Part 2: Quantitative contributions of cyanobacterial alkaline phosphatases to biogeochemical rates in the subtropical North Atlantic

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Abstract

Microbial enzymes alter marine biogeochemical cycles by catalyzing chemical transformations that bring elements into and out of particulate organic pools. These processes are often studied through enzyme rate-based estimates and nutrient-amendment bioassays, but these approaches are limited in their ability to resolve species-level contributions to enzymatic rates. Molecular methods including proteomics have the potential to link the contributions of specific populations to the overall community enzymatic rate; this is important because organisms will have distinct enzyme characteristics, feedbacks, and responses to perturbations. Integrating molecular methods with rate measurements can be achieved quantitatively through absolute quantitative proteomics. Here, we use the subtropical North Atlantic as a model system to probe how a combination of traditional bioassays and absolute quantitative proteomics can provide a more comprehensive understanding of nutrient limitation in marine environments. The experimental system is characterized by phosphorus stress and potential metal-phosphorus co-limitation due to dependence of the organic phosphorus scavenging enzyme alkaline phosphatase on metal cofactors. We performed nutrient amendment incubation experiments to investigate how alkaline phosphatase absolute abundance and activity is affected by trace metal additions and develop an inventory of cyanobacterial alkaline phosphatases. We show that

the two most abundant picocyanobacteria, *Prochlorocccus* and *Synechococcus* are minor contributors to total alkaline phosphatase activity as assessed by a widely used enzyme assay, with *Prochlorococcus* accounting for 3-35% and *Synechococcus* contributing 0.5-5% of alkaline phosphatase activity depending on location and metal cofactor. This was true even when trace metals were added, despite both species having the genetic potential to utilize both the Fe and Zn containing enzymes, PhoX and PhoA respectively. Serendipitously, we also found that the alkaline phosphatases responded to cobalt additions suggesting possible substitution of the metal center by Co in natural populations of *Prochloroccocus* (substitution for Fe in PhoX) and *Synechococcus* (substitution for Zn in PhoA). This integrated approach allows for a nuanced interpretation of how nutrient limitation affects marine biogeochemical cycles and highlights the benefit of building quantitative connections between rate and "-omics" based measurements.

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Introduction

Microbial enzymes alter marine biogeochemical cycles by catalysing chemical transformations and facilitating the movement of elements through planetary reservoirs. On one hand, enzyme contributions from different groups of microbes can be considered collectively, for instance in rate-based or bioassay incubation experiments where the activities of the entire microbial community are aggregated. On the other hand, we anticipate that the enzymes of different organisms will have different activities and responses to perturbations; this means that resolving enzyme provenance could enhance the quantitative connection between microbial activity and biogeochemical rates (e.g. the goals of the fledgling Biogeoscapes program (Saito et al., 2024)). "-Omics" based methods, particularly proteomics which directly resolves protein/enzyme concentrations, can provide a window into the relationships between microbial abundance, enzyme concentration, and biogeochemical rates.

In this work we use quantitative proteomics to constrain the relative contributions of different microbes (*Synechococcus* and *Prochlorococcus*) to biogeochemical rates of alkaline phosphatase activity in the oligotrophic subtropical North Atlantic gyre. In this region, primary production is constrained by availability of dissolved inorganic nitrogen (DIN) and phosphorus (DIP), but inputs of atmospherically derived iron (Fe) from Saharan desert dust create a niche for nitrogen fixation, partially alleviating nitrogen limitation but driving the system to DIP depletion (Martiny et al., 2019; Moore et al., 2013). Lack of DIP then drives a shift towards the acquisition of the abundant yet less bioavailable dissolved organic phosphorus (DOP) by phytoplankton (Lomas et al., 2010; Mather et al., 2008). The DOP pool includes relatively labile phosphomono- and diesters (together ~75 to 85 % of DOP) that derive from ribonucleic acids, adenosine phosphates and phospholipids (Kolowith et al., 2001; Young and Ingall, 2010). These compounds cannot be directly assimilated but require the phosphate group to be cleaved from the ester moiety first. Cleaving is catalysed by a range of hydrolytic enzymes, such as alkaline phosphatases, which are common in marine microbes, including bacterial as well as eukaryotic phytoplankton (Dyhrman and Ruttenberg, 2006; Luo et al., 2009; Shaked et al., 2006). Reflecting this, alkaline phosphatase activity (APA) is high across the oligotrophic gyres(Browning et al., 2017; Davis et al., 2019; Duhamel et al., 2010; Mahaffey et al., 2014; Wurl et al., 2013).

Alkaline phosphatase activity is commonly regulated by intracellular phosphate levels (Santos-Beneit, 2015) and appears to be closely linked to low ambient DIP concentrations (Mahaffey et al., 2014). However, these enzymes also have a metal dependence, as metal co-factors are involved in the hydrolysis process at the active site. Different alkaline phosphatases exist that, while sharing function, evolved independently and have distinct metal requirements. For example, in *Escherichia coli* (*E. coli*) the alkaline phosphatase PhoA has two Zn^{2+} (zinc) or Co^{2+} (cobalt) ions and one Mg^{2+} (magnesium) ion at each active site per homodimer(Coleman, 1992), and in *Pseudomonas fluorescens*

the monomeric alkaline phosphatase PhoX has two Fe^{3+} ions and three Ca^{3+} (calcium) ions(Yong et al., 2014). The active sites of PhoA and PhoX in marine microbes have yet to be characterized but based on sequence homology are presumed to be like these model organisms, leading to the hypothesis that alkaline phosphatase activity to be limited by scarce Fe, Zn, or Co trace metals in the marine environment (Lohan and Tagliabue, 2018).

Global change is predicted to intensify phosphorus stress and alter trace metal and nutrient cycles in the ocean (Hoffmann et al., 2012; Kim et al., 2014). Throughout the North Atlantic, the utilisation of DOP is widespread (Mather et al., 2008) and whole community rates of APA are high compared with other oceanic regions (Duhamel et al., 2010; Mahaffey et al., 2014). At this time, it is not known which microbes and enzyme types are responsible for bulk APA in the North Atlantic and elsewhere. Resolving this could lead to a more quantitative understanding of how APA activity is regulated in the modern ocean, allowing better predictions of future changes in enzyme abundance and activity and the resulting influence on carbon export. In this study, we use field-based quantitative proteomics to develop an inventory of alkaline phosphatase activity and to identify nutrient-related regulatory controls on alkaline phosphatase that are distinct for different organisms. We use this as a proof of concept for developing quantitative connections between biogeochemical rates and "-omics" based measurements of microbial enzymes, a topic that is of interest to ongoing international efforts to characterize ocean metabolism.

