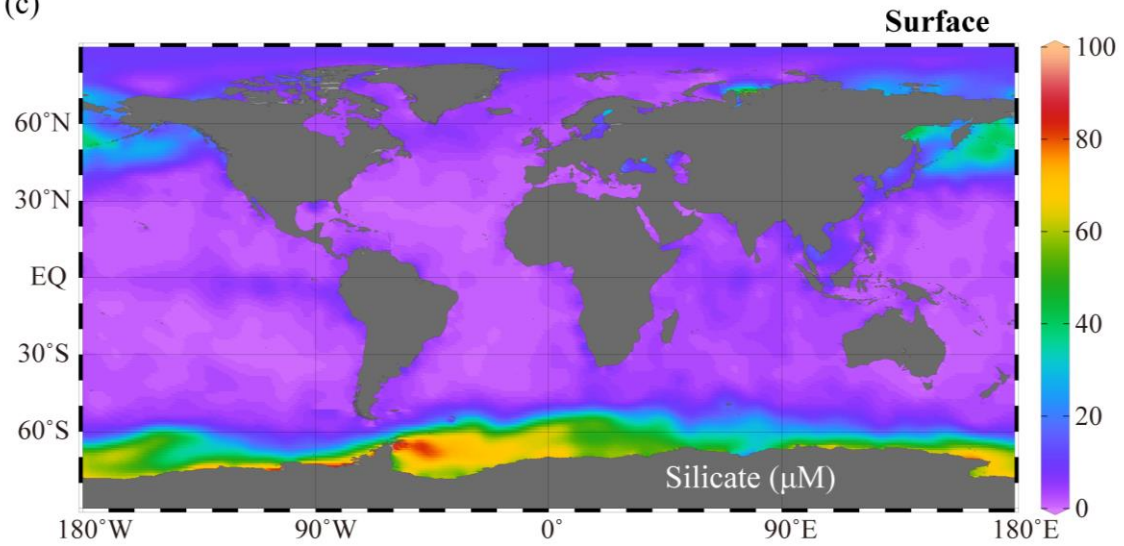
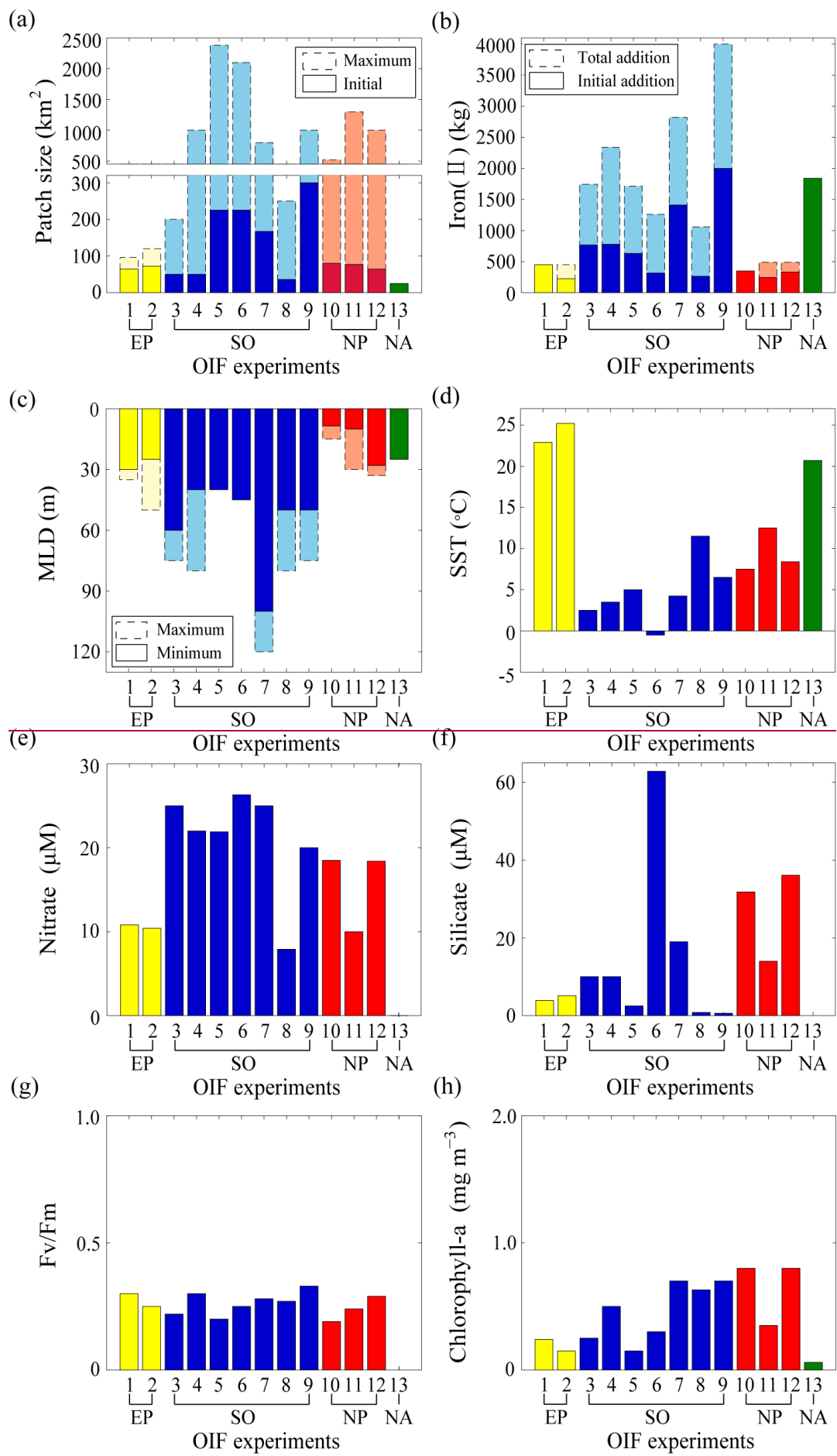
