

**Zinc stimulation of phytoplankton in a low carbon dioxide, coastal Antarctic environment:
evidence for the Zn hypothesis**

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Abstract.

The ocean acts as a carbon sink, absorbing carbon from the atmosphere and resulting in substantial uptake of anthropogenic CO₂ emissions. As biological processes in the oceans such as net primary production (NPP) contribute significantly to this sink, understanding how they will shift in response to increasing atmospheric CO₂ is necessary to project future ocean carbon storage capacity. Macronutrient and micronutrient resource limitation within the oceans regulates NPP, and while some micronutrients such as zinc (Zn) are present at very low concentrations, their ability to limit NPP has remained unclear. Zn is a key micronutrient used by phytoplankton for a multitude of metabolic functions, yet there have been few observations of its influence on natural oceanic phytoplankton populations. In this study, we observed Zn limitation of growth in the natural phytoplankton community of Terra Nova Bay, Antarctica, in addition to primary iron (Fe) limitation. Shipboard incubation experiments amended with Zn and Fe resulted in significantly higher chlorophyll *a* content and dissolved inorganic carbon drawdown compared to Fe addition alone. Zn and Fe stress response proteins detected in incubation and environmental biomass provided independent verification of algal co-stress for these micronutrients. We consider total biomass and low surface ocean pCO₂ as potential drivers of environmental Zn stress. This study definitively establishes that Zn limitation can occur in the modern oceans, opening up new possibility space in our understanding of nutrient regulation of NPP through geologic time, and we consider the future of oceanic Zn limitation in the face of climate change.

1 Introduction

46 Primary productivity in the oceans is a key component of the global carbon cycle and is
47 largely controlled by the availability of nitrogen (N), phosphorus (P), and iron (Fe). Yet there is
48 increasing evidence that other micronutrients such as zinc (Zn), cobalt (Co), and vitamin B₁₂ can
49 also influence phytoplankton productivity, often as secondary limiting nutrients after N, P, or Fe
50 are added (Moore et al. 2013; Browning and Moore 2023). Zn can be particularly scarce in the
51 photic zone (Bruland 1980; Jakuba et al. 2012) where total dissolved Zn (dZn_T) can be below 0.2
52 nM in seawater due to biological uptake and complexation by organic ligands (Bruland 1989;
53 Lohan et al. 2002; Baars and Croot 2011; Middag et al. 2019), which further lowers Zn
54 bioavailability (Sunda and Huntsman 2000; Saito et al. 2008; Lhospice et al. 2017). Marine
55 eukaryotic algae and copiotrophic bacteria possess a large metabolic demand for Zn that is on
56 par with that of Fe (Sunda and Huntsman 2000; Mazzotta et al. 2021).

57 Vertical profiles of dZn in the Southern Ocean have been measured previously. Zn has
58 not historically been considered as a limiting micronutrient in the Southern Ocean due to the
59 upwelling of nutrient-rich waters that bring dZn to nanomolar concentrations only a couple
60 hundred meters below the surface. Yet nutrient-like profiles of dZn are evident throughout this
61 region, with surface depletion due to biological uptake decreasing this large inventory in the
62 upper water column (Fitzwater et al. 2000; Coale et al. 2005; Baars and Croot 2011; Sieber et al.
63 2020; Kell et al. 2024). ~~(Coale et al. 2005; Baars and Croot 2011)~~. Additionally, both model-
64 based estimates (Roshan et al. 2018) and direct field measurements (Kell et al. 2024) of Zn
65 uptake in this region have demonstrated a substantial biological demand for Zn in surface waters,
66 leading to significant dZn drawdown. This is consistent with and genomic and laboratory studies
67 indicating an elevated Zn demand in polar phytoplankton (Twining and Baines 2013; Ye et al.
68 2022).

69 -Despite the scarcity of bioavailable Zn in the surface ocean and its high cellular demand,
70 relatively few experimental studies have examined the ability of Zn addition to stimulate natural
71 phytoplankton communities (**Supplementary Table 1**). These results have been variable with
72 findings that include negative results (Scharek et al. 1997; Coale et al. 2003; Ellwood 2004),
73 slight Zn stimulatory results (Crawford et al. 2003b), a “very small increase” relative to controls
74 in an unreplicated experiment (Coale et al. 2003), Zn stimulation within Fe and Si uptake
75 experiments (Franck et al. 2003), Zn primary and secondary limitation in the North Pacific in an
76 unreplicated experiment (Jakuba et al. 2012), secondary Zn limitation after primary Si limitation
77 in the Costa Rica Dome (Dreux Chappell et al. 2016), and enhanced Zn uptake rates under low
78 pCO₂ (Xu et al. 2012). Whether due to the early negative results, the few positive findings, or the
79 practical constraints of co-limitation studies in the field that limit the number of micronutrients
80 that can be tested, ~~a general prejudice against considering additional factors in controlling marine~~
81 ~~productivity,~~ it is our experience that there is currently no broad community recognition that zinc
82 limitation is a process that could affect primary productivity in any region of the oceans, leaving
83 the original ‘zinc hypothesis’ unresolved (Morel et al. 1994).

84 In ~~contrast~~juxtaposition, laboratory studies have unequivocally demonstrated that marine
85 phytoplankton can easily be Zn-limited in culture, and that Zn stress is exacerbated by low CO₂
86 due to an inability to synthesize the metalloenzyme carbonic anhydrase and resultant carbon co-
87 limitation (Morel et al. 1994; Buitenhuis et al. 2003; Sunda and Huntsman 2005). In this study,
88 we reconcile these perspectives with a comprehensive, multipronged study of the natural
89 phytoplankton assemblage in Terra Nova Bay (TNB), Antarctica, documenting evidence of Fe
90 and Zn stress in a low pCO₂ coastal environment.

91 **2 Results**

2.1 Biogeochemical characterization of Terra Nova Bay

Twenty-six stations within Terra Nova Bay (TNB) were, Antarctica, was temporally sampled temporally sampled over the course of one month (January 9 – February 18, 2018) during the 2017-2018 CICLOPS expedition (Fig. 1a; Supplementary Table 2) to concurrently characterize the natural progression of the phytoplankton bloom and biogeochemical changes in the water column (Kell et al. 2024). These stations were spatially distinct (each unique station was sampled once), but given that all stations were in relatively close proximity to each other within TNB (within a 52 km radius), we have combined all TNB station data to create a temporal analysis of the region.

Surface waters within TNB had low (~ 200 μatm) seawater pCO_2 (Fig. 1b) which contrasted with measurements >400 μatm further from the study site (Fig. 1c). A large phytoplankton bloom was present as indicated by high (> 3000 ng L^{-1}) chlorophyll fluorescence concentrations in January that waned into February (Fig. 1d). This observation of high productivity is characteristic of Antarctic polynya environments, which are recurring regions of open water surrounded by sea ice (Arrigo et al. 2012). This phytoplankton community initially consisted of a mixed assemblage of both diatoms as indicated by fucoxanthin (fuco, Fig. 1e) and the haptophyte *Phaeocystis* as verified by shipboard microscopy and as indicated by 19'-hexanoyloxyfucoxanthin (19'-hex, Fig. 1f). Surface fucoxanthin concentrations >200 ng L^{-1} were observed at the late TNB stations (Fig. 1e) while 19'-hex decreased to ~ 20 ng L^{-1} (Fig. 1f), indicating that the stations sampled in late February were dominated by diatoms rather than *Phaeocystis*. This was consistent with historical observations of phytoplankton succession patterns in TNB (DiTullio and Smith 1996; Smith et al. 2006; Mangoni et al. 2019). Additionally, we observed pronounced depletion of total dissolved Zn in surface waters across

all TNB stations, with an average concentration of 0.82 ± 0.47 nM at 10 m (Fig. 1g). Notably, as the bloom progressed, this depletion extended progressively deeper into the water column (Fig. 1g), indicative of strong Zn uptake and export from the euphotic zone.

Pronounced and progressively deepening total dissolved Zn (dZn_T) depletion over time was observed, with dZn depleted down to an average of 0.82 ± 0.47 nM at 10 m over all TNB stations (Fig. 1g).

The Zn-depleted area of the water column progressively deepened to 50 m and then to ~100 m throughout January and February (Fig. 1g), indicative of strong Zn uptake and export from the euphotic zone. Total Zn uptake (ρZn , measured concurrently using a stable isotope tracer method) (Kell et al. 2024) was highest in the shallow euphotic zone in early January and waned into February (Fig. 1h), following trends seen in chlorophyll fluorescence (Fig. 1d) and ^{19}N -hex (Fig. 1f). This ρZn trend was consistent with laboratory studies demonstrating the substantial Zn requirements of both diatoms and *Phaeocystis antarctica* (Saito and Goepfert 2008; Kellogg et al. 2020). Across all TNB stations, total dissolved Fe (dFe_T) in the upper 50 m remained below 1 nM (Fig. 1i) as observed previously in this region (Fitzwater et al. 2000). In the Ross Sea, dissolved iron (dFe) has previously been demonstrated to be the primary limiting nutrient for phytoplankton growth (Martin et al. 1990; Coale et al. 2003; Sedwick et al. 2011).

2.2 Biogeochemical characterization of the incubation study site

Within TNB station, station 27 (referred to as the “experimental site” herein) was chosen for the multifactor shipboard incubation experiment (Fig. 1a,b; red star). This site harbored a coastal bloom and was biologically and chemically characterized as having high *in situ* chlorophyll *a* levels (maximum of 3259 ng L^{-1} at 30 m; Fig. 1j) and was comprised of diatoms as

indicated by fucoxanthin and *Phaeocystis* as indicated by 19'-hex (**Fig. 1k**). A decrease in surface total dissolved inorganic carbon (DIC_T; 2181 $\mu\text{mol kg}^{-1}$ at 15 m compared to the deep water (200-1065 m) average of $2224 \pm 2.1 \mu\text{mol kg}^{-1}$, **Fig. 1l**) was also observed. Within the water column, dZn demonstrated a pronounced decrease from 5.1 nM at 50 m to 0.9 nM at 10 m, representing an 82% decrease (and a 76% decrease comparing the minimum dZn_T value at 10 m to the average deepwater (210 – 1000 m) concentration of $3.9 \text{ nM} \pm 0.4$; **Fig. 1m**), consistent with prior observations of surface dZn depletion in this region (Fitzwater et al. 2000).

Observations of rapid Zn uptake ($46 \text{ pmol L}^{-1} \text{ d}^{-1}$ at 10 m) at the experimental site (**Fig. 1n**) likely contributed to this surface depletion, as Zn uptake rates of this magnitude are of the appropriate scale to induce the multi-nanomolar surface water depletion during the austral spring and summer season (Kell et al. 2024). Consistent with high macronutrient abundance in this region, surface macronutrient concentrations (~~nitrate and nitrite, phosphate, and silicic acid~~) were partially depleted at the experimental site with 64%, 46%, and 29% decreases in ~~nitrate+nitrite (N+N), phosphate (P), and silicate (Si)~~nitrate+nitrite (N+N), phosphate (P), and silicate (Si), respectively, comparing 10 m and average deep water (200 – 1000 m) values (**Fig. 1o**).

2.3 Evidence for Zn stimulation of phytoplankton: experimental site shipboard incubations

A multifactor incubation experiment was conducted using surface waters collected at the experimental site by trace metal clean fish sampler (7 m) fed into a shipboard cleanroom to examine controls on net primary productivity, with triplicate treatments of Zn amended (+Zn; 2 nM as ZnCl_2), Fe amended (+Fe; 1 nM as FeCl_2), and Fe and Zn amended (+Fe+Zn) incubations, in addition to unamended controls. Addition of Fe alone (+Fe) resulted in significantly higher chl *a* content compared to controls ($p = 9.5\text{e-}5$) after six days (T6) (**Fig. 2a**), demonstrating primary

Fe limitation as observed previously in the Ross Sea (Martin et al. 1990; Mangoni et al. 2019). However, addition of Zn alone (+Zn) also resulted in significantly higher chl *a* content compared to the controls ($p = 0.011$), implying that a subset of the incubated phytoplankton population benefitted from the addition of Zn alone, without additional Fe (**Fig. 2a**). This observation is consistent with independent co-limitation, and may thus have been (Saito et al. 2008), where two nutrients (such as Fe and Zn) each independently limit different subpopulations or processes, and adding either nutrient alone yields a response experiencing primary Zn limitation (**Fig. 2a**). The combined addition of +Fe+Zn resulted in the highest average chl *a* content among all treatments at T6, with $4.5 \pm 0.21 \mu\text{g L}^{-1}$ compared to $3.9 \pm 0.35 \mu\text{g L}^{-1}$ and $2.8 \pm 0.15 \mu\text{g L}^{-1}$ achieved by +Fe and +Zn alone, respectively, demonstrating additive co-limitation between Zn and Fe (Sperfeld et al. 2016). The Zn stock solution was analyzed to confirm these results were not caused by inadvertent Fe contamination (see Methods). Significant differences in seawater chemistry were also observed within these incubations over time, with larger decreases in DIC_T in all metal treatments compared to the control ($-12.7 \mu\text{mol kg}^{-1}$ for +Fe ($p = 5.3\text{e-}6$), $-8.2 \mu\text{mol kg}^{-1}$ for +Zn ($p = 5\text{e-}5$), and $-18.5 \mu\text{mol kg}^{-1}$ for +Fe+Zn ($p = 2.2\text{e-}16$); **Fig. 2b**). The decrease in DIC_T observed with +Fe+Zn was significantly larger than that achieved with +Fe alone ($p = 4.4\text{e-}3$; **Fig. 2b**). Statistically significant differences in measured parameters among treatments are summarized in **Supplementary Table 3**.