Methods

Shipboard bioassays

All samples for this study were collected on board the *RRS James Cook* during research cruise JC150 (GEOTRACES process study GApr08), on a zonal transect at 22 °N leaving Guadeloupe on June 26th and arriving in Tenerife on August 12th, 2017, with multiple stations occupied for bioassays. A detailed description of the bioassays and analysis of environmental parameters is presented in Mahaffey et al. (submitted as a companion to this article).

Briefly, surface seawater was collected and processed according to trace metal clean protocols and before dawn. For each location, duplicate or triplicate 24 L polycarbonate (Nalgene) carboys were filled and spiked with additions of Fe, Zn or Co, as detailed in Table 1. The seawater was incubated at ambient sea surface temperature and 50 % surface light level for 48 h from dawn to dawn with a 12:12 h simulated light cycle using white daylight LED panels.

Table 1 Bioassay details at each station, showing the types of treatments, the amount of metal added, and the number of replicates per treatment for which proteomics analyses were conducted. Note that one of the three replicates of the Fe addition at the Station at 31 °W (*) was removed as an outlier from all further analysis.

	Station at 54 °W	Station at 50 °W	Station at 45 °W	Station at 31 °W

Treatment					
	Control	-	-	-	-
	Fe	+ 1.0 nM	+ 1.0 nM	+ 1.0 nM	+ 1.0 nM
	Zn	+ 1.0 nM	+ 1.0 nM	+ 0.5 nM	+ 1.0 nM
	Co	+ 50 pM	+ 50 pM	+ 50 pM	+ 20 pM
Replicates per treatment		2	2	2	3*

After the incubation period, subsamples for proteins were collected into acid cleaned 10 L polycarbonate carboys (Nalgene) and immediately filtered, collecting the >0.22 µm fraction on polyethersulfone membrane filter cartridges (Millipore, Sterivex) and recording the filtered volume. Any remaining water was pressed out with an air-filled syringe, the filtration unit was sealed with clay and then frozen at -80 °C. This procedure was repeated for the second (and third where applicable) replicate of each treatment.

Alkaline phosphatase activity (APA) rate measurements

Total APA was measured in unfiltered seawater samples using the synthetic fluorogenic substrate 4- methylumbeliferyll-phosphate (MUFP, Sigma Aldrich, Ammerman 1993, Davis et al 2019). MUFP stock solutions (100 mM in 2-methoxyethanol) were diluted with Milli-Q deionized water (200 µM stock). Unfiltered seawater was spiked with the MUFP substrate to final concentrations of 500 nM or 2000 nM MUFP for single substrate additions, or a series of replicates were incubated over a final MUFP concentration range from 100 nM to 2000 nM for the determination of enzyme kinetic parameters, Vmax and Km. Once spiked, samples were incubated in polycarbonate bottles in triplicate in the temperature and light adjusted reefer container for up to 12 hours.

MUFP hydrolysis to the fluorescent product, 4- methylumbelliferone (MUF), was measured at regular intervals (typically every 90 minutes) over a period of up to 8 - 12 hours using a Turner 10Au field fluorometer (365 nm excitation, 455 nm emission) after the addition of a buffer solution (3 : 1 sample: 50 mM sodium tetraborate solution, pH 10.5). A calibration was produced at the start and end of the cruise using MUF standards (concentration range 0–1000 nM) to ensure linearity of the fluorescence of MUF over the expected concentration range. Fluorescence response factors were determined daily using freshly prepared 200 nM MUF stocks and was used to convert the rate of change in fluorescence to MUFP hydrolysis rate, here considered to be synonymous with volumetric APA (nM P h-1). Boiled seawater blanks (500 nM MUFP) were incubated in parallel with samples to ensure that there was no significant change in fluorescence due to abiotic degradation or hydrolysis over time. Enzyme kinetic parameters were determined using a range of MUFP concentrations. Michaelis-Menten equation was transformed to produce substrate-response curves or linear regression plots and the maximum hydrolysis rates (Vmax) and half saturation constant (Km)

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were determined using the Hanes-Woolf plot graphical linearization of the Michaelis-Menten equation following Duhamel et al., 2011.

Protein extraction and digestion

All plastics materials were washed with ethanol and dried before usage. All samples from one station were processed together in one extraction and digestion cycle. The frozen Sterivex filter cartridges were transported to the laboratory on ice and cut open with a tube cutter. The filters were cut out from their holders with razor blades and placed into 2 ml microfuge tubes (Eppendorf). Following previously established protocols(Held et al., 2020; Saito et al., 2014)Click or tap here to enter text., proteins were extracted in a 1 % sodium dodecyl sulfate (SDS) buffer for 15 min at 20 °C, followed by 10 min at 95 °C for denaturation, and 1 h at 20 °C while shaking at 350 rpm. The protein extract was then centrifuged at 13.5 rpm for 20 min, with the impurities-free supernatant collected and then spin-concentrated for 1 h in 5 kD membrane filters (Vivaspin, GE Healthcare). Total protein concentrations were then measured by bicinchoninic assay (BCA) (Pierce) on a Nanodrop ND-1000 spectrophotometer (ThermoScientific). Proteins were left to precipitate in a 50:50 solvent mixture of methanol and acetone (Fisher) with 0.004 % concentrated HCl (Sigma, ACS 37 %) for 5 days at -20 °C. At the end of the precipitation period, samples were centrifuged at 13.5 rpm at 4 °C, supernatants were removed, and the remaining protein pellets were vacuum-dried (DNA110 Savan SpeedVac, ThermoFisher). Pellets were redissolved in 50 µl SDS buffer, and the post-precipitation total protein concentrations were measured via a second BCA assay to assess recovery. The protein extracts were digested with the proteolytic enzyme trypsin (1 µg per 20 µg protein; Promega #V5280) in a polyacrylamide tube gel(Lu and Zhu, 2005). The digested samples were concentrated by vacuum drying and stored at -20 °C until analysis. The final volume was recorded to calculate the total protein concentration in the processed sample, typically ~1 µg μl^{-1}.

Target protein selection

Protein biomarkers for *Synechococcus* and *Prochlorococcus* were chosen to detect DIP stress (PstS) and related coping mechanisms via DOP hydrolysis (PhoA and PhoX) in our samples (Table 2). PstS is the substrate-binding protein of the high-affinity phosphate ABC (ATP-Binding Cassette) transporter, which is upregulated under low intracellular phosphate concentrations via the *pho* regulon and has previously been used as an indicator of DIP stress (Cox and Saito, 2013; Martiny et al., 2006; Scanlan et al., 1993). PhoA and PhoX are the Zn/Co-dependent and Fe-dependent alkaline phosphatases, respectively, which facilitate the acquisition of phosphorus from the DOP pool.

Table 2 Details on the quantified peptide biomarkers that are used to represent each protein in the subsequent plots and discussions. For Prochlorococcus strains, 'HL' and 'LL' refer to high-light and low-light adapted strains, respectively.