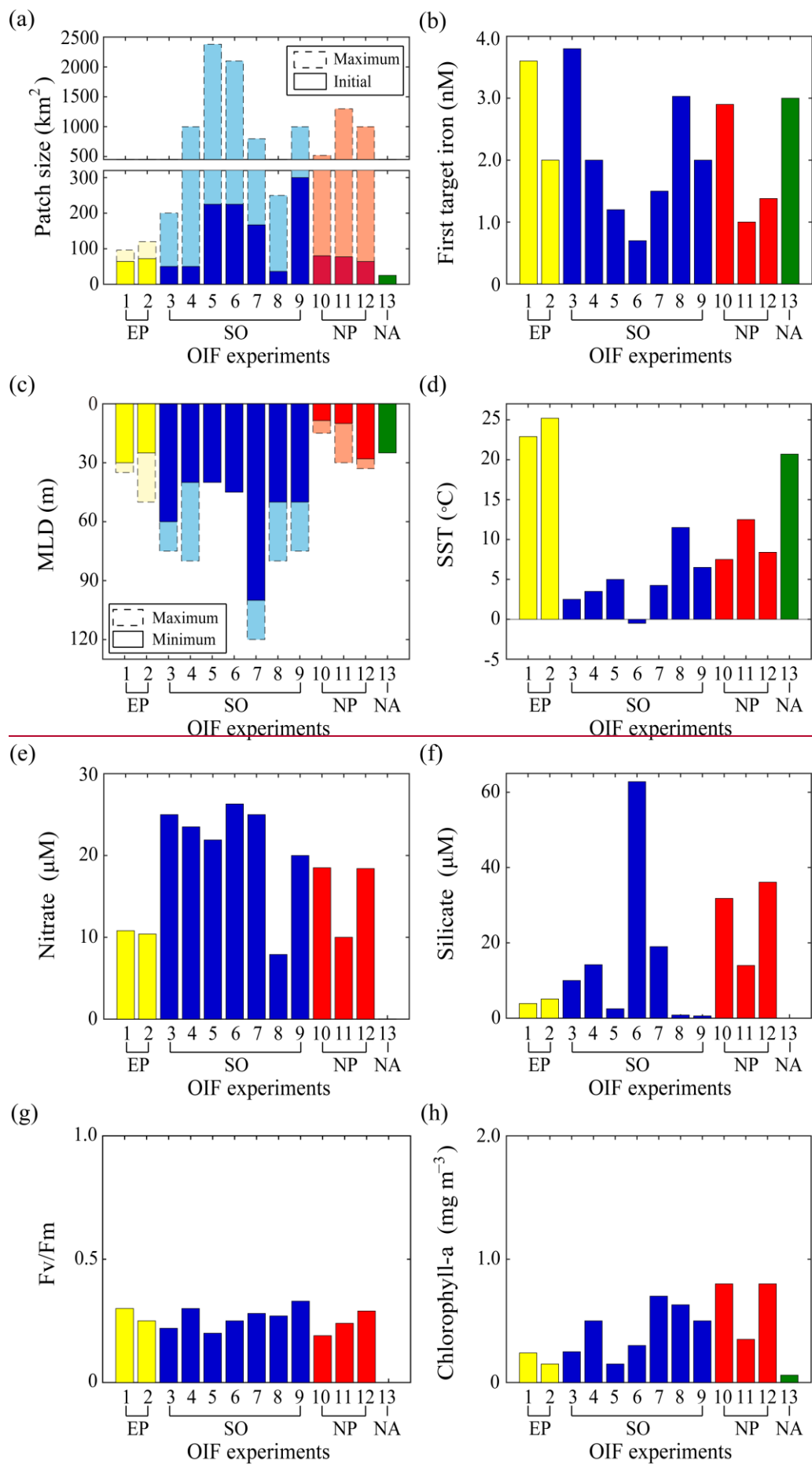