Further consistent with the observed Zn stimulation of biomass in the incubations, the largest decreases in macronutrient (P and N+N) concentrations in these incubations at T6 were observed in the +Fe+Zn treatment (**Supplementary Figure 1a,b**), as was the largest increase in particulate organic carbon (POC; **Supplementary Figure 1c**). POC collected from the +Zn and +Fe+Zn incubations was characterized by larger C:N atomic ratios (5.9 and 6.2, respectively)

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compared to the +Fe and T6 control (5.2 and 5.3; **Supplementary Figure 1d**). Significantly higher bacterial abundances in both +Fe ($p = 9.1\text{e-}4$) and +Fe+Zn ($p = 6.3\text{e-}4$) treatments relative to the T6 control (**Supplementary Figure 1e**) indicated the alleviation of bacterial Fe limitation, consistent with prior reports (Obenosterer et al. 2015; Fourquez et al. 2020; Sun et al. 2021).

At the conclusion of the incubation experiments, biomass was collected [by serial filtration through 5 mm and 0.2 mm filters, and the 0.2-5 \$\mu\text{m}\$ fraction was extracted for proteomic analysis](#) ~~was performed~~ (see Methods) and analyzed for biomarkers of Zn and Fe stress. We detected both algal Fe- and Zn-stress proteins, which provided an independent line of evidence corroborating the results described above (**Fig. 2c**). This included the detection of the Zn/Co responsive protein ZCRP-A (a putative Zn chaperone) (Kellogg et al. 2022a) as a biomarker of Zn stress as well as the iron starvation-induced proteins ISIP1A, ISIP2A and ISIP3 (ISIPs) as biomarkers of Fe stress (**Supplementary Table 4**). The ISIPs represent a group of unrelated proteins that are upregulated under Fe limitation in various algal species. ISIP1 proteins are responsible for endocytosis of siderophore-bound iron, ISIP2 proteins are involved in Fe^{3+} uptake, and ISIP3 has been suggested to act as an Fe storage protein (Allen et al. 2008; Behnke and LaRoche 2020). RUBISCO abundance within each treatment is shown in **Fig. 2c** as a proxy for the potential phytoplankton production. Within the T6 incubation biomass, there was an increased abundance of ISIPs in the control and +Zn treatment, and a decrease in ISIP protein abundance within the +Fe and +Fe+Zn treatments, consistent with primary Fe limitation and the expected response to Fe addition (**Fig. 2c**). ISIPs were taxonomically assigned to diatoms, *Phaeocystis*, and dinoflagellates (**Fig. 2d**). The strongest expression of ZCRP-A protein was detected in the +Fe treatment (**Fig. 2c,d**) indicative of Fe addition driving the community towards increased Zn stress. Notably, ZCRP-A was still detected in the +Fe+Zn treatment (**Fig.**

207 **2c,d**), implying that the added Zn was unable to completely satiate Zn demand as phytoplankton
208 biomass increased (as indicated by the increase in chl *a* at T6, **Fig. 2a**), despite added Zn (2 nM)
209 being double that of added Fe (1 nM). Sequence analysis of the contigs identified as ZCRP-A
210 homologs in these incubations revealed that all contigs contained one or more canonical
211 conserved motifs found in COG0523 family proteins such as ZCRP-A (**Supplementary Figure**
212 **2**). Coupled with evidence from prior laboratory studies (Kellogg et al. 2022a), this provides
213 further support for the role of ZCRP-A in responding to Zn scarcity. ZCRP-A proteins were
214 taxonomically assigned to chlorophytes, dinoflagellates, and *Phaeocystis*, with the detection of
215 *Phaeocystis* ZCRP-A only in the +Fe treatment (**Fig. 2d**). The detection of ZCRP-A attributed to
216 *Phaeocystis*, but the nondetection of ZCRP-A attributed to diatoms, implies that either ample
217 diatom biomass was not captured on the analyzed filters due to being filtered out by the >5 μ m
218 pre-filter, or that diatoms present in these incubations (as indicated by diatom RUBISCO; **Fig.**
219 **2d**) were outcompeting *Phaeocystis* for Zn. Our observations of Fe and Zn biomarkers shifting in
220 abundance in response to their respective metal treatment provides independent evidence for
221 Zn/Fe co-limitation.

223 2.4 Taxonomic characterization of incubation results

224 To characterize the phytoplankton species responding to metal amendment, we measured
225 phytoplankton pigments within the shipboard incubations over time, which revealed a diverse
226 taxonomic response to metal amendments. Measured pigments included fucoxanthin (fuco), 19'-
227 hexanoyloxyfucoxanthin (19'-hex), prasinoxanthin (prasino), chlorophyll *b* (chl *b*), and
228 chlorophyll *c3* (chl *c3*). Fuco is produced by both diatoms and by *Phaeocystis* under certain
229 conditions, while 19'-hex and chl *c3* are indicative of *Phaeocystis* in the Southern Ocean

230 (DiTullio et al. 2007). Fuco:19'-hex ratios significantly increased in the +Fe ($p = 4.2\text{e-}4$) and
 231 +Fe+Zn treatments ($p = 2.7\text{e-}3$) (**Supplementary Figure 3a**) due to no significant change in
 232 fuco (**Supplementary Figure 3b**) and decreased 19'-hex (**Supplementary Figure 3c**) relative to
 233 the T6 control. *Phaeocystis* contributions to total fuco concentrations are typically minimal at the
 234 low Fe levels of the Ross Sea, though *Phaeocystis* can revert to making fuco rather than 19'-hex
 235 when released from Fe limitation (DiTullio et al. 2007), as was evident in these incubations by
 236 decreased 19'-hex:chl *c3* ratios within the +Fe and +Fe+Zn treatments (**Supplementary Figure**
 237 **3d**). *Phaeocystis* therefore likely contributed to total fuco by responding to Fe addition. Notably,
 238 significant decreases in both fuco:chl *a* and 19'-hex:chl *a* (**Supplementary Figure 3e,f**) in all
 239 treatments compared to the T6 control indicated that other phytoplankton groups contributed to
 240 chl *a* (**Fig. 2a**) without contributing fuco nor 19'hex. Increases in chl *b* (**Supplementary Figure**
 241 **3g**) and prasinoxanthin (**Supplementary Figure 3h**) suggest that small green algae such as
 242 chlorophytes and prasinophytes also responded to +Fe and +Zn independently, consistent with
 243 the detection of chlorophyte ZCRP-A in these incubations (**Fig. 2d**). Photosynthetic efficiency of
 244 photosystem II (Fv/Fm) significantly increased with +Fe ($p = 0.011$) and with +Fe+Zn ($p =$
 245 0.0036) at T4 (day 4) compared to T4 controls, but did not significantly increase with +Zn alone,
 246 implying Fv/Fm may not be useful as a diagnostic for Zn stress and that caution should be used
 247 in interpreting its signals universally (**Supplementary Figure 3i**). No significant difference in
 248 Fv/Fm was observed among treatments at T6. Selective zooplankton grazing on small diatoms
 249 and solitary *Phaeocystis* cells may have played a role in affecting phytoplankton biomass and the
 250 observed pigment:chl *a* ratios. For instance, higher ratios of phaeophytin:total phaeopigments
 251 were observed in +Fe and +Zn amended incubations (**Supplementary Figure 4**) which may
 252 reflect grazing on solitary *Phaeocystis* cells, as high phaeophytin:total phaeopigments ratios were

previously observed in *Phaeocystis* dominated waters of the Ross Sea (DiTullio and Smith 1996).

2.5 Detection of Zn- and Fe-stress protein biomarkers in the water column

Metaproteomic and metatranscriptomic analyses of biomass within the water column at the experimental site provided additional confirmation of the incubation results, as we detected Zn- and Fe- stress-response proteins present within the water column, which were therefore naturally present without influence from incubation conditions. In addition to ISIPs and ZCRP-A, we detected ZCRP-B (a putative membrane-tethered Zn-binding protein) (Kellogg et al. 2022a), Zrt/Irt-like (ZIP) ~~Zn~~ transporters (which are known to be used by marine phytoplankton for uptake of Zn^{2+} and other divalent metal cations (Allen et al. 2008; Milner et al. 2013; Bender et al. 2018)), and θ (theta) and δ (delta) carbonic anhydrases (CAs). θ -CAs with Zn^{2+} coordination sites have been documented in diatoms (Jensen et al. 2020a), including the polar diatom *Chaetoceros neogracile* RS19 (Kellogg et al. 2022b), but no studies to date have investigated enzyme activity nor efficiency with Co^{2+} or Cd^{2+} . In contrast, δ -CA (i.e., *Thalassiosira weissflogii* TWCA1) is known to function with either Co^{2+} or Zn^{2+} as a cofactor (Lane and Morel 2000b) conferring metabolic flexibility when Zn^{2+} is scarce.

Both proteins and transcripts of Zn and Fe stress biomarkers (ZCRP-A and ISIPs) were observed throughout the water column at the experimental site. RUBISCO, ZCRP-A, and ISIP protein spectral counts were most abundant at the surface and decreased with depth within the within the 3 μm size fraction (**Fig. 3a-c**), consistent with the depletion of trace metals in the photic zone due to high-biomass bloom conditions. ZCRP-A was detected in both 3 and 51 μm filter pore-size fractions (**Fig. 3b**) and was predominantly attributed to *Phaeocystis* and the diatom genus *Chaetoceros* in the euphotic zone, and predominantly to *Phaeocystis* and the

diatom genus *Pseudo-nitzschia* in the mesopelagic zone (**Fig. 3i, Supplementary Figure 5a**).

The presence of *Phaeocystis* below the photic zone is consistent with prior observations of rapid export of *Phaeocystis* cells (DiTullio et al. 2000). Throughout the water column, ISIPs were predominantly attributed to *Phaeocystis* and to the diatom genera *Fragilariopsis*, *Chaetoceros*, and *Pseudo-nitzschia* (**Fig. 3i, Supplementary Figure 5b**). We note that ZCRP-B, a protein also found to be upregulated in marine diatoms under low Zn/Co and characterized as a putative membrane-tethered Zn/Co protein ligand (Kellogg et al. 2022a) was most abundant in the 0.2 μm fraction throughout the water column (**Fig. 3d**). As ZCRP-B shares ~30% similarity to the bacterial ABC-type nickel transporter component NikA, spectral counts within the bacterial 0.2 μm fraction most likely reflect true bacterial NikA. BLAST analysis of all ZCRP-B contigs confirmed that all ZCRP-B hits across all size fractions corresponded to bacteria (**Fig. 3i**).

~~ZIPs were almost solely detected in the 51 μm fraction, likely due to the capture of abundant *Phaeocystis* colonies and chain-forming diatoms (**Fig. 3e; Supplementary Figure 5c**).~~

The assignment of the majority of ZCRP-A and ISIP proteins to *Phaeocystis* in the upper water column provides additional evidence that *Phaeocystis* was likely Zn/Fe co-limited at the study site, consistent with incubation results (**Fig. 2d**).

~~ZCRP-A belongs to the phylogenetically complex COG0523 family, with some family members showing functional divergence (that is, activity using different metal cofactors) among paralogs (Blaby-Haas and Merchant 2012; Edmonds et al. 2021). Here, we infer a Zn-responsive function for the identified ZCRP-A contigs based on their homology to *T. pseudonana* and *P. tricornutum* ZCRP-A proteins, which we have previously characterized as Zn-responsive (Kellogg et al. 2022a). To further support this inference, we used SHOOT (Emms and Kelly 2022) to place each ZCRP-A contig within a phylogenetic context. Of the 21 unique contigs~~

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299 assigned as ZCRP-A homologs, 19 were confirmed to be *T. pseudonana* orthologs, while 2 were
300 assigned as orthologs to the Zn-related COG0523 *E. coli* proteins YjiA and YeiR, implying a
301 minor prokaryotic source (**Supplementary Table 5**). The placement of the majority of these
302 contigs within diatom clades supports our interpretation that these homologs are Zn-related.
303 ZIP proteins were almost solely detected in the 51 µm fraction, likely due to the capture
304 of abundant *Phaeocystis* colonies and chain-forming diatoms (**Fig. 3e; Supplementary Figure**
305 **5c**). ZIP family transporters are functionally diverse and capable of transporting multiple
306 divalent metal cations, including both Zn²⁺ and Fe²⁺ (Blaby-Haas and Merchant 2012), with
307 diatom homologs of ZIP1 known to be upregulated under Fe stress (Lampe et al. 2018). Given
308 the co-limitation of Fe and Zn at the study site, it is difficult to determine which metal these ZIP
309 transporters were primarily mediating.

310 The increased abundance of diatom θ-CA and δ-CA proteins within the water column
311 (**Fig. 3f,g**), as well as transcripts for the diatom Cd carbonic anhydrase CDCA, which can
312 replace Zn²⁺ with Cd²⁺ as the catalytic cofactor (Lane and Morel 2000a), in the 3 µm and 51 µm
313 fractions at 200 m (**Fig. 3h**) was indicative of a sinking, prior diatom bloom event (Subhas et al.
314 2019). θ-CA and δ-CA were predominantly taxonomically assigned to the diatom genera
315 *Chaetoceros* and *Pseudo-nitzschia*, respectively, while CDCA transcripts belonged to the diatom
316 genera *Chaetoceros* and *Corethron* (**Fig. 3i**). The presence of θ-CA, but lack of δ-CA, assigned
317 to *Chaetoceros* is consistent with proteomic analysis of the polar diatom *Chaetoceros neogracile*
318 RS19 grown in culture under Zn limiting conditions (Kellogg et al. 2022b).