Protein	Quantified peptide (amino acid sequence)	Isolate strains with this peptide

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Synecho-coccus	**PhoA**	HYIAVALER	WH8102 (clade III)
	PhoX	SQAGAELFR	WH8102 (clade III)
	PstS	WFQELAAAGGPK	RCC307 (clade X)
Prochlorococcus	**PhoA**	IYVIDPSSSPALLER	MIT9311 (clade HL II) MIT9312 (clade HL II) MIT9314 (clade HL II)
	PhoX	GNLWIQTDGK	MIT9314 (clade HL II)
	PstS	LSGAGASFPAK	MIT9301 (clade HL II) MIT9302 (clade HL II) MIT9311 (clade HL II) MIT9312 (clade HL II) MIT9314 (clade HL II) SB (clade HL II) NATL1A (clade LL I) NATL2A (clade LL I)

185

186 The criteria for a peptide of the protein biomarker to be used for quantification were as
187 follows. Firstly, we attempted to minimise the presence of methionine and cysteines because they are
188 subject to oxidation and cause modifications of the mass-to-charge ratio (m/z) during the analyses.
189 Secondly, the specificity and least common ancestor of each tryptic peptide was assessed using
190 METATRYP (https://metatryp.whoi.edu/) (Saunders et al., 2020). It has been demonstrated that
191 carefully selected tryptic peptides, screened by using tryptic peptides databases made from genome
192 sequences like METATRYP, can be used to identify specific proteins in mixed microbial assemblages
193 to the species or even sub-species (ecotype) taxonomic resolution (Saito et al., 2015). Finally, the
194 performance of each precursor ion was visually inspected in Skyline (MacLean et al., 2010) for peak
195 shape and signal to noise-ratio during uncalibrated test measurements using a target list containing
196 many peptides of cyanobacterial alkaline phosphatases on a subset of the incubation samples.

197 **Isotopically labelled standard peptides**

198 The absolute quantitation of the target peptides was achieved using heavy nitrogen isotope-labelled
199 peptide standards (Saito et al., 2020). Briefly, DNA was synthesized containing the reverse-translated
200 gene sequences for our target peptides interspaced with spacer sequences and ligated with a
201 PET30a(+) plasmid vector using the BAMHI 5' and XhoI 3 restriction sites (Novagen; obtained
202 through PriorityGENE, Genewiz). Different nucleotide sequences were used to encode for the spacer
203 (amino acid sequence: TPELFR) to avoid repetition. As per manufacturer instructions, the plasmid
204 was suspended in TE buffer (10 mM Tris-HCl, 1 mM ethylenediaminetetraacetic acid) to 10 ng μl^{-1}
205 and of this 1 µl was added to 20 µl competent Tuner(DE3)pLysS *E. coli* cells on ice. The cells were
206 heated to 42 °C for 30 sec to initiate transformation, followed by 2 min on ice. At room temperature,
207 80 µl ^{15}N-enriched (98 %, Cambridge Isotope Laboratories), kanamycin-containing (50 µl ml^{-1}) SOC
208 medium was added, and cells were incubated for 30 min at 37 °C at 300 rpm. Subsequently, 25 µl
209 were transferred to pre-heated (37 °C) 50 µg ml^{-1} agar plates and incubated overnight. One colony was

added to 500 µl ^{15}N-enriched SOC medium containing 50 µl ml^{-1} kanamycin as a starter culture and incubated for 3 h at 37 °C at 350 rpm. Next, 200 µl of the starter culture were transferred into 50 ml flat incubation flasks with 10 ml SOC medium and incubated for approximately 3 h at 37 °C and 350 rpm until the optical density at 600 nm reached 0.6. Protein production was induced by the addition of 100 mM isopropyl β-D-1- thiogalactopyranoside to the culture and incubating at 25 °C overnight. Inclusion bodies were initially harvested using BugBuster protein extraction protocols (Novagen). The remaining pellet containing the inclusion bodies, i.e. the insoluble protein fraction, was resuspended in 400 µl 6 M urea, left on the shaker table at 350 rpm at room temperature for 3 h, and then moved to the fridge overnight. The next morning, the proteins were reduced, alkylated, and digested with trypsin as outlined above for the bioassay samples, and stored frozen at -20 °C until use.

Absolute protein quantitation

To determine the absolute concentration of the peptides in the heavy peptide mixture, commercial standard peptides of known concentration were used. In addition to the peptides of interest, a range of tryptic peptide sequences from commercially available standards (apomyoglobin, Sigma; Pierce Bovine Serum Albumin, ThermoFisher) were included in the original plasmid design. Using these, the calibrated concentration of the heavy peptide mixture had a relative standard deviation of 57 %, with the standard deviation resulting from the cross-peptide and cross-replicate variability (n=3) (Fig. S1). A systematic method-focused study addressing the precision and accuracy of these measurements as well as the development of reference materials will be essential for using absolute quantitative proteomics in the marine environment in the future(Saito et al., 2024). The linear performance range of each heavy peptide standard was assessed using standard curves of the peptide mixture. Targeted proteomic measurements were made by high pressure liquid chromatography with tandem mass spectrometry (HPLC-MS/MS) on an Orbitrap Fusion Tribrid Mass Spectrometer (ThermoFisher). Two µg of each sample diluted to 10 µl in buffer B (0.1 % formic acid in acetonitrile) was spiked with 10 fmol µL^{-1} of the heavy peptide mixture and injected into the Dionex nanospray HPLC system at a flow rate of 0.17 µl min^{-1}. The chromatography consisted of a nonlinear gradient from 5 to 95 % of buffer B with the remaining concentration consisting of buffer A (0.1 % formic acid in LC-grade H_2O). Precursor (MS^1) ions were scanned for the m/z of the heavy peptide standards and their natural light counterparts. The mass spectrometer was run in parallel reaction monitoring mode and only peptides included in the precursor inclusion list were selected for fragmentation. Absolute peptide concentrations were calculated from the ratio of the peak areas of the product ions (MS^2) of the heavy peptide of known concentration to the natural light peptide (calculated in Skyline (MacLean et al., 2010)). Manual validation of peak shapes was performed for each peptide and sample. Differences between samples with regards to filtration volume, initial protein mass and recovery after precipitation were accounted for. Final peptide concentrations will hereafter be used to represent corresponding protein concentrations, with the caveat that the measurements are not able to discern

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active versus non-active proteins. The status of metalation and if the protein is correctly folded or functions as a polymeric complex cannot be determined from this method.