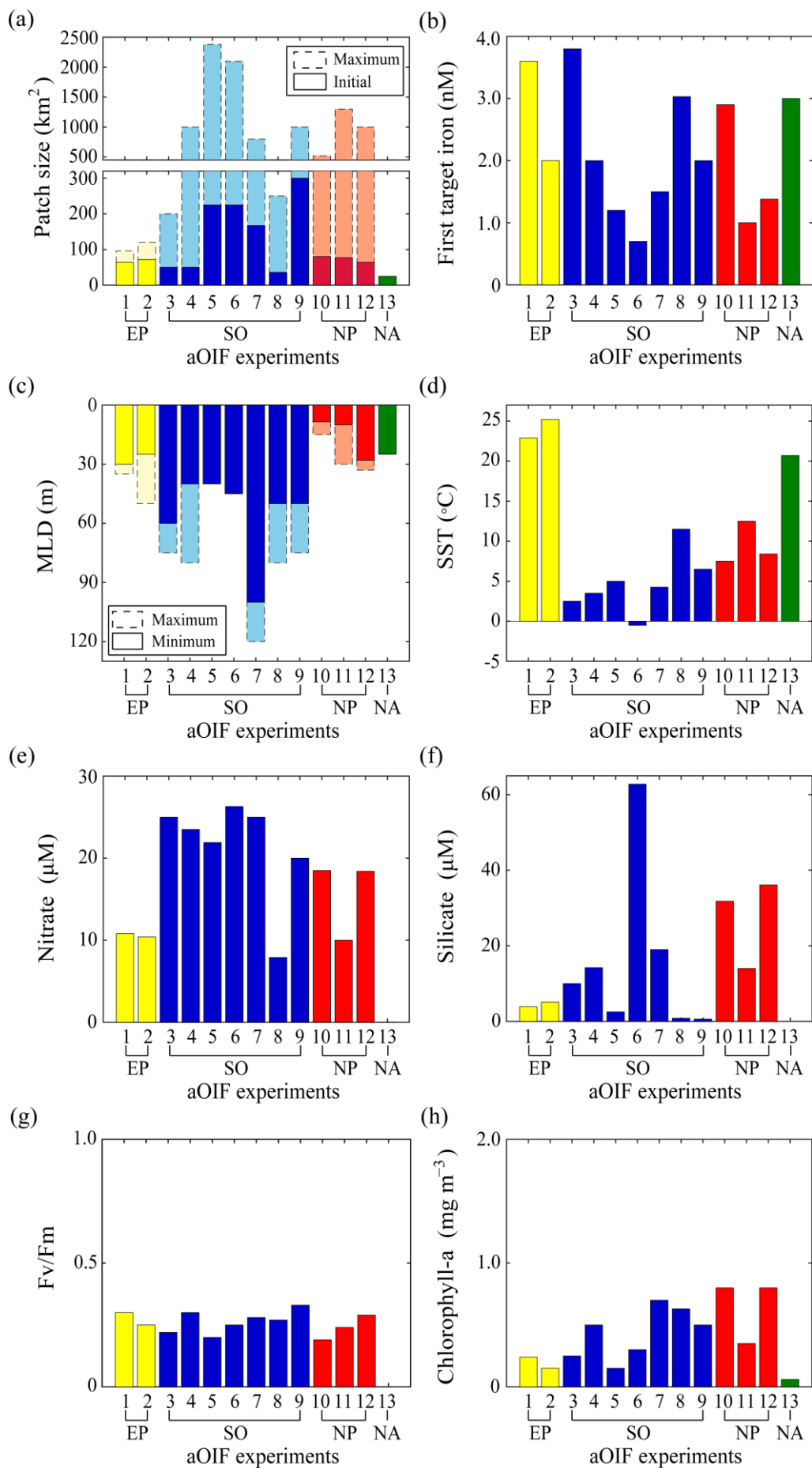


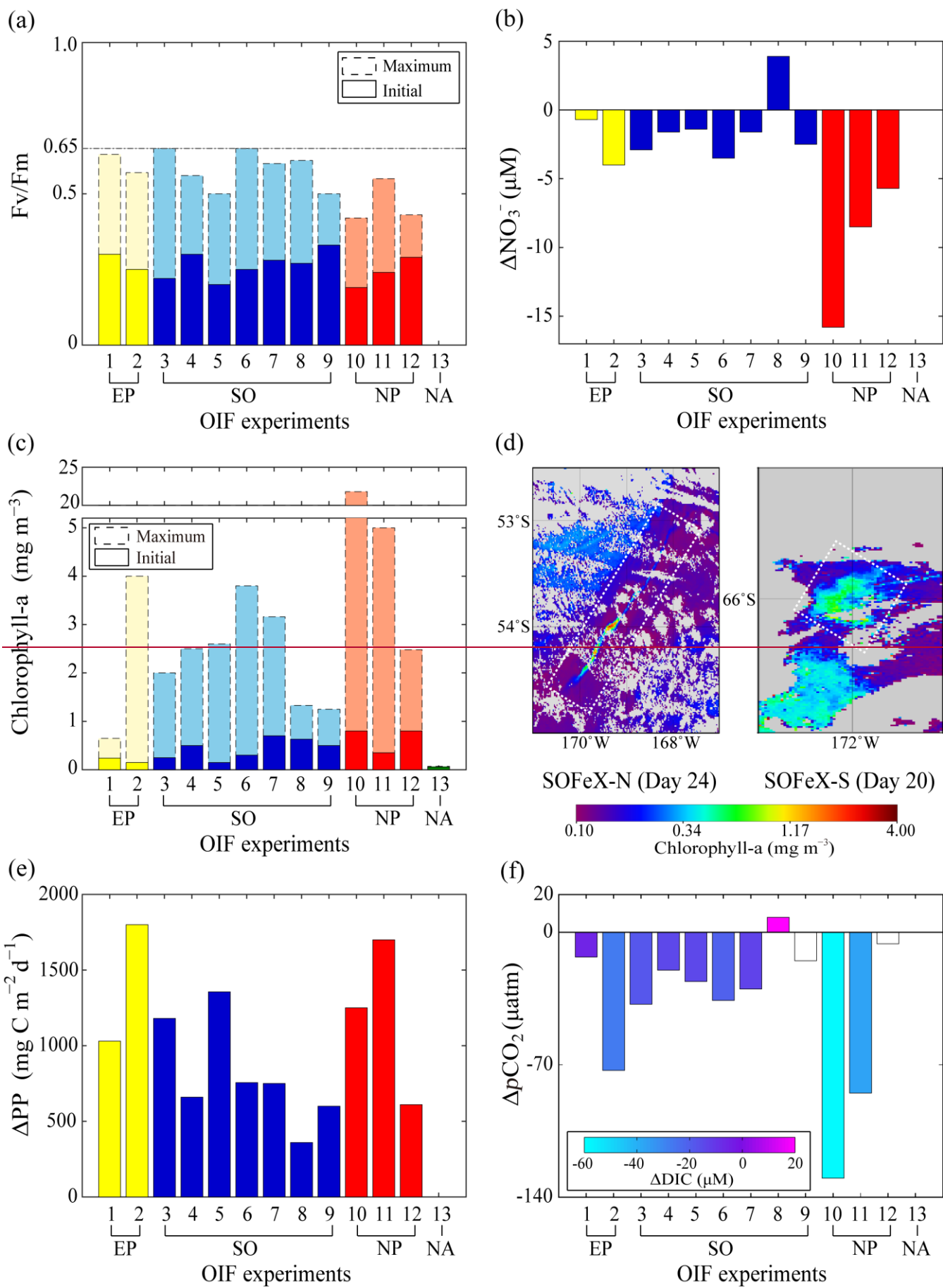