320 **2.6 Zn:P ratios of the surface seawater at the experimental site**

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A third independent line of evidence for the nutritional influence of Zn scarcity on TNB phytoplankton was obtained from *in situ* cellular stoichiometry. Particulate Zn:P ratios (Zn:P) analyzed from biomass collected at the surface of this experimental station were consistent with ratios from Zn-limited culture studies. Particulate Zn:C ratios reported previously in Zn-limiting culture studies of the diatom *Thalassiosira pseudonana* (Sunda and Huntsman 2005) were converted to Zn:P ratios using the Redfield ratio (Redfield 1958) (**Supplementary Table 65**). We then compared these ratios and associated growth rates with particulate Zn:P measured within biomass collected at 10, 25, 50 and 100 m at the experimental site. At each of these surface depths, Zn:P measured at the experimental site was $\sim 2\text{E-}4$ mol:mol, which, in comparison to cultured diatom Zn:P ratios, fell within the range of severely Zn-limited growth rates (**Supplementary Figure 6**), again demonstrating the propensity for Zn-limited growth in this region and corroborating the incubation results.

3 Discussion

Antarctic waters are generally considered to not be prone to Zn limitation, given that high ($> 1\text{nM}$) dZn concentrations are typically observed in surface Southern Ocean waters (Coale et al. 2003). -However, we observed multiple independent lines of evidence from both the field incubation experiment (chlorophyll, DIC, Zn and Fe biomarker proteins) and contextual environmental biogeochemical data of the water column at the incubation site (dZn, Zn uptake rates, pigments, cellular Zn:P stoichiometry, metaproteomic, and metatranscriptomic analyses) demonstrating that phytoplankton within Terra Nova Bay of the Ross Sea, Antarctica, were experiencing Zn and Fe nutritional stress.

343 Multiple factors could be considered as potential drivers in the creation of Zn-limiting
344 conditions in the field, including Zn demand imposed by total biomass and the species
345 comprising this biomass. The phytoplankton bloom observed during this expedition was
346 comprised primarily of diatoms and *Phaeocystis*, consistent with previous Ross Sea seasonal
347 blooms (Smith et al. 2006; Arrigo et al. 2012; Mangoni et al. 2019), and which contributed to the
348 observed high Zn uptake rates and thus surface Zn depletion (Kell et al. 2024), resulting in
349 nutrient-like dZn profiles throughout TNB.

350 Our field observation of Zn limitation was made in an environment characterized by
351 diminished pCO₂, which we consider as a factor potentially driving Zn stress. We observed a
352 substantial drawdown of surface seawater pCO₂ to 221 µatm at the incubation site (a ~45%
353 decrease compared to offshore waters in the Ross Sea measured during the same time frame; **Fig.**

354 **1b**). Biology was the driver of this decrease in pCO₂, rather than freshwater input from glacial
355 and sea ice melt. This is evident in the physicochemical data, where over the measured salinity
356 range (S=33.6-34.8), the effect of simple dilution by freshwater input (DIC=Total Alkalinity=0)
357 would result in a reduction of pCO₂ by only ~8-9 ppm. The signals we observe are much larger
358 than that, consistent with a large phytoplankton uptake driver. The total alkalinity (TA) also does
359 not change proportionally with DIC in this region, which is also not consistent with dilution
360 driving a conservative mixing of TA and DIC.

361 Laboratory studies have unequivocally demonstrated that marine phytoplankton can
362 easily be Zn-limited in culture due to their large Zn requirement, and that this effect is
363 exacerbated at low pCO₂ (Morel et al. 1994; Sunda and Huntsman 2005) due to the use of Zn as
364 a required catalytic cofactor within carbonic anhydrase (CA) metalloenzymes (Sunda and
365 Huntsman 2005). CAs catalyze the reversible dehydration of HCO₃⁻ to CO₂, the substrate

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required by the carbon fixing enzyme RUBISCO. As HCO_3^- constitutes about 90% of the dissolved inorganic carbon (DIC) pool in the surface ocean, sufficient CA activity prevents carbon stress in marine phytoplankton by ensuring adequate CO_2 supply to RUBISCO. It has therefore been hypothesized that the combination of high biomass and resulting low CO_2 may cause severe Zn depletion that may limit algal growth rates due to lack of Zn and thus reduced CA activity, and thus reduced availability of carbon for photosynthesis (Morel et al. 1994; Sunda and Huntsman 2005). This Zn-C limitation relationship is referred to as ‘biochemically dependent co-limitation’, in which the availability of one nutrient is essential for the acquisition or utilization of another nutrient, especially at low concentrations (Saito et al. 2008).

To explore this in the context of our field observations, using the available quantitative constraints on Zn and CO_2 co-limitation thresholds available from the literature (see Methods), we estimated that the threshold for Zn- CO_2 limitation in culture synthesized across many alga occurs at 259 $\mu\text{atm pCO}_2$. We then compared this laboratory-determined Zn/C limitation threshold estimate to both the *in situ* 221 $\mu\text{atm pCO}_2$ measured at our field study site, and to the historical, global trend in surface ocean pCO_2 (**Fig. 4a,b**). Global surface ocean pCO_2 levels are rapidly rising above both the laboratory-estimated 259 $\mu\text{atm pCO}_2$ Zn/C limitation threshold and our field observation value of 221 μatm (Jiang et al. 2023) (**Fig. 4a,b**). Though only a fraction of the modern-day surface ocean is currently at ≤ 250 ppm pCO_2 (predominantly comprised of polar regions; **Fig. 4c**), this represents a large decrease in oceanic extent compared to only 100 years ago (**Fig. 4d**). Even though this may move the majority of oceanic regions farther from Zn and C limitation thresholds, there continue to be highly productive and episodic coastal blooming events that induce significant pCO_2 drawdown (Harrison et al. 2018; Dai et al. 2022). These coastal regions are increasingly recognized as being disproportionately significant contributors to

global ocean carbon export (with respect to their area), particularly at the high latitudes (Harrison et al. 2018; Dai et al. 2022), and will hence continue to be prone to Zn stress at low CO₂ as we have observed. Many other coastal regions have been observed to experience depressed CO₂ such as the Amundsen Sea (Tortell et al. 2012), Amazon River plume (Valerio et al. 2021), the west Florida Shelf (Robbins et al. 2018), the East China Sea (Shim et al. 2007), the Northern Gotland Sea (Schneider and Müller 2018), and Monterey Bay, California (Chavez et al. 2018) to name a few examples. On the other hand, it is likely that despite rising pCO₂ levels, some coastal regions will continue to experience episodic or persistent low pCO₂ due to high productivity (as observed in this study), freshwater inputs, or other regional processes. Though we do not attempt to model future pCO₂ dynamics in these areas, our results suggest that Zn status may continue to be an important physiological constraint under low pCO₂ conditions, particularly in productive coastal systems. As such, Zn limitation should be considered as part of the broader framework for understanding carbon cycling in these regions, especially as they play a disproportionate role in global carbon export.

4 Conclusions

Given the great challenge of conducting Zn manipulation experiments without contamination, we did not try to manipulate pCO₂ as an additional experimental treatment. Instead, we actively sought out a low pCO₂ environment for the study site, building on prior laboratory studies and a cadmium (Cd) pCO₂ field study (Cullen et al. 1999). The interaction of Zn (and Cd and Co) with CO₂ is an important area of future research, particularly in coastal environments. With the continuing rise in atmospheric and surface ocean pCO₂ levels, broader changes in the biogeochemical cycling of Zn and other bioavailable trace metals will likely occur

within the oceans, influencing NPP and thus total ocean carbon storage capacity. These low pCO₂ conditions environments that routinely occur in numerous coastal environments globally should be further examined for Zn effects in addition to carbon uptake dynamics in different temperature environments (Tortell et al. 2008; Dai et al. 2022). While there are elaborate biochemical capabilities available to many marine algae for dealing with Zn scarcity (Kellogg et al. 2022a), our results suggest that the geographic extent of possible Zn/C co-limiting environments may further decrease in the coming decades with rising anthropogenic CO₂ emissions. Despite this, the biochemical demand for Zn in marine organisms remains substantial, with cellular demand rivaling that of Fe. The multitude of metabolic functions requiring Zn, including but not limited to carbonic anhydrase activity, implies the need for further exploration of Zn influences on primary productivity in a changing ocean environment.

5 Materials and Methods

5.1 Study area and sample collection

Sample collection occurred during the CICLOPS (Cobalamin and Iron Co-limitation of Phytoplankton Species) expedition (NBP18-01) aboard the RVIB *Nathaniel B. Palmer*, December 11, 2017 – March 3, 2018 in the Amundsen Sea and Ross Sea of the Southern Ocean (Fig. 1a). Station metadata is provided in Supplementary Table 2. All stations were assumed to be representative of TNB during this temporal study (as evident in the total dissolved metal, macronutrient, and chlorophyll *a* datasets). Water samples for dissolved trace metal analyses were collected using trace metal sampling protocols described previously (Cutter and Bruland 2012). A trace metal clean rosette suspended on a Kevlar line and equipped with twelve 8L X-Niskin bottles (Ocean Test Equipment) was used to collect seawater at depths ranging from 10 –

435 600 m. Niskin bottles were transported to a positive-pressure trace metal clean shipboard van for
436 filtration upon surfacing. Total fluorescence on the vertical profiles was measured using an ECO
437 chlorophyll fluorometer (Wet Labs) equipped to the rosette. The rosette also included
438 instrumentation for measuring conductivity and temperature (Sea-Bird Electronics).

439

440 **5.2 Preparation of plasticware**

441 Polyethylene and polycarbonate sampling and incubation bottles were rigorously cleaned
442 to remove trace metal contaminants before use. Bottles were rinsed with 18.2 Ω Milli-Q water
443 (Millipore), soaked for 72h in <1% Citranox detergent, rotated, soaked for an additional 72h, and
444 then rinsed five times with Milli-Q water. Bottles were then filled with 10% HCl (Baker instar-
445 analyzed) by volume and soaked for a minimum of one week, rotated, and soaked for another
446 week. Bottles were then rinsed five times with dilute acid (HCl, pH 2) and stored double-bagged
447 in plastic zip bags. All cleaning work was conducted in a Class 100 clean room.

448

449 **5.3 Underway seawater pCO₂ partial pressure**

450 Surface water pCO₂ measurements were conducted aboard the RVIB *Nathaniel B.*
451 *Palmer* using an underway method consisting of an air-water equilibrator and IR CO₂ analyzer
452 developed and operated by the Lamont-Doherty Earth Observatory (LDEO) group (Takahashi et
453 al. 2020). A complete data report and sensor list are
454 available: <https://service.rvdata.us/data/cruise/NBP1801/doc/NBP1801DATA.pdf> (last access:
455 14 December 2024) (<https://www.rvdata.us/>, last access: 14 December 2024).

456

457 **5.4 TDIC and POC measurements**

Total alkalinity (TA) and dissolved inorganic carbon (DIC) were measured on CTD and incubations samples in near real-time aboard the NBP. Dissolved inorganic carbon (DIC) and total alkalinity (TA) samples were collected following ~~previously established~~the protocols of ~~Dickson et. al. 2007~~(Dickson et al. 2007). DIC analyses were conducted within ~4 h of collection. We acidified 1.25 mL of sample using an automated custom-built injection and bubble stripping system coupled to an infrared gas analyzer (LICOR LI7000) and integrated the infrared absorption signal versus time for each stripped gas sample to yield a total mass of CO₂. Each sample was analyzed in triplicate or greater. Since microbubbles regularly formed as samples warmed between sample acquisition and DIC analysis, every integration curve was visually inspected and those curves that exhibited evidence for bubbles were rejected. Certified reference materials (Dickson CRM batch 169) were analyzed between every 3 to 4 unknowns. The estimated precision based upon unknowns (>860 samples run in triplicate) and CRM replicates (n = 738) was $\pm 2.0 \mu\text{mol kg}^{-1}$ (± 1 SD). ~~As described by DeJong et al. 2015~~(DeJong et al. 2015); ~~A~~analyses for TA on filtered samples were completed within ~12 h of collection by using a potentiometric titrator (Metrohm 855 Robotic Titrator) ~~(DeJong et al. 2015)~~. The estimated precision based on replicate analyses of CRMs (n = 195) was $\pm 2.6 \mu\text{mol kg}^{-1}$ (± 1 SD).

5.5 Analysis of historical atmospheric and surface ocean pCO₂ trends

Decadal surface ocean pCO₂ reconstructions ~~from Jiang et al. 2023~~ (Jiang et al. 2023) were downloaded, binned by decade, and plotted using the ‘violinplot’ in MATLAB. Atmospheric pCO₂ data was assembled from the running Mauna Loa record (Keeling et al. 1976), and from measurements made on Antarctic firn ice (Etheridge et al. 1996).