Identification of significant responses to metal additions

Changes in protein concentrations in response to metal additions were compared relative to the unamended control treatment after 48 h. This approach accounts for any bottle effects. Due to the unique challenges of ocean proteomics sampling and large-scale trace-metal clean bioassays, treatment replication was limited to n = 2 at 54 °W, 50 °W and 45 °W and to n = 3 at 31 °W. Many statistical tests assume normal distributions, which for n = 2 is not assessable. Therefore, in our case, significant differences in protein concentrations were evaluated using a two-fold change criterion, in which the concentrations in all replicates of the metal treatments must lie outside a two-fold change in the average ± one standard deviation of the control to be deemed a significant response. The fold-change in expression and in particular the two-fold change is a commonly used metric to identify proteins that are significantly more or less expressed across different conditions(Carvalho et al., 2008; Lundgren et al., 2010; Zhang et al., 2006).

For the biogeochemical parameters measured in the bioassays, i.e. Chl-*a*, APA and cell counts replication was not limited to n=2 in most cases. Where n=3, ANOVA (a=0.05) followed by Tukey posthoc tests were applied to compare the Control treatments with other treatments.

Results and Discussion

Biogeochemical setting

The oligotrophic subtropical North Atlantic is marked by high deposition of Saharan desert dust, delivering large amounts of Fe and other lithogenic trace metals to the surface ocean (Kunde et al., 2019). During JC150, contrasting biogeochemical regimes existed in the western and eastern basin with high-metal, low-phosphorus, low-nitrogen surface waters at the 54 °W and lower-metal, higher-phosphorus, higher-nitrogen surface waters 31 °W Mahaffey et al. (2025; companion article). Furthermore, *Synechococcus* was two-fold more abundant in the west than in the east, whilst *Prochlorococcus* was more than six-fold more abundant in the east than in the west and numerically more abundant than *Synechococcus* throughout. Overall, the stations at 50 °W and 45 °W exhibited biogeochemical intermediates to the conditions in the east and west. The confluence of gradients in both DIP and trace element availability, as well as clear shifts in microbial community structure, provide a natural field laboratory to probe how environmental drivers differentially influence the contributions of dominant microbes to whole-ecosystem enzyme activity.

Table 3 Date, location and biogeochemical conditions at 40 m depth at the start (t_0) of the bioassays. Biogeochemical parameters are presented as the average ± one standard deviation of replicate t_0 samples, except for the singlet samples of

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DOP at Station 4 and dCo at all stations. Mixed layer depths (MLD; defined after [52]) averages over multiple days, as these were not always determined on the exact day of bioassay set-up.

	Parameter	Station at 54 °W	Station at 50 °W	Station at 45 °W	Station at 31 °W
General	Date	11th July 2017	15th July 2017	19th July 2017	5th August 2017
	Location	22 °N 54 °W	22 °N 50 °W	23 °N 45 °W	22 °N 31 °W
	SST (°C)	27	27	26	25
	MLD (m)	24 ± 3 (5th to 8th July)	33 ± 1 (12th to 15th July)	42 ± 9 (17th to 20th July)	51 ± 8 (4th to 8th August)
Macronutrients	DIP (nM)	3.7 ± 2.1	3.7 ± 1.0	3.4 ± 0.8	14 ± 0.70
	DOP (nM)	87 ± 7.5	137 ± 39	112	129 ± 29
	DIN (nM)	1.5 ± 1.9	1.66 ± 0.56	3.36 ± 1.0	6.2 ± 0.0
	APA (nM h^{-1})	2.8 ± 0.21	2.86	2.48 ± 0.10	1.15 ± 0.08
Trace metals	dFe (nM)	1.26 ±0.06	0.53 ± 0.06	0.83 ± 0.00	0.23 ± 0.05
	dZn (nM)	0.25 ± 0.14	0.46 ± 0.09	0.14 ± 0.01	0.04 ± 0.01
	dCo (pM)	11.0	11.1	13.0	13.9
Phytoplankton community	*Synechococcus* (cells ml^{-1})	$3.4 \pm 0.55 \cdot 10^3$	-	-	$1.6 \pm 0.26 \cdot 10^3$
	Prochlorococcus (cells ml^{-1})	$29 \pm 0.37 \cdot 10^4$	-	-	$181 \pm 0.37 \cdot 10^4$
	Chl-*a* (µg L^{-1})	0.064 ± 0.01	0.055 ± 0.01	0.110 ± 0.06	0.149 ± 0.005

Alkaline phosphatase responded differently in the traditional vs proteomic bioassays

We measured responses in alkaline phosphatase activity (traditional bioassay) and enzyme identity and provenance (proteomic assay) to metal additions across the North Atlantic gyre. Both assays were performed on matched samples at four locations (St 2, 3, 4, 7; Figure 1), allowing direct comparison of the results. In all cases for the traditional assay (shown as circles in Figure 1), bulk alkaline phosphatase activity did not increase significantly upon metal additions (see also Figure 2). However, the proteomic assay revealed that specific alkaline phosphatases did respond positively to the metal additions (squares in Figure 1), depending on the location and metal added. The discrepancy between the traditional bioassay (no response) and proteomic assay (specific, albeit patchy responses) merited additional exploration of the two approaches, which we detail below. One important explanation emerges from the fact that the APA assay covers the entire microbial community (i.e. everything from bacteria to eukaryotes) but the proteomics measurements are specific to a subset of *Prochlorococcus* and *Synechococcus*.

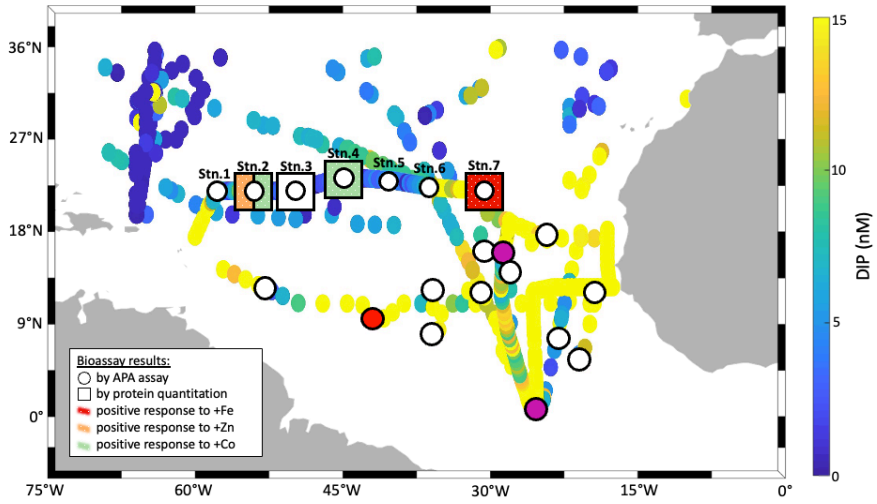

Figure 1 Summary of traditional and molecular bioassay results from this manuscript and others. Map of the North Atlantic showing surface phosphate concentrations (compiled by Martiny et al., 2019 and augmented with data from Browning et al. 2017. Overlain are locations of bioassays, where the response of APA to metal additions was tested (circles), and of bioassays, where the absolute concentration of the alkaline phosphatase proteins was measured in response to metal additions (squares). Bioassays of the present study include longitudinal station labels. The others are from Mahaffey et al. (2014)) and Browning et al. (2017) as well as from additional bioassays during JC150 Mahaffey et al. (submitted as a companion to this article). but where no protein measurements were made. Symbols at bioassay locations are coloured in orange, green or red, if a positive response was observed upon addition of Fe, Zn or Co respectively.