Fig. 5

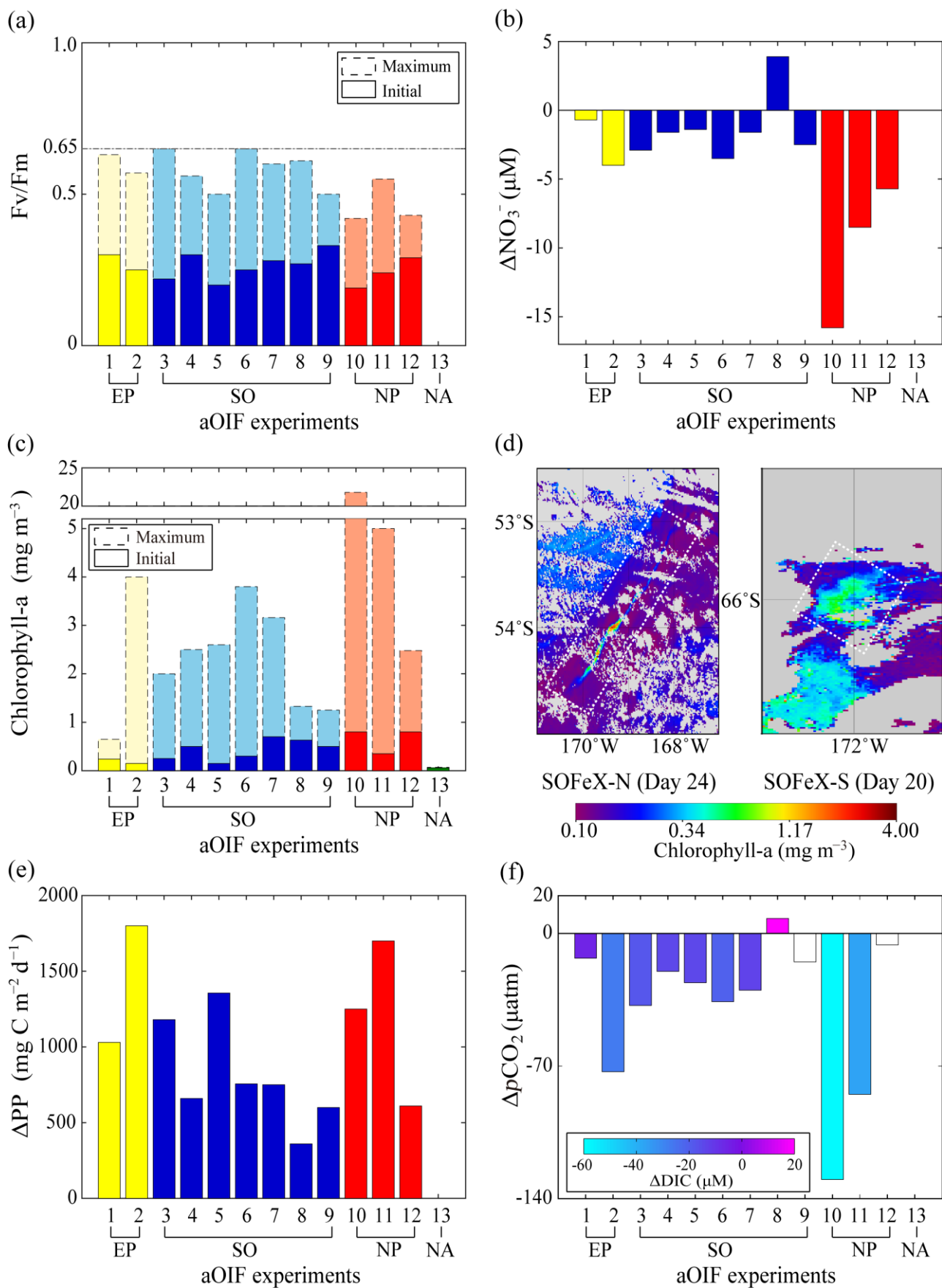