5.6 Calculation of Zn-and pCO₂ co-limitation of phytoplankton thresholds

There are few experimental measurements of Zn- and pCO₂-co-limitation, either in the lab or *in situ*. This study documented Zn of a natural phytoplankton assemblage in the field at a pCO₂ of ~220 ppm. In the literature, several models exist to interpret co-limitation (Buitenhuis et al. 2003; Saito et al. 2008). For this study we chose to use the biochemically dependent co-limitation model for growth rate (μ):

$$\mu = V_{max} \frac{[CO_{2(aq)}]}{\phi K_s + [CO_{2(aq)}]}, \quad (1)$$

where V_{max} is the maximum growth rate, $[CO_{2(aq)}]$ is the aqueous CO₂ concentration of the growth medium in micromoles per kilogram of seawater, K_s is the half-saturation constant in micromoles per kilogram of seawater, and ϕ is a Zn-dependent growth term:

$$\phi = \frac{[dZn] + K_{s,Zn}}{[dZn]} \quad (2)$$

Here, the dissolved Zn concentration in the growth medium $[dZn]$ is modified by a Zn-dependent saturation constant ($K_{s,Zn}$). Few studies have enough experimental data to robustly establish a kinetic relationship between $[dZn]$ and $[CO_{2(aq)}]$, so we compiled several estimates for these terms from the literature. Diatom growth rates under pCO₂ limitation were taken from Riebesell et al. (1993) (Riebesell et al. 1993). Reported pH and temperature measurements for each treatment, a total alkalinity of 2300 $\mu\text{mol kg}^{-1}$, and a salinity of 35 were used to calculate aqueous CO₂ concentrations using CO2SYSv3.2.1.(Sharp et al. 2023). Reported V_{max} and K_s for *D. brightwellii*, *T. punctigera*, and *R. alata* were 1.46, 1.30, and 0.93 d⁻¹, and 1.4, 1.2, and 2.1 $\mu\text{mol kg}^{-1}$, respectively. Values for coccolithophore growth ($K_s = 0.97 \mu\text{mol kg}^{-1}$, $V_{max} = 4.7 \text{ d}^{-1}$) were taken from Krumhardt et al. (2017) (Krumhardt et al. 2017). A value for $K_{s,Zn}$ of 300 pmol

L⁻¹ was taken from Buitenhuis et al. (2003). This value is for the coccolithophore *E. huxleyi* generated under varying CO₂ and Zn conditions, and Zn response growth curves under single CO₂ conditions (ambient) are similar to other diatoms like *T. pseudonana* (Sunda and Huntsman 1995). The value of 300 pmol L⁻¹ appears high, but is tied to the functional form of the biochemically co-limitation equation (Buitenhuis et al. 2003; Saito et al. 2008). Based on the same dataset and different models for co-limitation, Buitenhuis et al. (2003) arrived at K_{s,Zn} values of ranging from 38 pmol L⁻¹ to 300 pmol L⁻¹. They calculated Zn-limitation alone, at CO₂-replete conditions, of 19 pmol L⁻¹. Thus, the chosen value of K_{s,Zn} is not a reflection of high Zn demand but determined by the functionality of biochemical co-limitation by Zn and C.

To calculate ϕ , a surface ocean Zn concentration of 50 pmol L⁻¹ was assumed (Bruland 1980; Wyatt et al. 2014). While these concentrations reflect total dissolved Zn, the relationship between bioavailable free Zn and dZn, especially in the field, remains unclear. Eq. 1 was then used to calculate effective CO₂ concentrations (and thus pCO₂ values) at which growth is halved, or in other words, $\mu = 0.5V_{max}$. We note that this calculation is distinct from the CO₂ half-saturation constant because of the co-limitation by Zn. The median pCO₂ threshold for 50% growth from the three diatom species was 278 ppm. Including coccolithophores decreases the median to 259 ppm. These values are slightly higher than the *in situ* evidence for Zn limitation at 220 ppm presented in the present study. Our results cannot be considered as an upper bound for Zn-CO₂ limitation, but serve as evidence for growth limitation under those specific environmental conditions.

5.7 Analyses of total dissolved metals using isotope dilution

The analysis of total dissolved metals for this expedition has been described previously (Kell et al. 2024). Briefly, seawater collected shipboard by pressure-filtering X-Niskin bottles through an acid-washed 142 mm, 0.2 μm polyethersulfone Supor membrane filter (Pall) within 3 hours of rosette recovery using high purity (99.999%) N_2 gas and stored at 4°C. All sample collection occurred shipboard within an on-deck trace metal clean van. Samples were acidified to pH 1.7 with high purity HCl (Optima) within 7 months of collection and were stored acidified at room temperature for over 1 year prior to analysis. This extended acidification time was used to counteract any loss of metal due to adsorption to the bottle walls (Jensen et al. 2020b).

Quantification of total dissolved Fe, Mn, Ni, Cu, Zn, and Cd was performed using isotope dilution. Acidified seawater samples were spiked with a stable isotope spike solution artificially enriched in ^{57}Fe , ^{61}Ni , ^{65}Cu , ^{67}Zn , and ^{110}Cd (Oak Ridge National Laboratory). Concentrations and spike ratios were verified by ICP-MS using a multi-element standard curve (SPEX CertiPrep). Preconcentration of spiked seawater samples for total dissolved metal analysis was performed using the automated solid phase extraction system seaFAST-pico (Elemental Scientific) in offline concentration mode with an initial volume of 15mL and elution volume of 500 μL (Rapp et al. 2017; Wuttig et al. 2019). Following preconcentration, multielemental quantitative analysis was performed using an iCAP-Q inductively coupled plasma-mass spectrometer (ICP-MS) (Thermo Scientific). Concentrations of Mn, Fe, Ni, Cu, Zn and Cd were determined using a six-point external standard curve of a multi-element standard (SPEX CertiPrep), diluted to range from 1-10 ppb in 5% nitric acid. An indium standard (SPEX CertiPrep) was similarly added to these standard stocks, diluted to range 1-10 ppb. Instrument injection blanks consisted of 5% nitric acid in Milli-Q. Standard curve R^2 values were ≥ 0.98 for all metals monitored. Method accuracy and precision were assessed using the 2009

547 GEOTRACES coastal surface seawater (GSC) standard (n = 8; **Supplementary Table 76**),
548 which produced values consistent with consensus results (Kell et al. 2024).

550 **5.8 Macronutrient, pigment, and Fv/Fm analyses**

551 Seawater for macronutrient (silicate, phosphate, nitrate, and nitrite) analyses were filtered
552 through 0.2 µm pore-size Supor membrane filters and frozen at sea in acid-washed 60-mL high-
553 density polyethylene bottles until analysis. Macronutrient analyses were conducted by nutrient
554 autoanalyzer (Technicon Autoanalyzer II) by Joe Jennings at Oregon State University. The
555 chemotaxonomic distribution of phytoplankton pigments was determined using HPLC as
556 described previously (DiTullio et al. 2003). Photosynthetic efficiency of photosystem II (Fv/Fm)
557 was measured using a Phyto PAM phytoplankton analyzer (Walz, Effeltrich, Germany) as
558 described previously (Schanke et al. 2021).

560 **5.9 Bacterial abundance**

561 One ml samples for heterotrophic prokaryotes abundance (HPA) analysis were fixed for
562 10 min with a mix of paraformaldehyde and glutaraldehyde (1% and 0.05% final concentration,
563 respectively), frozen in liquid nitrogen and stored at -80°C until analysis. After thawing,
564 samples were stained with SYBR Green (Invitrogen Milan, Italy) using 10⁻³ dilution of stock
565 solution for 15 min at room temperature. Cell concentrations were assessed using a FACSVerse
566 flow cytometer (BD BioSciences Inc., Franklin Lakes, USA) equipped with a 488 nm Ar laser
567 and standard set of optical filters. FCS Express software was used for analyzing the data and HP
568 were discriminated from other particles on the basis of scatter and green fluorescence from
569 SYBR Green (Balestra et al. 2011).

570

571 **5.10 ICP-MS analysis and Zn uptake rates using ^{67}Zn**

572 ^{67}Zn stable isotope uptake experiments were performed to quantify the movement of
573 dissolved Zn to the particulate phase (in units of $\text{pmol L}^{-1} \text{d}^{-1}$); see (Kell et al. 2024). Briefly,
574 unfiltered seawater was collected using the trace metal rosette over a depth range of 10 – 600 m
575 into 250mL trace metal clean polycarbonate bottles. Bottles were spiked with ^{67}Zn such that the
576 total added (spiked) concentration of Zn was 2 nM. Immediately after spiking, incubation bottles
577 were sealed, inverted to mix, and transferred to flow-through on-deck incubators for 24hr.
578 Biomass was collected after 24hr by vacuum filtering each incubation sample at 34.5 kPa (5 psi)
579 onto an acid-cleaned $3\mu\text{m}$ pore-size acrylic copolymer Versapore filter (Pall) mounted on an
580 acid-cleaned plastic filtration rig. Sample filters were retrieved from storage at -80°C , removed
581 from cryovials using plastic acid-washed forceps, and transferred into trace metal clean 15 mL
582 PFA vials with 4 mL of 5% HNO_3 (Optima) containing a 1 ppb Indium (In) internal standard.
583 Filters were digested for $\sim 3.5\text{h}$ at 140°C using a HotBlock® heating block (Environmental
584 Express, USA) before they were removed and discarded. After evaporating the remaining
585 solution to just dryness, the residue was resuspended in 2 mL of 5% HNO_3 (Optima) by light
586 vortexing. Process blank filters were digested and processed as sample filters were. This
587 experiment was also carried out using ^{110}Cd as a tracer of Cd uptake in separate incubation
588 bottles (data not shown here). Digests were analyzed in duplicate by ICP-MS using a Thermo
589 ICAP-Q plasma mass spectrometer calibrated to a multi-element standard curve (Spex Certiprep)
590 over a range of 1 – 20 ppb. Natural Cd and Zn isotope abundances of the standards were
591 assumed to calculate concentrations of ^{110}Cd , ^{111}Cd , ^{114}Cd , ^{67}Zn , ^{66}Zn , and ^{68}Zn . Total Zn uptake
592 ($\text{pmol L}^{-1} \text{d}^{-1}$) was calculated using particulate ^{67}Zn and total water column dZn measurements as

described previously (Cox et al. 2014). The particulate metal measurements captured contributions from the active transport of metal into cells, nonspecific metal adsorption to cell surfaces, metal adsorption to non-living particulate organic matter, and metal adsorption to particulate inorganic matter, though we expect active transport into cells to dominate the measured particulate isotopic signal due to the high abundance of actively growing autotrophic cells in the photic zone observed in Southern Ocean during austral summer. Particulate Zn:P measurements were calculated using particulate Zn measured on Cd-spiked filters and thus do not include any pZn contribution from Zn tracer addition. Particulate phosphorus concentrations were measured by ICP-MS simultaneously and were calibrated to a standard curve ranging from 100 to 3200 ppb using a 1 ppm certified P stock (Alfa Aesar Specpure). All SPEX and P standard curves had R^2 values > 0.99 . The Zn stock solution used in the incubation experiments was similarly analyzed by ICP-MS to confirm that the stock was not Fe contaminated—this analysis showed that less than 2.3 pM (which was near the instrument blank level for this analysis) of iron was added for every 2 nM of zinc, far less than needed to stimulate phytoplankton to the extent observed in our experiments.

5.11 Shipboard incubation experiments

Incubation experiments were conducted at station 27 (-74.9870°N, 165.8898°E). Raw surface seawater was pumped directly into a cleanroom container van, collected into acid-cleaned 50L carboys, and dispensed into acid-washed 1L polycarbonate bottles using a trace metal sampling system with acid-washed polypropylene tubing and a teflon diaphragm pump. Incubation bottles were first rinsed with seawater then filled. Seawater was collected at 16:05 UTC. Triplicate incubation bottles were amended with +Fe (1 nM), +Zn (2 nM) and +Fe+Zn,

sealed, and placed into a flow-through on-deck incubator with light screens that shaded the incubator to 20% percent ambient surface irradiance. Incubations were sampled at 0, 48, 96, and 144 hours (corresponding to T0, T2, T4, and T6 timepoints) for analysis by filtering onto GFF filters for chlorophyll (all time points, biological triplicates), pigment analyses (T6, biological triplicates), and proteomic analyses (T6, pooled biological triplicates). Chlorophyll was extracted immediately, otherwise samples were frozen at -80°C until further analyses, with pigment and protein samples kept in -80°C freezers, liquid nitrogen dewars, or dry ice coolers at all times during transport back to the laboratories. All amendments and sampling were conducted in a positive-pressure, clean room van with laminar flow hoods and plastic sheeting to minimize trace-metal contamination.

5.12 Metaproteomic analysis

Water column metaproteomic biomass was collected onto 0.2, 3, and 51 µm pore-size filters (“field filters”) using in-situ battery operated McLane pumps. Half of each field filter was processed for metaproteomic analysis. Incubation metaproteomic biomass samples were serially filtered through a 5µm pore prefilter followed by a 142mm GFF filter. Three-fourths of each GFF filter was used for subsequent metaproteomic analysis of the incubations. All filters were frozen at -80°C and stored until laboratory extraction. To extract proteins, filters were placed into extraction buffer (1% SDS, 0.1M Tris/HCL pH 7.5, 10mM EDTA). 8 mL of buffer was used for each field filter, and 15 mL of buffer was used for each GFF incubation filter. All reagents were made with HPLC-grade water. Samples were heated at 95°C for 10 minutes and shaken at room temperature for 30 minutes. Filters were removed and protein extracts were filtered through 5.0 µm Millex low protein binding filters (Merck Millipore #SLSV025LS).

639 Millex filters were rinsed with 1 mL of extraction buffer to ensure no loss of protein. Samples
640 were then spun for 30 minutes at 3220 rcf in an Eppendorf 5810 centrifuge. The supernatant was
641 transferred to Vivaspin 5K MWCO ultrafiltration columns (Sartorius #VS0611). Protein extracts
642 were concentrated to approximately 300 µL, washed with 1 mL of extraction buffer, and
643 transferred to a 2 mL ethanol-washed microtube (all tubes from this point on are ethanol-
644 washed). Vivaspin columns were rinsed with small volumes of protein extraction buffer to
645 remove all concentrated protein and samples were brought up to 400 µL with extraction buffer.
646 Samples were incubated with 2 µL benzonase nuclease (EMD Millipore 70746-3) for 30 minutes
647 at 37°C.