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Variability in the response to metal additions in the traditional bioassays

As mentioned above, there was no significant response in bulk alkaline phosphatase activity to metal additions at any of the stations. Here we focus on Stations 2, 3, 4, and 7, where matched proteomic assays were also conducted. However, alkaline phosphatase assays were conducted on incubations at all seven stations, and there was no response to metal addition at any of them, nor in APA rates normalized to chl-a (Figures S2-S5 and see Table S6). Given this, we sought to address whether there were other observable shifts in microbial activity as a result of the metal additions, including in Chl-A (a proxy for phytoplankton growth) and cell counts for *Prochloroccocus* and *Synechococcus* (Figure 2). While there were no statistically significant changes, either positive or negative, in any of these conventional assays, there were differences between replicate incubation bottles and within the basal conditions across the stations. Despite our efforts to homogenize the incubations and work in large volumes, this variation seems to result from stochasticity of sampling the low biomass system of the subtropical North Atlantic gyre, particularly since there is clear variation among the control bottles as well as in the amended conditions. It is also consistent with past literature including Browning et al., 2017 in which only one in eight experiments showed a metal driven response in APA and Mahaffey et al., 2014 in which there was a positive response of APA to

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Zn only in the eastern basin (see Figure 1). One possible explanation for the presence of the many null responses across the basin is that organisms could be re-allocating metals towards use in alkaline phosphatases when under phosphorus stress. Supporting this idea, a comparable re-allocation mechanism of cellular Fe between metalloproteins involved in biological N_2 fixation and photosynthesis has previously been demonstrated in the diel cycle of *Crocosphaera watsonii* (Saito et al., 2011b). Regardless, the absence of significant responses in the biogeochemical parameters contrasted notably with the observed responses in protein data detailed below.

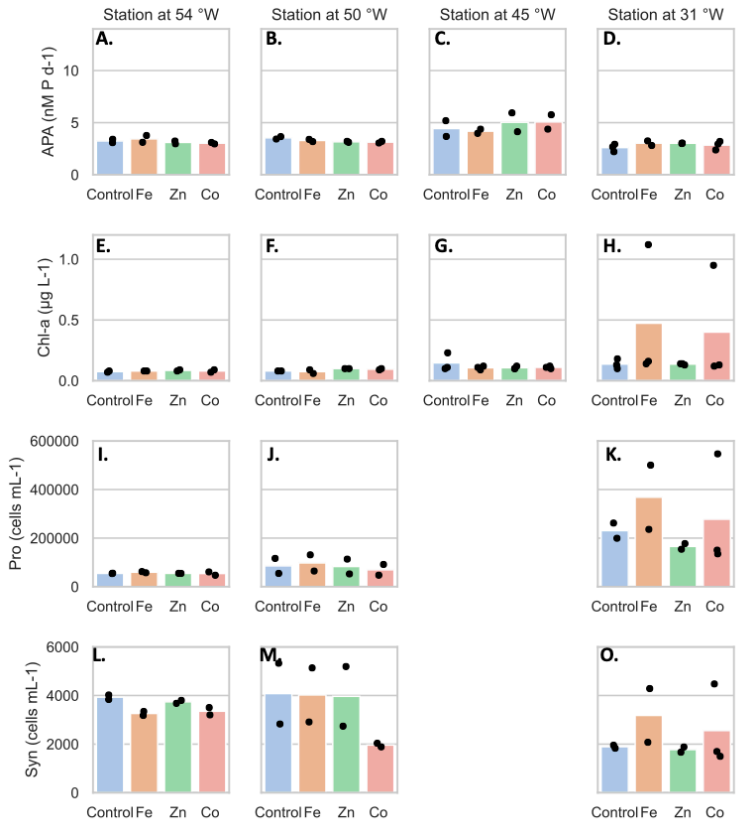

Figure 2. Mean concentrations (bars) of the bioassay parameters after the addition of Fe, Zn and Co at the four stations), specifically concentrations of chlorophyll a (µg L^{-1}), (A-D) rates of alkaline phosphatase (nM d^{-1}), (E-H) Prochlorococcus *abundance (cells mL^{-1}) (I-K) and* Synechococcus *abundance (cells mL^{-1}) (L-O). Dots represent the concentrations of each replicate. Note the data gap for cell counts at the Station at 45 °W.*

Strain-resolved cyanobacterial alkaline phosphatases did respond to metal additions

In contrast to the bioassay results, there were clear changes in proteomically-resolved alkaline phosphatase concentrations after metal additions. We focused on the enzymes PhoA and PhoX and

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used peptides that were specific to one or more strains of either *Prochlorococcus* or *Synechococcus* (; 2) and represent a subset of the population of alkaline phosphatase enzymes in the ocean. We note that marine alkaline phosphatases are found at different subcellular localizations and are also known to be secreted to the environment (i.e. into the dissolved phase) (Li et al., 1998; Luo et al., 2009). Our measurements focus on the alkaline phosphatase associated with microbial cells (i.e. the particulate phase). Coming from an overview of the enzyme concentrations across isoforms, taxa and bioassays, we will discuss how these compare to the APA assay involving fluorogenic substrates.

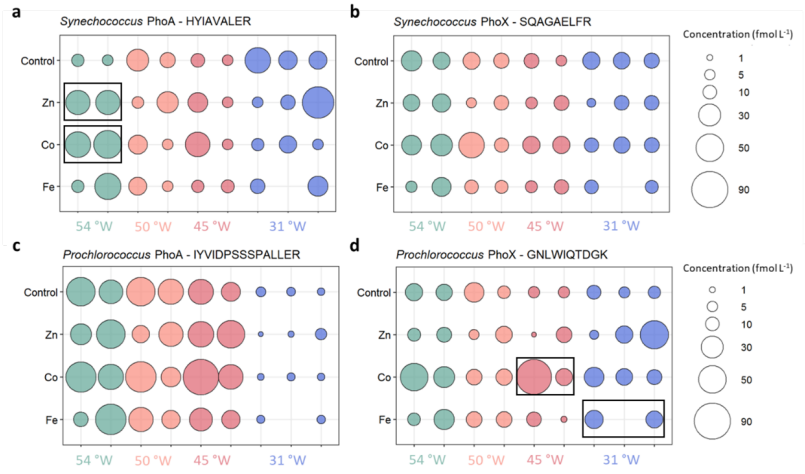

Figure 3 Absolute concentrations of the alkaline phosphatases PhoA (left column) and PhoX (right column) of Synechococcus (top) and Prochlorococcus (bottom) in the different metal treatments or the unamended control at the four probed stations. Bubbles of the same colour are replicates of the same treatment and show the concentrations as fmol enzyme per L seawater. Black boxes indicate significant change from Control treatment.