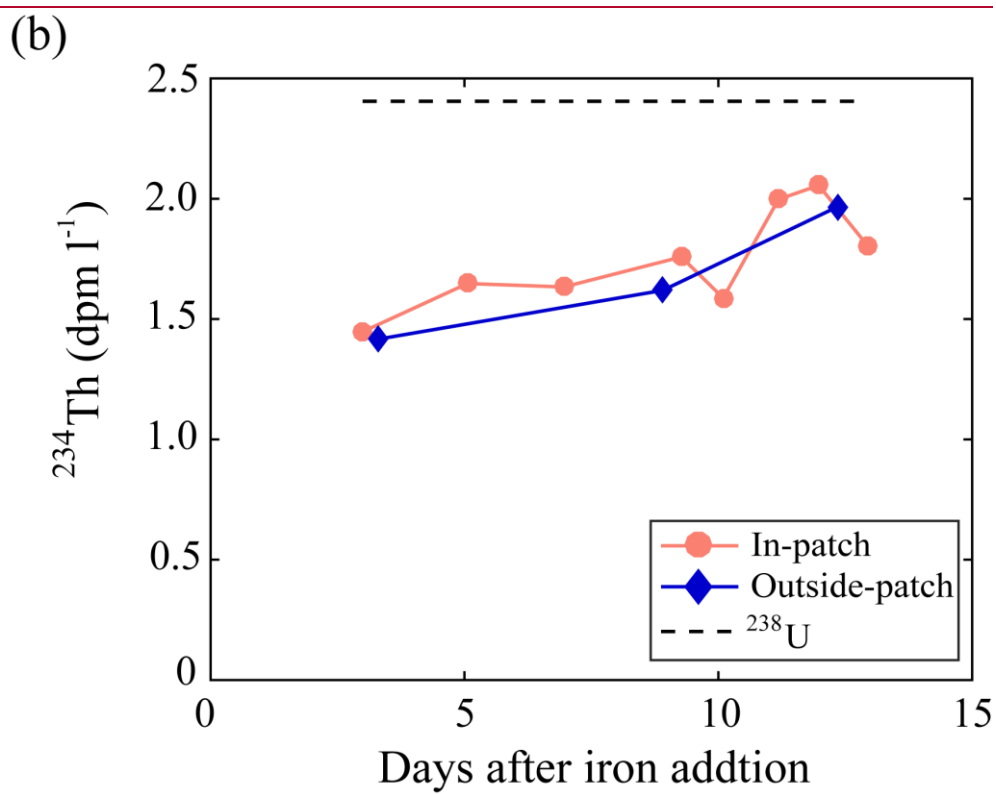
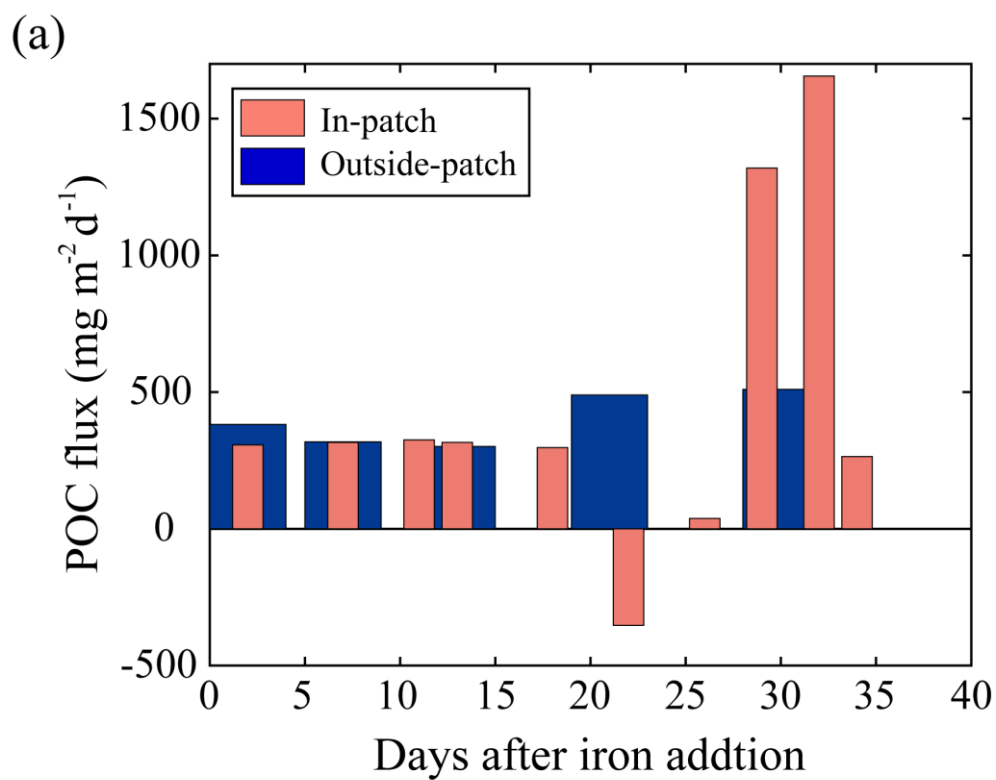