648 Extracted proteins were purified from SDS detergent, reduced, alkylated and digested
649 with trypsin while embedded within a polyacrylamide tube gel, using a modified, previously
650 published method (Lu and Zhu 2005). A gel premix was made by combining 1 M Tris HCL (pH
651 7.5) and 40% Bis-acrylamide L 29:1 (Acros Organics) at a ratio of 1:3. The premix (103 µL) was
652 combined with 50-100 µg of the extracted protein sample, Tris-EDTA, 7 µL 1% APS and 3 µL
653 of TEMED (Acros Organics) to a final volume of 200 µL. After 1 hour of polymerization at
654 room temperature, 200 µL of gel fix solution (50% ETOH, 10% acetic acid in LC/MS grade
655 water) was added to the top of the gel and incubated at room temperature for 20 minutes. Liquid
656 was then removed and the tube gel was transferred into a new 1.5 mL microtube containing 1.2
657 mL of gel fix solution before incubating at room temperature, 350 rpm in a Thermomixer R
658 (Eppendorf) for 1 h. Gel fix solution was removed and replaced with 1.2 mL of destain solution
659 (50% MeOH, 10% acetic acid in LC/MS grade water) and incubated at 350 rpm, room
660 temperature for 2 h. Liquid was removed, gels were cut up into 1 mm cubes and added back to
661 tubes containing 1 mL of 50:50 acetonitrile:25 mM ammonium bicarbonate (ambic) and

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incubated for 1 h, 350 rpm at room temperature. Liquid was removed and replaced with fresh 50:50 acetonitrile:ambic solution and incubated at 16°C, 350 rpm overnight. The above step was repeated for 1 hour the following morning. Gel pieces were then dehydrated twice in 800 µl of acetonitrile for 10 min at room temperature and dried for 10 min in a ThermoSavant DNA110 speedvac after removing the solvent. Proteins were reduced in 600 µL of 10 mM DTT, 25 mM ambic at 56°C, 350 rpm for 1 h. The volume of unabsorbed DTT solution was measured prior to removal. Gel pieces were washed with 25 mM ambic, and 600 µL of 55 mM iodoacetamide was added to alkylate proteins at RT, 350 rpm for 1 h. Gel cubes were then washed with 1 mL ambic for 20 minutes, 350 rpm at RT. Acetonitrile (1mL) dehydrations and speedvac drying were repeated as described above. Trypsin (Promega #V5280) was added in an appropriate volume of 25 mM ambic to rehydrate and submerge gel pieces at a concentration of 1:20 µg trypsin:protein. Proteins were digested overnight at 350 rpm, 37°C. Unabsorbed solution was removed and transferred to a new tube. 50 µl of peptide extraction buffer (50% acetonitrile, 5% formic acid in water) was added to gels, incubated for 20 min at RT, then centrifuged at 14,100 x g for 2 min. The supernatant was collected and combined with the corresponding unabsorbed solution. The above peptide extraction step was repeated again, combining corresponding supernatants. Combined digested peptides were centrifuged at 14,100 x g for 20 minutes, supernatants transferred into a new tube and dehydrated down to approximately 20 µL in the speedvac. Total digested peptides were quantified (Bio-Rad DC protein assay, Hercules, CA) with BSA as a standard. Peptides were then diluted in 2% acetonitrile, 0.1% formic acid in LC/MS grade water to a concentration of 1µg/µL for storage until analysis. All water used in the tube gel digestion protocol was LC/MS grade, and all plastic microtubes were ethanol rinsed and dried prior to use.

684 Purified peptides were diluted to $0.1 \mu\text{g} \mu\text{l}^{-1}$ and $20 \mu\text{l}$ ($2 \mu\text{g}$) was injected onto a Dionex
685 UltiMate 3000 RSLCnano LC system (Thermo Fisher Scientific) with an additional RSLCnano
686 pump run in online two-dimensional active modulation mode coupled to a Thermo Fusion Orbitrap
687 mass spectrometer as described previously (McIlvin and Saito 2021).

688 ~~Frozen filters were thawed and extracted as described previously (Hughes et al. 2014).
689 Briefly, filters were extracted using SDS detergent to solubilize both membrane and soluble
690 proteins and subjected to a 10-min heating incubation at 95°C . Protein material was subsequently
691 purified, alkylated, reduced and digested with trypsin (mass spectrometry grade; Promega
692 Corporation) at a trypsin:protein ratio of 1:20 and peptides were purified using C18 tips (Pierce
693 C18 Tips, $100 \mu\text{l}$ bed; Thermo Fisher Scientific). Purified peptides were diluted to $0.1 \mu\text{g} \mu\text{l}^{-1}$ and
694 $20 \mu\text{l}$ ($2 \mu\text{g}$) was injected onto a Dionex UltiMate 3000 RSLCnano LC system (Thermo Fisher
695 Scientific) with an additional RSLCnano pump run in online two-dimensional active modulation
696 mode coupled to a Thermo Fusion Orbitrap mass spectrometer.~~

697 A translated metatranscriptome (see below) was used as a reference protein database and
698 peptide spectra matches were performed using the SEQUEST algorithm within Proteome
699 Discoverer v.2.1 (Thermo Fisher Scientific) with a fragment tolerance of 0.6 Da and parent
700 tolerance of 10 ppm . Identification criteria consisted of a peptide threshold of 95% (false
701 discovery rate (FDR) = 0.1%) and protein threshold of 99% (1 peptide minimum, FDR = 0.8%)
702 in Scaffold v.5 (Proteome Software) resulting in $5,387$ proteins identified in the incubation
703 experiment and $27,924$ proteins identified in the water column. To avoid double-counting mass
704 spectra, exclusive spectral counts were used for the downstream proteomic analysis. Exclusive
705 spectral counts were normalized using the normalized spectral abundance factor (NSAF)
706 calculation (Zhang et al. 2010) to allow for a comparison of protein abundance across samples

707 while remaining consistent with the metatranscriptomic procedure, see Cohen et al. 2021 for
708 details. Counts associated with redundant ORFs (sharing identical taxonomic and functional
709 assignments) were summed together. The stand-alone command line application BLAST+ from
710 the National Center for Biotechnology Information (NCBI) was used to identify proteins of
711 interest in the metaproteomic data. Metaproteomes were BLAST searched ($E = 5e-5$) against the
712 known sequences of proteins of interest acquired from annotated proteomic databases
713 (**Supplementary Table 4**) and combined with further annotation data based on contig ID (see
714 below).

715

716 **5.13 Metatranscriptomic analysis**

717 RNA sequencing was performed using the Illumina HiSeq platform. Transcriptomic
718 assemblies were generated for biomass collected using McLane pumps filtered through 0.2, 3,
719 and 51 μm pore-size filters. In order to enrich metatranscriptomic libraries derived from 0.2 μm
720 filters in prokaryotic transcripts and libraries derived from 3 μm and 51 μm filters in eukaryotic
721 transcripts, 0.2 μm libraries were generated from total rRNA-depleted mRNA and 3 μm and 51
722 μm libraries were generated from polyA mRNA. Total RNA was extracted from 0.2 μm , 3 μm ,
723 and 51 μm filters using Macherey-Nagel a NucleoMag RNA kit (Macherey-Nagel GmbH &
724 Co.KG). Cleared lysate was loaded into a 96 deep-well plate and put on an epMotion 5075 TMX
725 liquid handler to complete the RNA extraction following the Machery-Nagel standard protocol.
726 For 3 μm and 51 μm samples with total RNA greater than 1 μg , 800 ng of total RNA was used
727 for preparing poly A libraries with an Illumina Stranded mRNA Prep Ligation kit (Illumina),
728 following the manufacturer's protocol. For the 3 μm and 51 μm samples with total RNA less
729 than 1 μg , 20 ng of total RNA was used as input for the SMART-Seq v4 Ultra Low Input RNA

730 kit (Takara Bio USA, Inc), which converts poly(A) RNA to full-length cDNA using a modified
 731 oligo (dT) primer with simultaneous cDNA amplification. The resulting double-stranded cDNA
 732 was then fragmented using a Covaris E210 system with the target size of 300bp. Libraries were
 733 prepared from fragmented double-stranded cDNA using an Illumina Stranded mRNA Prep
 734 Ligation kit (Illumina). For RNA obtained from 0.2 μ m filters, ribosomal RNA was removed
 735 using a riboPOOL Seawater Kit (Galen Laboratory Supplies, North Haven, Connecticut, USA).
 736 The riboPOOL Seawater Kit is a customized mixture of Removal Solutions: Pan-Prokaryote
 737 riboPOOL, Pan-Plant riboPOOL and Pan-Mammal in a ratio of 6:1:1. The rRNA-depleted total
 738 RNA was used for cDNA synthesis by Ovation RNA-Seq System V2 (TECAN, Redwood City,
 739 USA). Double stranded cDNA was then prepared for the libraries using an Illumina Stranded
 740 mRNA Prep Ligation kit (Illumina). Ampure XP beads (Beckman Coulter) were used for final
 741 library purification. Library quality was analyzed on a 2200 TapeStation System with an Agilent
 742 High Sensitivity DNA 1000 ScreenTape System (Agilent Technologies, Santa Clara, CA, USA).
 743 Resulting libraries were subjected to paired-end Illumina sequencing via NovaSeq S4.

744 The input paired-end fastq sequences are trimmed of sequencing adapters, primers and
 745 low quality bases by using either BLASTN (NCBI, v2.2.25) (Altschul et al. 1990) or
 746 trimmomatic, v0.36 (Bolger et al. 2014). The trimmed paired and unpaired sequences were then
 747 depleted of rRNA sequences with riboPicker v0.4.3 (Schmieder et al. 2012). The command-line
 748 program clc_assembler, v5.2.1 (Qiagen) was used to assemble processed sequences into contigs
 749 and ORFs were identified by FragGeneScan, v1.31 (Rho et al. 2010). The trimmed sequences
 750 were mapped to the predicted ORFs using the command-line program clc_mapper, v5.2.1
 751 (Qiagen) to generate mapped raw read counts for each ORF. The raw counts were normalized
 752 initially to RPKM values, to account for variations in inter-sample sequencing depth and the

ORF sequence length (Mortazavi et al. 2008). The RPKM values were subsequently converted to TPM (transcripts per million) units for estimation of the relative RNA abundance among samples (Li and Dewey 2011). The ORFs were annotated for putative function by several programs in parallel using BLASTP against PhyloDB, hidden Markov models PFAM and TIGRFAM by HMMER, v3.3.2 (Eddy 2011), KEGG Ortholog HMM by kofamscan, v1.3.0 (Aramaki et al. 2020), and transmembrane HMM by TMHMM (Krogh et al. 2001). Additional annotations were generated by similarity searches using BLASTP to transporter (PhyloDB), organelle (PhyloDB) and KOG (Tatusov et al. 2003) databases. The ORFs are assigned to the best taxonomic species/group as determined by LPI (Lineage Probability Index) analysis (Podell and Gaasterland 2007). The final list of curated ORFs was generated by removing ORFs with low mapping coverage (< 50 reads total over all samples) and with no BLAST hits and no known domains.

5.14 Statistical analysis and data visualization

ANOVA and Dunnett tests were performed using MATLAB 2019a. Statistics are summarized in **Supplementary Table 3**. Figures were made using matplotlib (version 3.5.0), Ocean Data View (version 5.5.2), Excel (2019), and RStudio (version 1.3.1093). Color palettes used in Ocean Data View section plots (<https://doi.org/10.5281/zenodo.1243862>) are inverse “roma” for trace metal concentrations, “thermal” for Zn and Cd uptake rates, and “algae” for chlorophyll fluorescence (Crameri 2023).

Data Availability

775 CICLOPS (NBP18-01) conductivity–temperature–depth (CTD) hydrography data
 776 including pressure, temperature, total dissolved oxygen, conductivity, fluorescence, and beam
 777 transmission (<https://doi.org/10.1575/1912/bco-dmo.783911.1>) and total dissolved metal, Zn and
 778 Cd uptake rate, macronutrient, and pigment datasets are available through the NSF Biological
 779 and Chemical Oceanography Data Management Office (BCO-DMO) repository
 780 (<https://doi.org/10.7284/907753>). Underway pCO₂ data collected during cruise NBP1801 are
 781 available through R2R at <https://doi.org/10.7284/139318>. The mass spectrometry global
 782 proteomics data for CICLOPS bottle incubations and water column analyses have been deposited
 783 with the ProteomeXchange Consortium through the PRIDE repository under the project name
 784 “Zinc-iron co-limitation of natural marine phytoplankton assemblages in coastal Antarctica” with
 785 project accession number PXD037056
 786 (<https://www.ebi.ac.uk/pride/archive/projects/PXD037056>). This data is accessible for review by
 787 using the following login information: username reviewer_pxd037056@ebi.ac.uk, password:
 788 IFdOUoEb. The translated transcriptome used for spectrum to peptide matching has been
 789 deposited in the National Center for Biotechnology Information sequence read archive under
 790 BioProject accession no. PRJNA890306
 791 (<https://www.ncbi.nlm.nih.gov/bioproject/?term=PRJNA890306>) and RNA-Seq BioSample
 792 accession nos. SAMN31286421-SAMN31286522
 793 (<https://www.ncbi.nlm.nih.gov/biosample/?term=SAMN31286421>).
 794

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804 **Author contributions statement**

805 All authors contributed to data acquisition and analysis. RMK, NLS, LEL, RJC, DR, DMM,
806 MRM, FB, RBD, GRD and MAS implemented the shipboard incubation study. RMK, MMB,
807 RJC, DR, TJH, and AVS contributed to the formal analysis. NLS, LEL, FB, OM, RF, and GRD
808 contributed to pigment datasets and interpretations. CB provided bacterial abundance data. RBD
809 contributed DIC and *in situ* pCO₂ data. AVS contributed analyses and discussion regarding
810 historical pCO₂ data and Zn-C growth limitation estimates. AEA contributed the
811 metatranscriptome reference database used for proteomic analyses. RMK and MAS wrote the
812 original draft. RMK, MAS, GRD, AVS, TJH, and RBD contributed to review and editing. All
813 authors approved the final submitted manuscript.