The results of all measured alkaline phosphatase concentrations are shown in Fig. 3 and all data is compiled in Table S5. *Synechococcus* PhoA and PhoX concentrations in the control treatments ranged from 6 to 43 fmol L^{-1} and 6 to 26 fmol L^{-1}, respectively, with no clear cross-basin trend despite a strong west-to-east decreasing gradient in *Synechococcus* cell abundance (Table 3). Similarly, *Prochlorococcus* PhoA and PhoX concentrations in the control treatments ranged from 2 to 55 fmol L^{-1} and 6 to 23 fmol L^{-1}, respectively, but with elevated PhoA at in the west and the lowest concentrations at 31 °W, which is opposite to the west-to-east increasing gradient in *Prochlorococcus* cell abundance. This suggests that we observed a gradient in DIP/trace metal nutrient stress for *Prochlorococcus*, but not for *Synechococcus*.

Our measured alkaline phosphatase concentrations were similar, albeit at the lower end, to concentrations reported for other cyanobacterial enzymes and nutrient regulators from the North Pacific (~10^{-1} to 10^{3} fmol L^{-1}) (Saito et al., 2014). Interestingly, our alkaline phosphatase

concentrations occurred at the same concentration range as other macronutrient stress indicators (response regulator protein PhoP, sulfolipid biosynthesis protein SqdB, nitrogen regulatory protein P-II), all of which did not exceed tens of fmol L^{-1} (Saito et al., 2014). In contrast, concentrations of the *Prochlorococcus* PstS transporter protein were higher, ranging from 95 to 472 fmol L^{-1} (Table S5). This is within the concentration range of other cyanobacterial nutrient transporters, such as the urea transporter UrtA, measured previously (Saito et al., 2014). In mediating nutrient stress, particularly phosphorus stress, the relative role of transporter proteins (such as PstS) versus other strategically deployed enzymes like alkaline phosphatase in the oligotrophic specialists *Synechococcus* and *Prochlorococcus*, represents an interesting avenue for future research.

Evidence for direct biochemical regulation of certain alkaline phosphatases by metals

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Our strain-specific, quantitative proteomics approach allowed us to resolve contrasting responses across the sites. The responses differed with varying phytoplankton species, alkaline phosphatase form and stimulating metal addition, consistent with differences in the biogeochemical regimes (Table 3). At the iron-rich westernmost station (Station 4; 54 °W), the *Synechococcus* PhoA concentration increased six- and seven-fold upon addition of Zn (to 38 ± 0.56 fmol L^{-1}) and Co (to 47 ± 6.8 fmol L^{-1}) relative to the control (6.7 ± 1.5 fmol L^{-1}), respectively. At one intermediate Station (45 °W), the *Prochlorococcus* PhoX concentration increased 8-fold upon addition of Co relative to the control. Notably, a direct response of alkaline phosphatase to an addition of Co has not been shown in the field before. In contrast, at the low iron easternmost station, the *Prochlorococcus* PhoX increased over two-fold upon Fe addition (to 18 ± 2.6 fmol L^{-1}) relative to the control (8.2 ± 2.4 fmol L^{-1}).

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At least three scenarios are possible to explain the increased alkaline phosphatase concentrations of *Synechococcus* and *Prochlorococcus* in seawater in these treatments – two biochemical and one growth driven hypotheses. First, the metal addition may stimulate the production of the alkaline phosphatase enzyme via a direct or indirect metal regulation on the expression of this enzyme, as was previously observed for PhoA with Zn additions in *Synechococcus* cultures (Cox and Saito, 2013). Second, the metal addition may prevent the degradation of the existing alkaline phosphatases by filling empty metal co-factor sites (Bicknell et al., 1985), with the caveat that PhoA is likely to be periplasmic and hence unlikely to be actively degraded (Luo et al., 2009). Both biochemical scenarios allow for increased alkaline phosphatase concentrations at a constant cell abundance. The third explanation is that the alkaline phosphatase concentration increases because the metal addition stimulates overall cell growth, resulting in higher phosphorus demands and hence more production of alkaline phosphatase proteins by the cell. This could manifest itself as higher cell abundances in addition to increased alkaline phosphatase concentration per unit biomass.

While the different scenarios are not mutually exclusive, our quantitative proteomic approach allowed us to discern between biochemical and growth mechanisms by normalising the alkaline

phosphatase concentrations to the total cell counts of *Prochlorococcus* and *Synechococcus*, caveating that the cell counts are not strain-specific, unlike the peptide-based protein measurements. Cell counts did not change significantly across these treatments Mahaffey et al. (submitted as a companion to this article). which means that the trends of increased alkaline phosphatase concentration per L seawater persisted in bioassays (i.e. +Zn and +Co at 54 °W and +Fe at 31 °W; cell counts do not exist for 45 °W) even when converted to the number of alkaline phosphatase enzymes per cell, indicating biochemical regulation as opposed to simply growth of the responsible organism. Specifically, the concentration of *Synechococcus* PhoA increased to 8418 ± 673 enzymes cell^{-1} upon Co addition and to 6057 ± 48 enzymes cell^{-1} upon Zn addition relative to 1025 ± 257 enzymes cell^{-1} in the control at 54 °W, while the concentration of the *Prochlorococcus* PhoX increased to 59 enzymes cell^{-1} upon Fe addition relative to 19 ± 7 enzymes cell^{-1} in the control at 31 °W. Therefore, a direct biochemical metal control on the alkaline phosphatase concentrations during the bioassays is plausible (i.e. either of the first two explanations) and adds weight to the hypothesis for the localised metal-phosphorus co-limitation in the subtropical North Atlantic (Browning et al., 2017; Jakuba, R. Wisniewski et al., 2008; Mahaffey et al., 2014; Saito et al., 2017; Shaked et al., 2006).

These estimates of enzyme copies per cell are potentially underestimates as multiple *Prochlorococcus* and *Synechococcus* ecotypes co-exist and the alkaline phosphatase peptide sequences probed here do not encompass all of them (Table 2). Moreover, it is also possible that there are additional isoforms of alkaline phosphatase present in these organisms that have yet to be identified (Bradshaw et al., 1981). Yet in these marine cyanobacteria, the cellular concentration of alkaline phosphatase was much higher compared to a measurement in the model bacterium *E. coli*, which contained ~4 PhoA copies cell^{-1} (Wiśniewski and Rakus, 2014). This underscores the ecological demand for alkaline phosphatases due to the significant depletion of phosphorus in the marine environment. It is yet to be determined whether the per-cell estimates of alkaline phosphatases presented here are the norm for marine cyanobacteria, or whether these estimates are exceptionally high due to the prevalence of phosphorus stress in our study region.