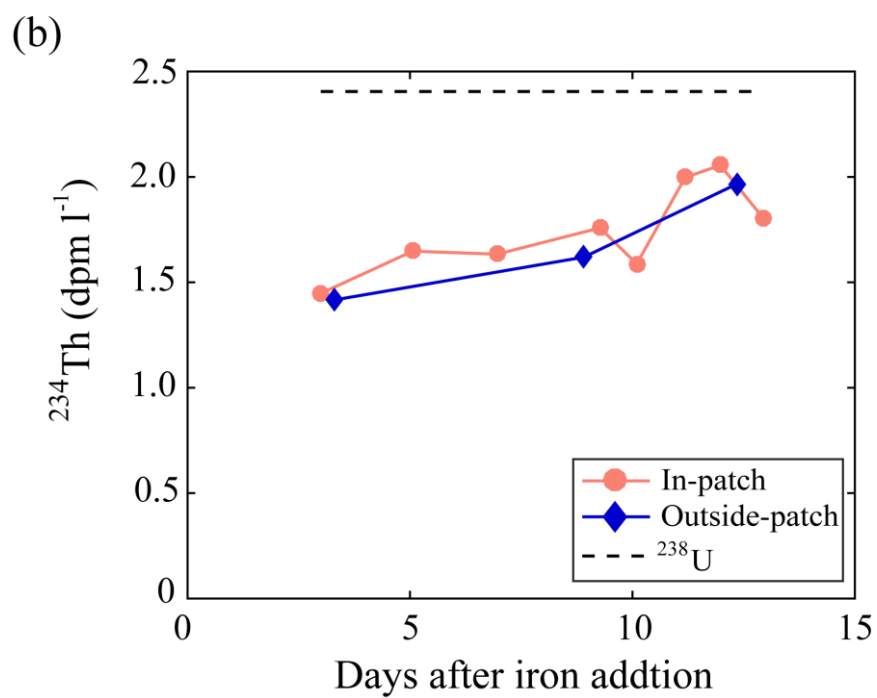