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815 **Competing interests statement**

816 The authors declare no competing interests.

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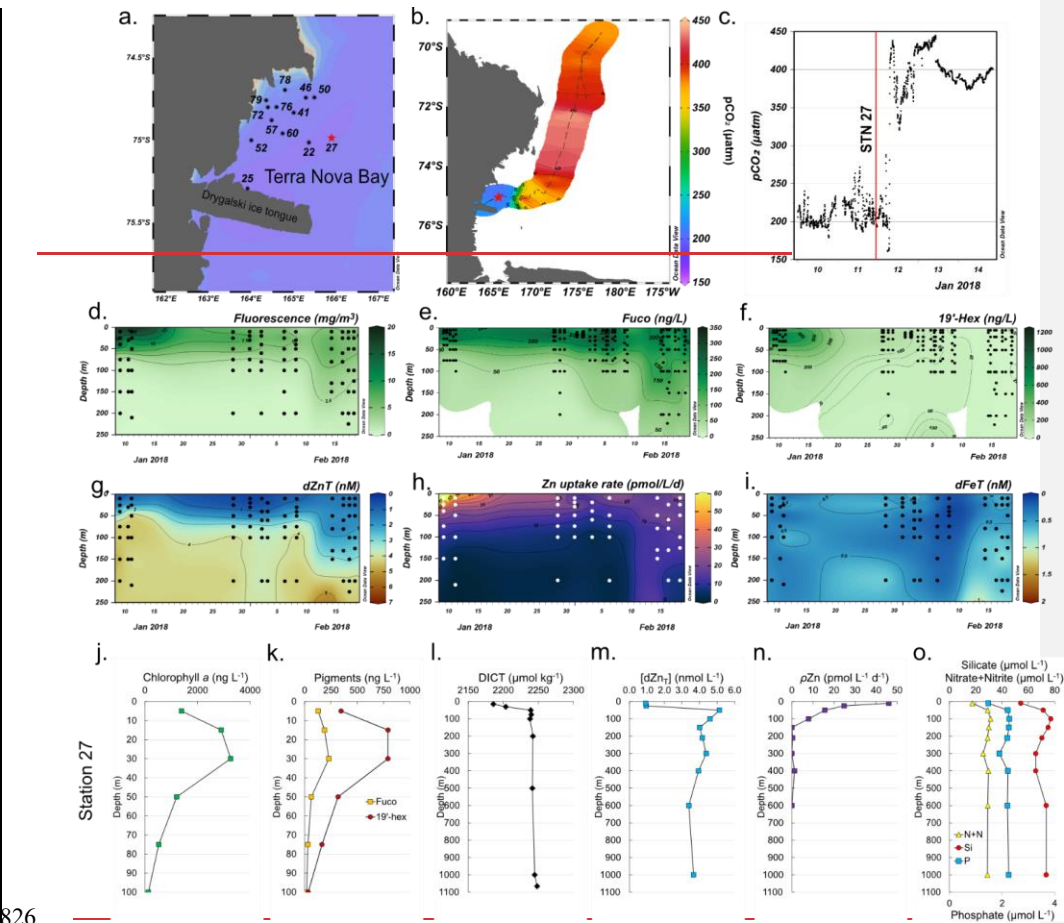
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Figures



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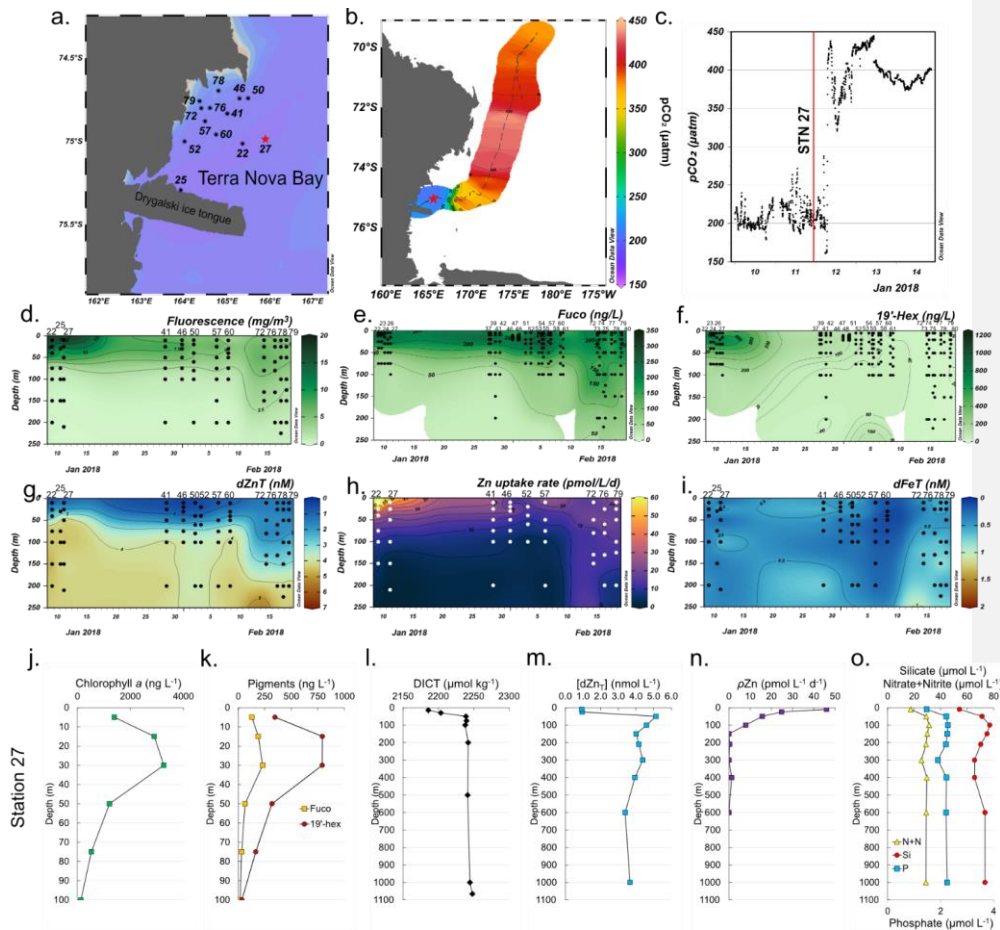
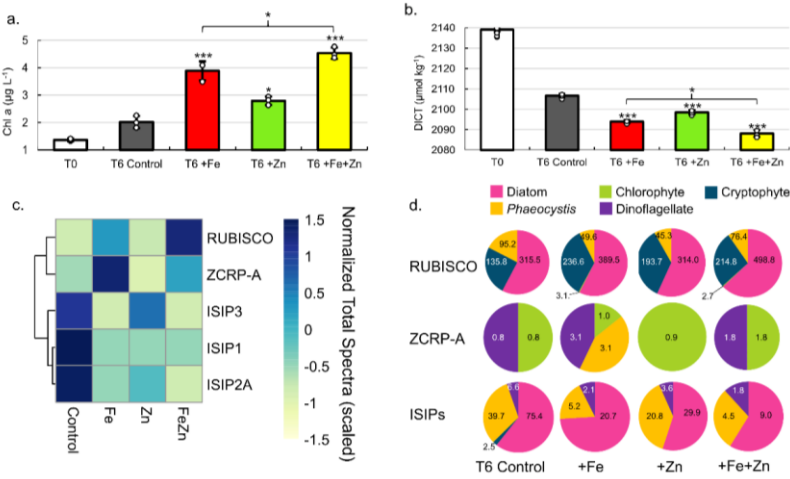


Figure 1. Temporal biogeochemistry of Terra Nova Bay and characterization of the experimental site at Station 27. (a) Sampling locations over the Ross Sea shelf in Terra Nova Bay, Antarctica. (b) Location of station 27 (red star) and surrounding seawater pCO₂ measured over a three-day transit northwards represented in color scale. (c) pCO₂ measured over time within TNB during the three-day transit shown in (b). The vertical red line denotes the pCO₂

834 level at the time of initial seawater collection at station 27. **(d)** Total chlorophyll fluorescence,
835 **(e)** fucoxanthin (fuco), **(f)** 19' hexanoyloxyfucoxanthin (19'-hex), **(g)** total dissolved Zn, **(h)** total
836 Zn uptake rates, and **(i)** total dissolved Fe measured in the upper 250 m represented on a color
837 scale. Station data is presented in order of sampling date, from the earliest (Stn 22, early January)
838 to the latest (Stn 79, late February). The data gap between January 13-23 occurred when the ship
839 was unable to sample due to icebreaking duties for the McMurdo Station resupply ship. Stations
840 indicated in (a) are those where the trace metal rosette (TMR) was deployed; pigment data was
841 supplemented with additional TNB stations using a CTD (Table S2). Depth profiles of **(j)**
842 chlorophyll *a*, **(k)** the pigments fuco and 19'hex, **(l)** total dissolved inorganic carbon (DICT), **(m)**
843 total dissolved Zn, **(n)** total Zn uptake rates, and **(o)** the macronutrients nitrate+nitrite (N+N),
844 phosphate (P), and silicate (Si) at the study site. Panels **(d),(g),(h)** and **(i)** were originally
845 presented in Kell et al. (2024) and are reprised here to introduce the environmental context of the
846 study site.

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871 **Figure 2. Evidence for Zn co-limitation with Fe in bottle incubations.** (a) Chlorophyll *a* and
872 (b) total DIC (DICT) at T0 (day 0) and in each treatment at T6 (day 6). Significant differences
873 among groups were found using one-way ANOVA and post-hoc Dunnett test (** $p < 0.001$, **
874 $p < 0.01$, * $p < 0.05$). Error bars are the standard deviation of biological triplicates ($n=3$).
875 Individual data points are overlaid (white circles). (c) Heatmap of row-scaled exclusive protein
876 spectral counts (normalized total spectra) showing relative protein abundance in each treatment.
877 The dendrogram shows similarity in spectral abundance among samples based on Euclidean
878 distance and hierarchical clustering. Color gradients represent low (yellow) to high (blue) protein
879 expression. Ribulose-1,5-biphosphate carboxylase/oxygenase (RUBISCO), zinc/cobalt
880 responsive protein A (ZCRP-A), and iron starvation induced proteins (ISIP1, ISIP2A, ISIP3) are
881 shown. (d) Taxonomies assigned to RUBISCO, ZCRP-A, and ISIP proteins in each treatment at
882 T6. Counts (normalized total spectra) assigned to each taxa are shown. ISIPs are the combined
883 spectral counts of ISIP1A, ISIP2A and ISIP3.

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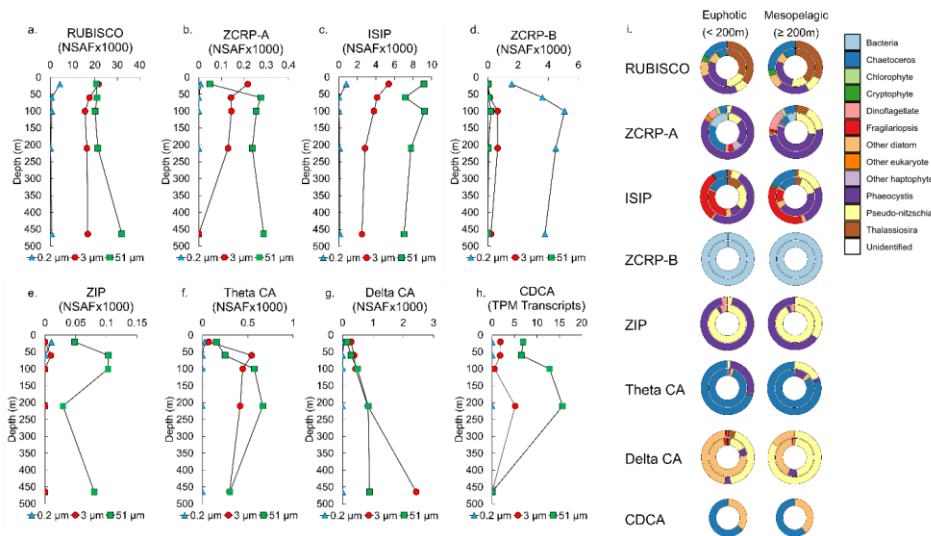


Figure 3. Metatranscriptomic and metaproteomic detection of Zn- and Fe-related proteins of interest at the experimental site. Depth profiles of summed NSAF-normalized protein spectral counts of (a) RUBISCO, (b) ZCRP-A, (c) iron starvation induced proteins (ISIPs), (d) ZCRP-B, (e) ZIP, (f) Theta CA, and (g) Delta CA detected from proteomic analysis of each filter size fraction (0.2, 3 and 51 μm). (h) TPM-normalized transcript read counts of CDCA. (i) Stacked pie charts depicting relative community composition for proteins of interest for euphotic (< 200 m) and mesopelagic (≥ 200 m) depths. The outer rings show community composition based on NSAF-normalized protein spectral counts while the inner rings are TPM-normalized transcript read counts. Protein and transcript counts plotted in (i) were summed across all size fractions. ISIPs are the combined spectral counts of ISIP1A, ISIP2A and ISIP3.