While PhoX enzymes are unknown to use Co as a metal co-factor and the response at 45 °W warrants further investigation, the substitution of Zn with Co in PhoA has been hypothesised previously based on the distributions of trace metals and phosphate in the Sargasso Sea (Jakuba, R. Wisniewski et al., 2008; Saito et al., 2017). The results from 54 °W support this hypothesis as the addition of both Zn and Co were associated with almost equal increases of the *Synechococcus* PhoA concentration relative to the control. It is thought that while Zn is the preferred metal centre for PhoA, it is possible to substitute Co for Zn in the protein, such as occurs in *Thermotoga maritima* (Wojciechowski et al., n.d.) *in vivo* and *in vitro* in *E. coli* (Gottesman et al., 1969). Metabolic substitution capabilities between Zn and Co in carbonic anhydrases have previously been identified in marine phytoplankton, with similar or slightly reduced growth rates for a range of marine diatoms and

coccolithophores, when Zn was replaced with Co in carbonic anhydrases (Dupont et al., 2006; Kellogg et al., 2020; Morel et al., 2020; Price and Morel, 1990; Sunda and Huntsman, 1995; Timmermans et al., 2001; Xu et al., 2007; Yee and Morel, 1996). However, in certain organisms such as in the coccolithophore *Emiliana huxleyi*, it is possible that Co is the preferred metal co-factor since the growth rate was higher under replete Co than under replete Zn. One reason could be the simultaneous development of ocean chemistry and cyanobacterial metabolism under the Co- and Fe-replete, but Zn-deplete conditions of the ancient ocean ~2.5 Gyr ago (Dupont et al., 2006; Johnson et al., 2024; Saito et al., 2003). Another explanation for Co use in alkaline phosphatases may require maintaining low intracellular availability of Zn to avoid toxicity through inhibition of Co insertion by high Zn into cobalamin (Hawco and Saito, 2018). Supporting this, the cyanobacterium *Synechococcus bacillaris* and *Prochlorococcus* were found to have absolute Co requirements for growth(Sunda and Huntsman, 1995). Together with these aspects, our insights from the bioassay response at 54 °W merits further investigations into whether *Synechococcus* can interchange Zn and Co in PhoA, and indeed which metal is preferred. This would be an important insight for considerations of stoichiometric plasticity and niche partitioning across the vast Zn- and Co-depleted regions of the ocean, especially where dZn can become depleted to levels similar or below dCo (Kellogg et al., 2020).

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Across all bioassays, the addition of Zn did not increase the concentration of any presumably Fe-dependent PhoX, and the addition of Fe did not increase the concentration of any presumably Zn- or Co-dependent PhoA. In other words, no significant unexpected responses were observed. Nevertheless, there are some non-significant trends upon the addition of Co that warrant further study. For example, the addition of Co increased the putative Fe-containing *Prochlorococcus* PhoX protein concentration dramatically in one of the replicates at 45 °W and hence, Co could be an efficient metal co-factor in PhoX (as in the bacterium *Pasteurella multocida* (Wu, Jin-Ru et al., 2007)), or at least, directly or indirectly stimulate production of PhoX. This contrasts with the results of Kathuria & Martiny (Kathuria and Martiny, 2011), who hypothesized an enzyme inhibiting role of Co (and Zn; with Fe untested) for the activity of both *Synechococcus* and *Prochlorococcus* PhoX by replacing Ca^{2+} at the active site.

Taken together, the results of our bioassays suggest that alkaline phosphatase enzymes are affected by trace metal concentrations, and that the response to Zn, Co or Fe may be species or strain specific. The metal effects differed between the responsive enzyme type (PhoA versus PhoX) and phytoplankton species (*Prochlorococcus* versus *Synechococcus*) at contrasting biogeochemical settings across the basin. It is plausible that the significant changes in protein concentration can result directly from the metal addition triggering more alkaline phosphatase production per cell. This demonstrates that the cycling of macronutrients and metals are intermittently linked and that the nature of that linkage depends on microbial community composition

Towards a quantitative, in situ marine metalloproteome of *Synechococcus*

An advantage of absolute quantitative measurements over relative proteomics data is the ability to relate the absolute protein concentrations to other data types, including biological rate measurements and cellular metal stoichiometry. To this end, the concentrations of the Zn, Co or Fe-dependent alkaline phosphatases measured in this study naturally lead to two questions: First, how much metal is allocated as alkaline phosphatase co-factors in the cell, and how does this compare to total cellular metal content? Second, how does the APA estimated from enzyme abundance compare to assay-based APA?

To address these questions, model calculations were performed. Variables other than the absolute concentrations of the alkaline phosphatases were either measured concomitantly during the bioassays, such as cellular metal quotas, cell abundance, APA and DOP concentration, or sourced from the literature such as strain specific contribution to cell abundance, phosphoester contribution to the DOP pool, enzyme kinetics parameters and subcellular enzyme localization (Table S7). We chose to use the *Synechococcus* PhoA concentrations in the control treatments at 54 °W after 48 h in these calculations for three reasons: First, cellular metal quotas of *Synechococcus* but not of *Prochlorococcus* were measured in this treatment, due to limited sampling capacity and small cell sizes for *Prochloroccocu (Sofen et al., 2022)*s. Second, enzyme kinetics parameters of the PhoA rather than the PhoX isoform are well documented in the literature(Lazdunski and Lazdunski, 1969). Third, estimates for the contribution of *Synechococcus* strain WH8102 (to which our measured PhoA is specific) to total *Synechococcus* counts exist from previous studies nearest to 54 °W(Ohnemus et al., 2016). A similar reasoning applied to the phosphoester contribution to the DOP pool. For ease, more detailed explanations, all values, and assumptions are in Tables S3 and S4.