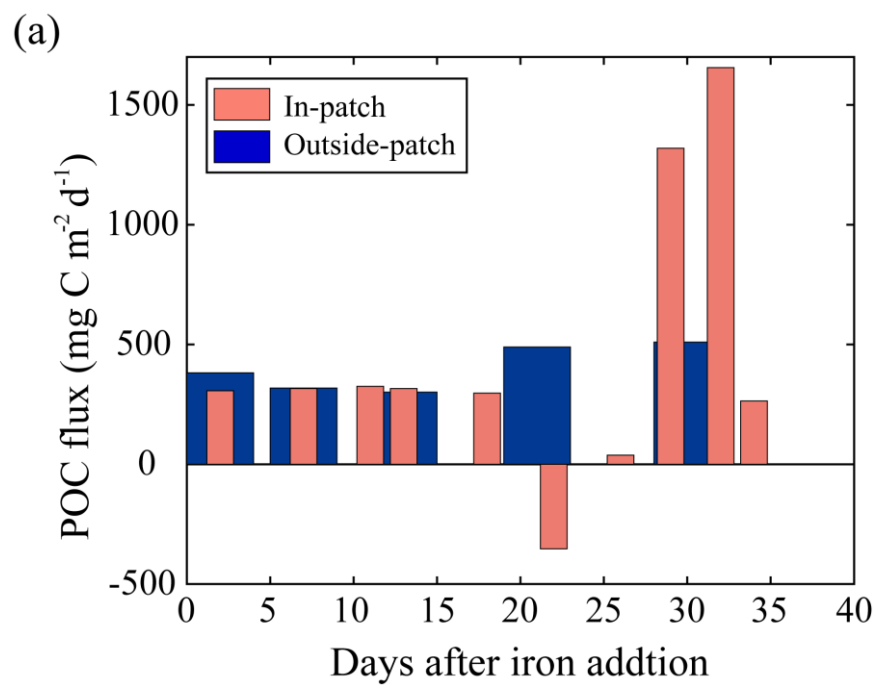


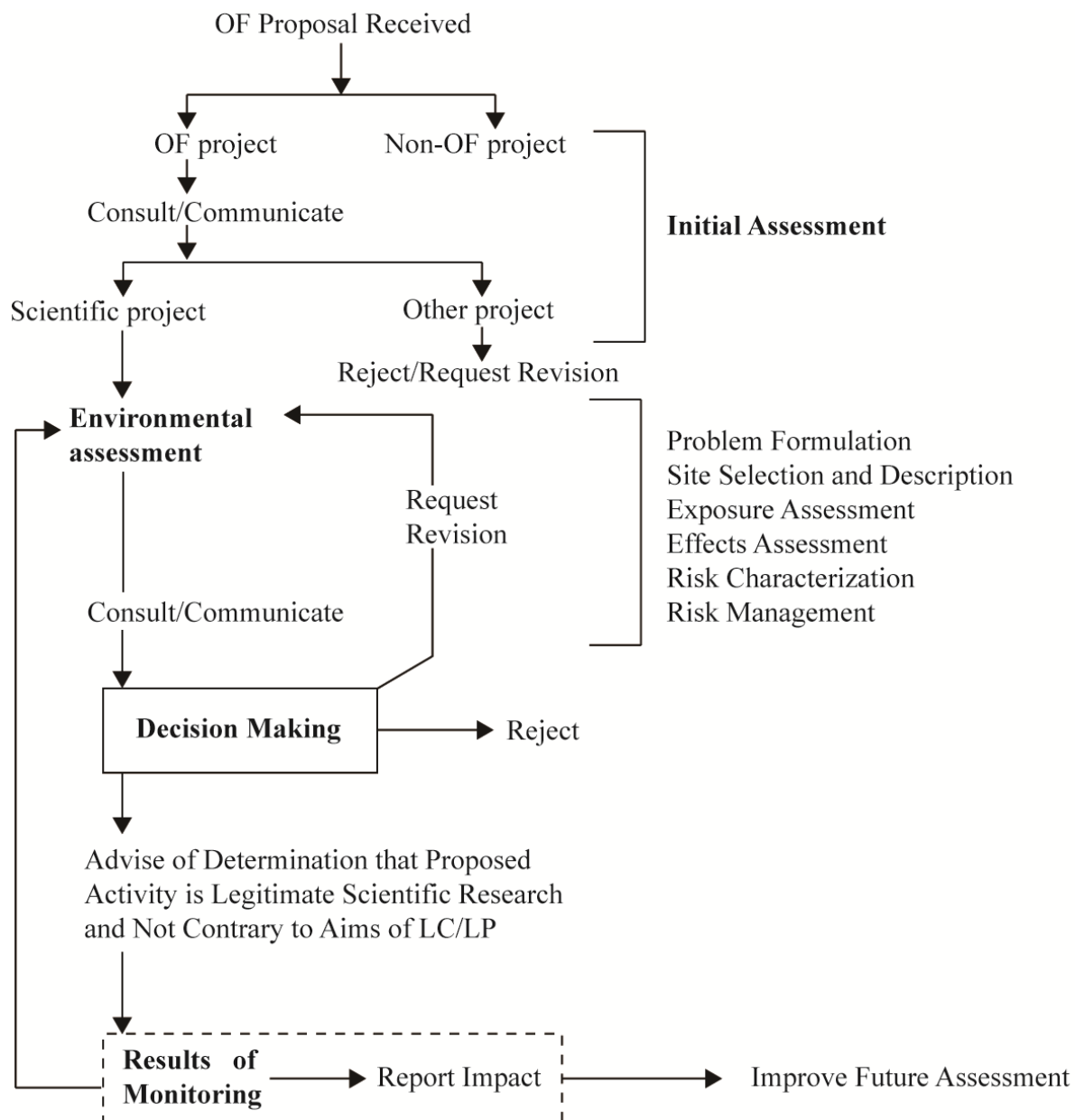