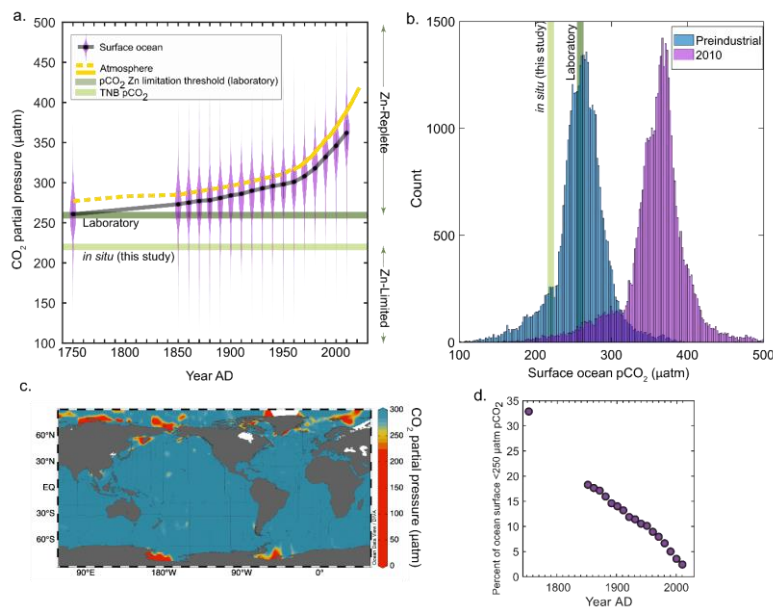


Figure 4. The partial pressure of CO₂ (pCO₂) and associated phytoplankton responses from this study and the literature. (a) Pre-industrial and decadal surface ocean pCO₂ reconstructions plotted as violin plots, with a running black line through the median values. The atmospheric curve is a composite of ice core data (dashed yellow line (Etheridge et al. 1996)) and the Mauna Loa record (solid yellow line (Keeling et al. 1976)). An estimated thresholds for zinc-limited

916 growth is plotted as the median of previous laboratory results (259 μatm , dark green line; see
917 Methods), and is compared to the *in situ* results of this study (220 μatm , light green line). **(b)**
918 Data in (a) plotted as a histogram comparing preindustrial and modern (2010) pCO_2 values, with
919 the same pCO_2 levels indicated. **(c)** Global map of surface ocean pCO_2 plotted using
920 GLODAPv2.2022 data (Lauvset et al. 2022). **(b)** Percentage of the ocean surface less than 250
921 μatm pCO_2 as a function of time. Surface ocean pCO_2 reconstructions taken from Jiang et al.
922 2023.

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 1239
 1240 **Supplementary information for “Zinc stimulation of phytoplankton in a low carbon**
 1241 **dioxide, coastal Antarctic environment: evidence for the Zn hypothesis”**
 1242
 1243 **Supplementary Table 1. Summary of zinc incubation experiments in marine environments.**
 1244 Note that all experiments focus on phytoplankton populations with the exception of Fukada et al
 1245 (2000) and Mazzotta et al. (2021) which focus on bacterial heterotrophy. This list represents our
 1246 knowledge of the literature but may be incomplete.
 1247

Study	Location	Zn Treatment Replication	pCO ₂ Data	Dissolved Environmental Zn Data	Zn Response
Scharek et al. 1997(Scharek et al. 1997)	Southern Ocean	Zn n=2, control n=1	No	No	Negative/“very small”
Fukada et al., 2000(Fukuda et al. 2000)	North Pacific	Integrated water column leucine aminopeptidase activity (0-100m)	No	Yes	Leucine aminopeptidase activity positively correlated with dZn
Cochlan et al. 2002(Cochlan et al. 2002)	Western Ross Sea	None (n=1 +Zn, +Zn+Fe)	No	No	Increase in specific uptake of NO ₃ ⁻
Coale et al. 2003(Coale et al. 2003)	Ross Sea, Southern Ocean	None (n=1 +Zn, +Zn+Fe)	No	Yes	Negative/“very small”
Crawford et al. 2003(Crawford et al. 2003a)	Northeast Pacific Ocean	Triplicate (n=3)	Yes	No	“Slightly but significantly” altered Chl, nitrate and phosphate

Franck et al. 2003(Franck et al. 2003)	Costa Rica Dome	Triplicate (n=3)	No	No	Secondary Zn limitation after Fe in diatom microscopy data
Ellwood 2004(Ellwood 2004)	Subantarctic	Duplicates (n=2)	No	Yes	Negative
Cullen et al. 1999; Cullen and Sherrell 2005(Cullen et al. 1999; Cullen and Sherrell 2005)	Coastal California	Duplicates (n=2)	No	No	Enhanced Cd:P in low pCO ₂ treatments, decrease in Cd:P with Zn addition
Jakuba et al. 2012(Jakuba et al. 2012)	North Pacific	Singlicate with timepoints sacrificed	No	Yes	Primary, no additive effect with iron.
Dreux Chappell et al. 2016(Dreux Chappell et al. 2016)	Costa Rica Dome	Triplicate (n=3)	No	Yes	Secondary limitation with Si

Sharma et al. 2020(Sharma et al. 2020)	Eastern Arabian Sea	Triplicate (n=3)	Yes	No	Negative/ “insignificant effects”
Mazzotta et al. 2021(Mazzotta et al. 2021)	Equatorial Pacific	Quintuplicate (n=5) sediment trap incubations	No	Yes in separate study (Cohen et al., 2021)	Enhancement of alkaline phosphatase activity with Zn addition in sediment trap samples
This study	Terra Nova Bay, Ross Sea	Triplicate (n=3)	Yes	Yes	Primary and Secondary limitation. Independent validation by Zn biomarkers.

Supplementary Table 2. Station metadata for the NBP18-01 cruise. Stations at which total Zn uptake rates were determined are indicated by asterisks (*).

Station	Latitude (°N)	Longitude (°E)	Sampling Date
			(yyyy-mm-dd hh:mm)
4*	-72.751	-116.001	2017-12-30 01:23

10	-73.054	-129.988	2018-01-03 03:39
11*	-74.047	-133.764	2018-01-03 19:58
15*	-75.864	-151.918	2018-01-05 14:51
20*	-76.714	179.819	2018-01-08 02:00
22*	-75.013	165.358	2018-01-09 15:56
<u>23</u>	<u>-75.003</u>	<u>165.781</u>	<u>2018-01-01 3:51</u>
<u>24</u>	<u>-75.031</u>	<u>165.551</u>	<u>2018-01-10 16:33</u>
25	-75.293	163.914	2018-01-11 01:26
<u>26</u>	<u>-75.139</u>	<u>164.836</u>	<u>2018-01-11 5:14</u>
27*	-74.987	165.890	2018-01-11 16:05
29*	-76.001	172.997	2018-01-16 03:00
31	-77.295	175.390	2018-01-17 04:39
32*	-76.750	172.000	2018-01-17 19:19
34	-77.147	168.503	2018-01-23 22:59
35*	-76.231	168.769	2018-01-26 20:17
<u>37</u>	<u>-75.001</u>	<u>164.999</u>	<u>2018-01-27 23:21</u>
<u>39</u>	<u>-75.001</u>	<u>164.170</u>	<u>2018-01-28 6:00</u>
41*	-74.833	165.002	2018-01-29 00:41
<u>42</u>	<u>-75.166</u>	<u>164.169</u>	<u>2018-01-29 19:30</u>
46*	-74.742	165.287	2018-01-31 21:32
<u>47</u>	<u>-74.877</u>	<u>164.249</u>	<u>2018-02-01 19:28</u>
<u>48</u>	<u>-74.873</u>	<u>164.670</u>	<u>2018-02-01 23:30</u>
50	-74.741	165.488	2018-02-02 21:17

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<u>51</u>	<u>-75.000</u>	<u>164.498</u>	<u>2018-02-03 0:46</u>
52*	-75.000	164.005	2018-02-03 21:43
<u>53</u>	<u>-74.980</u>	<u>164.441</u>	<u>2018-02-05 5:30</u>
<u>54</u>	<u>-74.750</u>	<u>164.169</u>	<u>2018-02-05 19:26</u>
<u>55</u>	<u>-74.751</u>	<u>165.498</u>	<u>2018-02-06 0:10</u>
57*	-74.879	164.482	2018-02-06 20:12
<u>58</u>	<u>-74.893</u>	<u>164.843</u>	<u>2018-02-07 5:26</u>
60	-74.959	164.739	2018-02-08 20:06
<u>61</u>	<u>-74.807</u>	<u>164.773</u>	<u>2018-02-09 6:35</u>
62*	-74.999	169.491	2018-02-09 19:27
67*	-76.454	167.919	2018-02-11 19:19
70	-74.744	170.374	2018-02-13 21:25
72*	-74.800	164.395	2018-02-14 22:28
<u>73</u>	<u>-75.037</u>	<u>164.503</u>	<u>2018-02-15 7:03</u>
<u>74</u>	<u>-74.839</u>	<u>164.531</u>	<u>2018-02-15 23:18</u>
<u>75</u>	<u>-74.790</u>	<u>164.677</u>	<u>2018-02-16 6:22</u>
76*	-74.799	164.597	2018-02-16 20:15
<u>77</u>	<u>-74.863</u>	<u>165.764</u>	<u>2018-02-17 6:59</u>
78	-74.696	164.796	2018-02-17 20:34
79*	-74.757	164.356	2018-02-18 19:29
<u>80</u>	<u>-74.637</u>	<u>164.792</u>	<u>2018-02-18 23:00</u>

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Supplementary Table 3. Summary of ANOVA statistics for T6 incubation treatments comparing measured parameters. Significant differences among groups were found using one-way ANOVA and post-hoc Dunnett test (** $p < 0.001$, * $p < 0.01$, $p < 0.05$, . $p < 0.1$). Chl *a*, chlorophyll *a*; DIC_T, total dissolved inorganic carbon; chl *b*, chlorophyll *b*; Prasino, prasinoxanthin; fuco, fucoxanthin; 19'hex, 19'-hexanoyloxyfucoxanthin; chl *c*3, chlorophyll *c*3. NA, no statistically significant difference. All parameters were measured in biological triplicate.

Treatment (T6)	Chl <i>a</i>	DIC _T	Chl <i>b</i>	Prasino	Fuco	19'hex	Fuco:hex	Hex:Chl <i>c</i> 3	Bacterial abundance
+Fe vs Ctrl	$p = 9.5\text{e-}5$ (***)	$p = 5.3\text{e-}6$ (***)	$p = 4.0\text{e-}3$ (**)	$p = 2.9\text{e-}2$ (*)	NA	$p = 8\text{e-}3$ (**)	$p = 4.2\text{e-}4$ (***)	$p = 2.0\text{e-}4$ (***)	$p = 9.1\text{e-}4$ (***)
+Zn vs Ctrl	$p = 1.1\text{e-}2$	$p = 5.0\text{e-}6$	$p = 8.0\text{e-}2$	$p = 7.4\text{e-}3$	NA	NA	NA	$p = 0.02520$	NA

	(*)	(***)	(.)	(**)				(*)	
+FeZn vs Ctrl	$p =$ 1.3e-7 (***)	$p =$ 2.2e-16 (***)	$p =$ 4.0e-4 (***)	$p =$ 5.7e-2 (.)	NA	NA	$p =$ 2.7e-3 (**)	$p =$ 1.9e-4 (***)	$p =$ 6.3e-4 (***)
+FeZn vs +Fe	$p =$ 3.4e-2 (*)	$p =$ 4.4e-3 (**)	NA	NA	NA	$p =$ 6.0e-2 (.)	NA	NA	NA

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1270 **Supplementary Table 4.** Representative proteins of interest, reference organism and IDs.

Protein of interest	Reference organism	Protein ID
ZCRP-A	<i>Thalassiosira pseudonana</i> CCMP1335	3054 (JGI Thaps3*)
ZCRP-B	<i>Thalassiosira pseudonana</i> CCMP1335	938 (JGI Thaps3_bd**)
RUBISCO	<i>Phaeodactylum tricornutum</i> CCMP632	AAF07200.1 (NCBI)
ISIP1A	<i>Thalassiosira oceanica</i> CCMP1005	K0RCT3 (Uniprot)
ISIP2A	<i>Phaeodactylum tricornutum</i> CCMP632	B7FYL2 (Uniprot)
ISIP3	<i>Phaeodactylum tricornutum</i> CCMP632	B7G4H8 (Uniprot)
<u>ZIP1</u>	<i>Phaeodactylum tricornutum</i> CCMP632	46780 (JGI Phatr2 [†])
CDCA	<i>Thalassiosira pseudonana</i> CCMP1335	25840 (JGI Thaps3*)

1271 *Joint Genome Institute (JGI) Thaps3 database

1272 (<https://mycocosm.jgi.doe.gov/Thaps3/Thaps3.home.html>)

1273 **Joint Genome Institute Thaps3_bd database

1274 (https://mycocosm.jgi.doe.gov/Thaps3_bd/Thaps3_bd.home.html)

1275 [†]Joint Genome Institute CCAP 1055/1 v2.0 Phatr2, all models database
1276 (<https://mycocosm.jgi.doe.gov/Phatr2/Phatr2.home.html>)

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1279 **Supplementary Table 5.** Summary of metaproteomic contigs detected in both the incubation
1280 biomass and in the water column at station 27 identified as ZCRP-A or ZIP homologs. For each
1281 contig, the top reference BLAST hit (see Supplementary Table 4) is listed with alignment
1282 statistics (E-value, percent identity). An example ortholog identified by SHOOT and the
1283 ortholog’s corresponding UniProt protein description is also presented.

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1286 **Supplementary Table 65.** Particulate Zn:P ratios measured in *T. pseudonana* cultured at various
1287 concentrations of Zn (log [Zn']) (Sunda and Huntsman 2005) compared to particulate Zn:P ratios
1288 measured at Station 27 of research cruise NBP18-01. Zn:C measurements by Sunda and
1289 Huntsman 2005 were converted to Zn:P using the Redfield ratio. Data is plotted in
1290 Supplementary Figure 6. Zn' refers to the sum of inorganic Zn metal complexes.

1291 [†]Particulate P data measured directly in this study

1292 *Particulate P from Sunda 2005 data was estimated by converting particulate C measurements to
1293 P using the Redfield ratio (106C:1P).