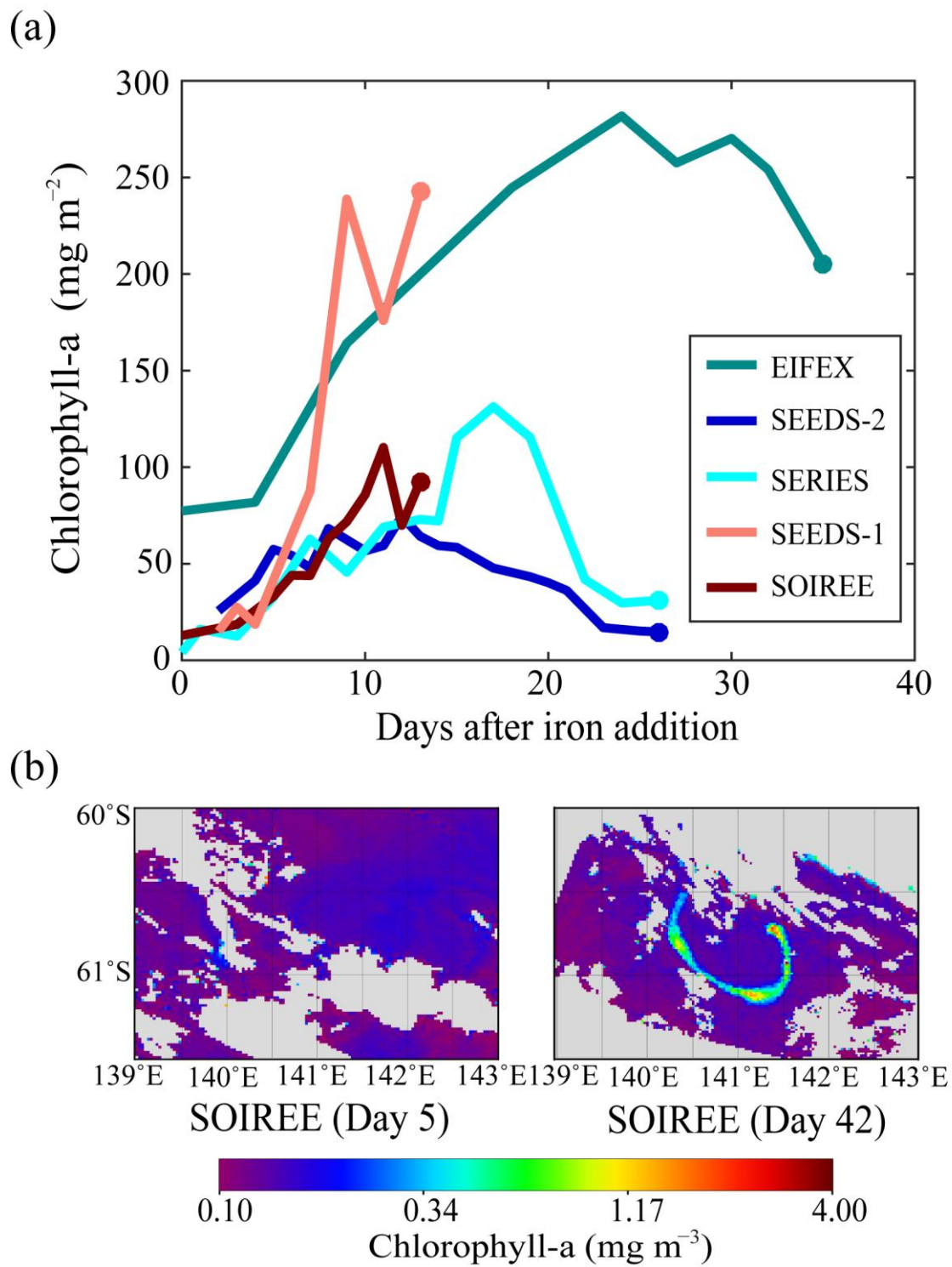


Fig. 11