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Particulate Zn:P ratios at Station 27

Depth (m)			Particulate Zn (mol)		Particulate P (mol) ⁺	Zn:P (mol:mol)	
100			1E-11		8E-08	1E-04	
50			1E-11		1E-07	1E-04	
25			3E-11		2E-07	2E-04	
10			4E-11		2E-07	2E-04	
Cellular Zn:P ratios of cultured <i>T. pseudonana</i> from Sunda and Huntsman 2005, Table 1.							
Experiment	Log [Zn ²⁺]	pH	Particulate Zn (mol)	Particulate C (mol)	Particulate P (mol)*	Zn:P (mol:mol)	Growth rate (d ⁻¹)
1	-12.05	8.2	3.3E-05	22	0.21	2E-04	0.1
1	-11.47	8.2	5.2E-05	15	0.14	4E-04	1.23
1	-10.87	8.2	1.2E-04	15	0.14	9E-04	1.62
1	-10.36	8.2	1.5E-04	15	0.14	1E-03	1.76
1	-9.82	8.2	1.8E-04	15	0.14	1E-03	1.76
1	-11.96	9	3.8E-05	15	0.14	3E-04	0.51
1	-11.38	9	8.5E-05	15	0.14	6E-04	1.12
1	-10.78	9	2.0E-04	15	0.14	1E-03	1.28
1	-10.27	9	4.1E-04	15	0.14	3E-03	1.42
1	-9.73	9	5.0E-04	15	0.14	4E-03	1.49
2	-11.82	8.2	4.9E-05	15	0.14	3E-04	0.64

2	-11.82	8.2	4.0E-05	15	0.14	3E-04	0.79
2	-10.87	8.2	1.3E-04	15	0.14	9E-04	1.45
2	-9.82	8.2	2.0E-04	15	0.14	1E-03	1.51
2	-11.71	9	5.8E-05	15	0.14	4E-04	0.73
2	-11.71	9	5.8E-05	15	0.14	4E-04	0.78
2	-10.76	9	1.9E-04	15	0.14	1E-03	1.39
2	-10.76	9	2.1E-04	15	0.14	1E-03	1.38
2	-9.71	9	4.3E-04	15	0.14	3E-03	1.44
2	-9.71	9	5.0E-04	15	0.14	4E-03	1.42
4	-11.82	8.2	2.8E-05	18.1	0.21	3E-04	0.72
4	-11.32	8.2	5.5E-05	15.2	0.13	4E-04	1.45
4	-10.82	8.2	1.3E-04	14.7	0.14	9E-04	2
4	-10.32	8.2	2.2E-04	15.1	0.14	2E-03	2.03
4	-9.82	8.2	2.6E-04	15.6	0.14	2E-03	2.04

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Supplementary Table 76. Reference seawater comparisons using the 2009 GEOTRACES coastal surface seawater (GSC) standard.

This study	GEOTRACES GSC consensus (nM)
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Metal	(n = 8)	
	(nM)	
Fe	1.6 ± 0.23	1.6 ± 0.12
Zn	1.4 ± 0.23	1.5 ± 0.10
Cd	0.4 ± 0.01	0.4 ± 0.02
Cu	1.3 ± 0.05	1.1 ± 0.15
Ni	4.2 ± 0.07	4.5 ± 0.21
Mn	2.1 ± 0.37	2.2 ± 0.08

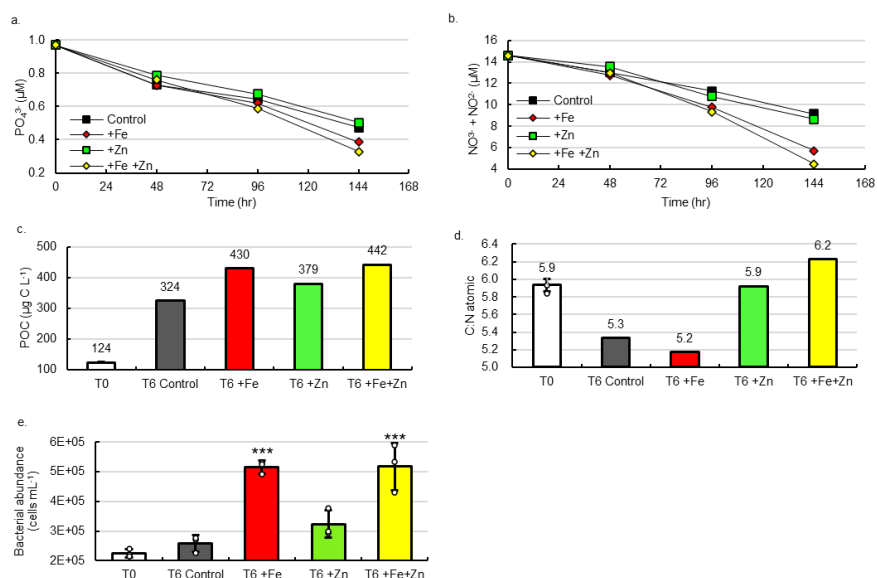
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1305 **Supplementary Figures**

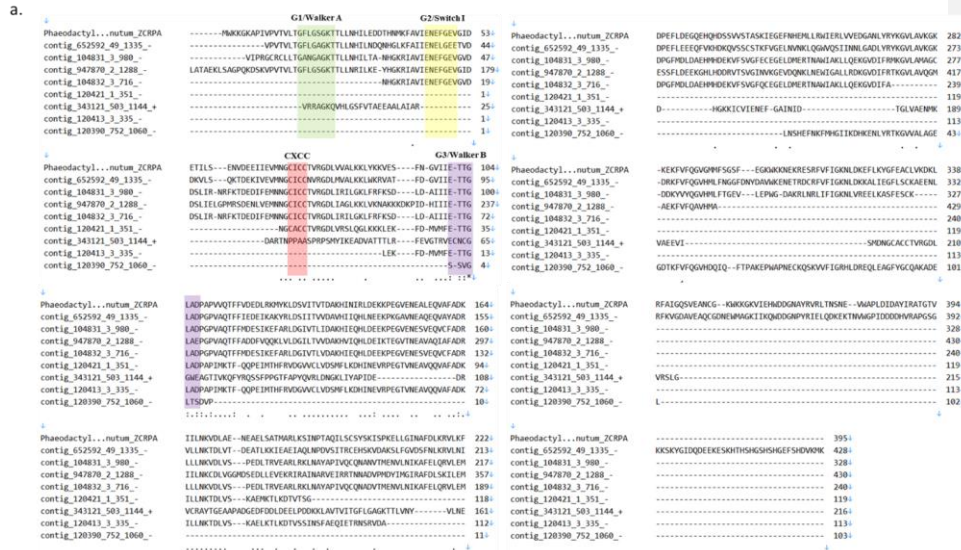
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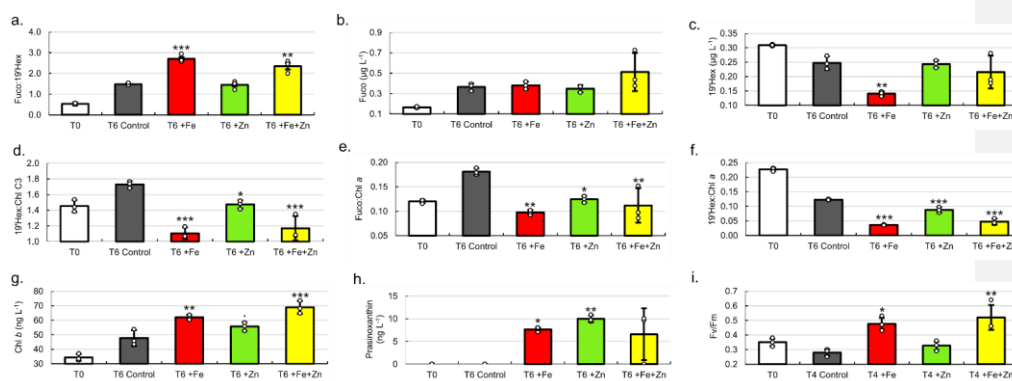
Supplementary Figure 1. Additional parameters measured in shipboard bottle incubations.

Time course decreases in (a) phosphate and (b) nitrite + nitrate. (c) total POC of biomass in each treatment at T6. (d) the atomic carbon:nitrogen (C:N) ratio of biomass in each treatment at T6, and (e) bacterial abundance. Significant differences among groups were found using one-way ANOVA and post-hoc Dunnett test (***) $p < 0.001$, ** $p < 0.01$, * $p < 0.05$). Error bars are the standard deviation of biological triplicates (n=3) with individual data points overlaid (white circles). Macronutrients, POC, and PON were measured in singlicate (n=1) from pooled biological triplicates.



Supplementary Figure 2. Sequence alignments of ZCRP-A peptides detected in T6 incubation biomass. (a) Sequence alignment of the *Phaeodactylum tricornutum* ZCRP-A protein compared to all ZCRP-A proteins detected in T6 incubation biomass. Alignment was generated using the MUSCLE algorithm with default parameters within MEGA11. Four conserved GTPase (G1/Walker A, G2/Switch I, CXCC metal binding, and G3/Walker B) are labeled. **(b)** E values and percent% identities of the identified proteins with significant sequence similarity to *P. tricornutum* ZCRP-A aligned above.

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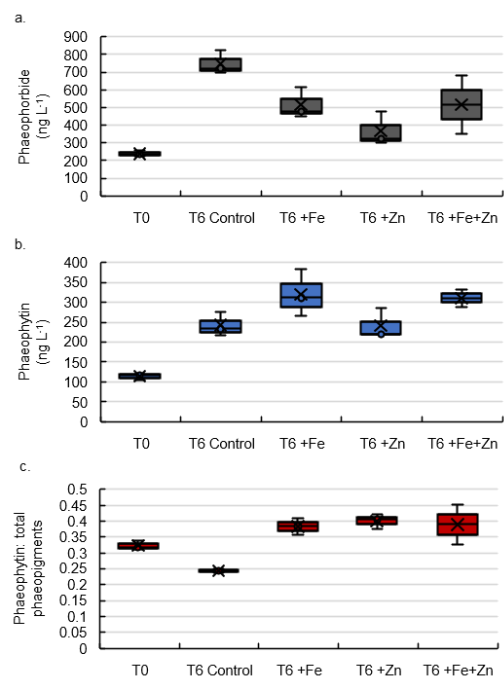
1338 **Supplementary Figure 3. Pigment analysis of T6 incubations.** (a) ratio of fucoxanthin: 19'-
1339 Hex, (b) fucoxanthin, (c) 19-hexanoyloxyfucoxanthin (19'-Hex), (d) ratio of 19'-Hex: chl c3, (e)
1340 ratio of fucoxanthin: chlorophyll a, (f) ratio of 19'-Hex: chlorophyll a, (g) chlorophyll b, (h)
1341 prasinoxanthin, and (i) maximum quantum efficiency (Fv/Fm) among treatments at T4.

1342 Significant differences among groups were found using one-way ANOVA and post-hoc Dunnett
1343 test (*** p < 0.001, ** p < 0.01, * p < 0.05, . p < 0.1). Data with error bars are presented as mean
1344 values ± the standard deviation of biological triplicates (n=3) with individual data points overlaid
1345 (white circles).

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1350 **Supplementary Figure 4.** Phaeopigments measured in T6 incubation biomass. Abundances of
1351 (a) phaeophorbide and (b) phaeophytin, and (c) the ratio of phaeophytin: total phaeopigments
1352 (the sum of phaeophorbide and phaeophytin). Error bars are the standard deviation of biological
1353 triplicates (n=3).

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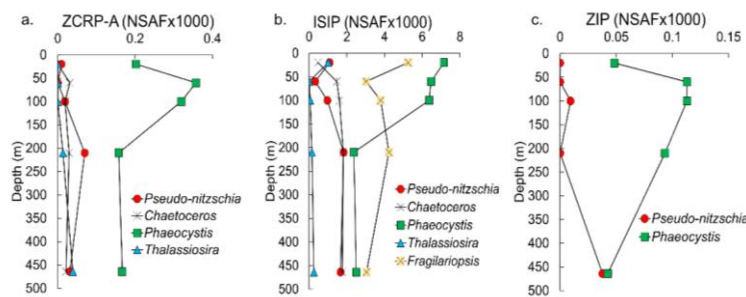
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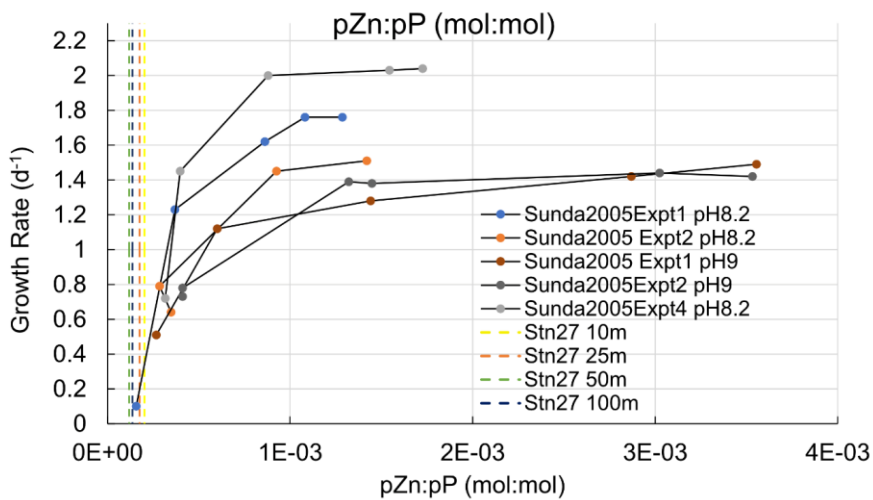
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Supplementary Figure 5. Depth profiles for proteins of interest at station 27 categorized by genus. Depth profiles of NSAF-normalized protein spectral counts of (a) ZCRP-A, (b), iron starvation induced proteins (ISIPs), and (c) ZIPs summed by genus. Proteins assigned to an individual genus were summed across all size fractions (0.2, 3 and 51 μ m). ISIPs are the combined spectral counts of ISIP1A, ISIP2A and ISIP3.



Supplementary Figure 6. Comparison of particulate Zn: particulate P ratios measured in the water column at the experimental site to those ratios measured in culture studies of a Zn-limited diatom. Ratio of particulate Zn (pZn) and particulate phosphorus (pP) (pZn:pP) measured in the upper water column at the study site (station 27; vertical dashed lines) compared to pZn:pP measurements in Zn limitation studies of the diatom *Thalassiosira pseudonana* in culture by Sunda and Huntsman 2005.

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