Equation 1a approximates the cellular Zn allocation towards the *Synechococcus* PhoA from the replicate-averaged PhoA concentration in seawater normalised to cell abundance, assuming full metalation of the enzyme with four metal ions per dimer(Coleman, 1992). The cell abundance is a function of *Synechococcus* cell counts and the fractional abundance of strain WH8102, to which the measured PhoA is specific. Equation 1b expresses the results of Equation 1a as a fraction of the total cellular Zn content.

$$\text{Allocated } Zn_{PhoA} = \text{metalated co-factors}_{PhoA} \times PhoA_{SW} / (\textit{Syn.}\text{ abundance} \times \text{WH8102 fraction}) \quad \textbf{(1a)}$$

$$\text{Fractional allocated } Zn_{PhoA} = Zn_{PhoA} / Zn_{\text{total cell.}} \quad \textbf{(1b)}$$

The amount of metal allocated to PhoA in *Synechococcus* is 3,054 atoms cell^{-1}. This translates to a maximum fractional contribution towards the total cellular Zn content of 0.66 % after dividing by cellular Zn measured using SXRF (Table S1). If the co-factor in PhoA is assumed to be occupied by

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Alkaline phosphatase activity and phytoplankton biomass (by Chl-*a* proxy) did not increase significantly upon the metal additions in the bioassays, neither together with the responses observed in the absolute enzyme concentrations at 54 °W, 45 °W and 31 °W, nor in any other treatments or locations (also Mahaffey et al.; submitted). A quantitative explanation for this – and hence a demonstration of the power of proteomics on the organism level - emerges from estimates of enzyme abundance-based enzyme rates, where the APA assay covers the entire microbial community (i.e. everything from bacteria to eukaryotes) but the proteomics measurements are specific to a subset of *Prochlorococcus* and *Synechococcus*. ¶

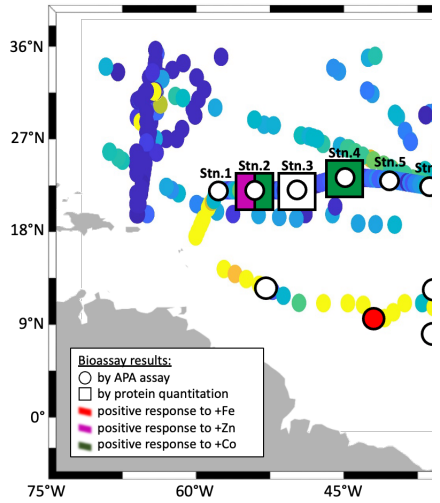

Figure 3 Map of the North Atlantic showing surface phosphate concentrations (compiled by Martiny et al., 2019(Martiny et al., 2019); augmented with data from Browning et al. 2017(Browning et al., 2017)). Overlain are locations of bioassays, where the response of APA to metal additions was tested (circles), and of bioassays, where the absolute concentration of the alkaline phosphatase proteins was measured in response to metal additions (squares). Bioassays of the present study include longitudinal station labels. The others are from Mahaffey et al. (2014)(Mahaffey et al., 2014) and Browning et al. (2017)(Browning et al., 2017) as well as from additional bioassays during JC150 Mahaffey et al. (submitted as a companion to this article). but where no protein measurements were made. Symbols at bioassay locations are coloured in red, purple or green, if a positive response was observed upon addition of Fe, Zn or Co respectively.¶

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Co^{2+} instead of Zn^{2+}, the fractional contribution to the cellular Co content is 38 %, due to the lower total cellular Co content of *Synechococcus* compared to Zn (Table S1). It is possible that the active sites of PhoA are occupied by a mixture of Zn and Co, incompletely metalated, or under competition by metals other than Zn or Co. Nevertheless, these low fractional contributions of PhoA-allocated Zn appear biochemically reasonable, as the majority of Zn in *Synechococcus* appears to be stored in metallothioneins to maintain Zn homeostasis and potentially supply alkaline phosphatases with Zn as needed (Cox and Saito, 2013; Mikhaylina et al., 2022). However, our bioassay results also suggest Zn may not be the preferred co-factor in *Synechococcus* PhoA: The larger response of this enzyme to Co additions at 54 °W suggests the effective substitution of or even preference for Co (Fig. 2). This aligns with evolutionary arguments (see '*Metal control on alkaline phosphatases*') and imply that PhoA is a potential major sink of cellular Co. This also implies that *Synechococcus* growth may be sensitive towards Co-phosphorus co-limitation in the oligotrophic ocean.

Equation 2a approximates the *Synechococcus* PhoA-abundance based hydrolysis rate as a function of the PhoA concentration (converting molarity units to grams using its molecular weight), phosphoester substrate concentration, and Michaelis-Menten kinetics parameters V_{max} and K_m, the maximum reaction rate and half-saturation constant, respectively, derived from an *E. coli* homologue of PhoA (Table S2). Equation 2b expresses the results of Equation 2a as a fraction of the 'total APA', a function of the measured MUF-P assay-based APA with a correction applied for the subcellular localisation of marine alkaline phosphatases, of which only the periplasmic-outwards fraction (~20 to 80 %) is detectable via the MUF-P assay. In other words, the calculated 'total APA' accounts for both the dissolved and particulate activity. Details on made assumptions are in the supplement (Table S2).

$$Rate_{PhoA} = PhoA_{SW} \times molecular\ weight \times V_{max} \times substrate\ /\ (substrate + K_m) \quad \textbf{(2a)}$$

$$where\ substrate = DOP \times phosphoester\ fraction$$

$$Fractional\ Rate_{PhoA} = Rate_{PhoA} \times periplasmic\text{-}outwards\ fraction\ /\ assayed\ APA \quad \textbf{(2b)}$$

The protein abundance-based rates range from 0.00517 nM h^{-1} to 0.213 nM h^{-1} for the Zn-dependent *Synechococcus* PhoA and from 0.00428 nM h^{-1} to 0.0187 nM h^{-1} for the Co-dependent PhoA (Fig. 4a and b), using the *E. coli* kinetics parameters for each metal that are slower under Co coordination. In terms of fractional contributions to total APA, the rate estimates translate to maximally 5.2 % for Zn-PhoA and 0.46 % for the Co-PhoA. Regardless of the choice of enzyme kinetics, it appeared that *Synechococcus* PhoA contributed a small component to the total APA in our bioassays. This concurs with the observed increase in concentration of *Synechococcus* PhoA upon metal addition versus the null response in APA. Applying the same calculation and kinetics parameters for the case of the *Prochlorococcus* PhoA yields abundance-based rates between 0.035

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nM h^{-1} and 1.4 nM h^{-1} for the Zn-PhoA and between 0.029 nM h^{-1} and 0.13 nM h^{-1} for the Co-PhoA, which translate to maximal contributions to the total APA of 35 % and 3.1 %, respectively. These higher rates and fractional contributions compared to the *Synechococcus* PhoA are due to the higher concentrations of the *Prochlorococcus* PhoA than the *Synechococcus* PhoA in the chosen samples. The taxon-specific alkaline phosphatase concentrations illustrate the challenge of interpreting bulk enzyme activities when the functional enzyme class is produced by many biological taxa (cyanobacteria, heterotrophic bacteria, diatoms etc.). In essence, the different bioassay responses shown here demonstrate the need to further develop a "meta-biochemistry' capability to understand biogeochemical reactions at the mechanistic level.

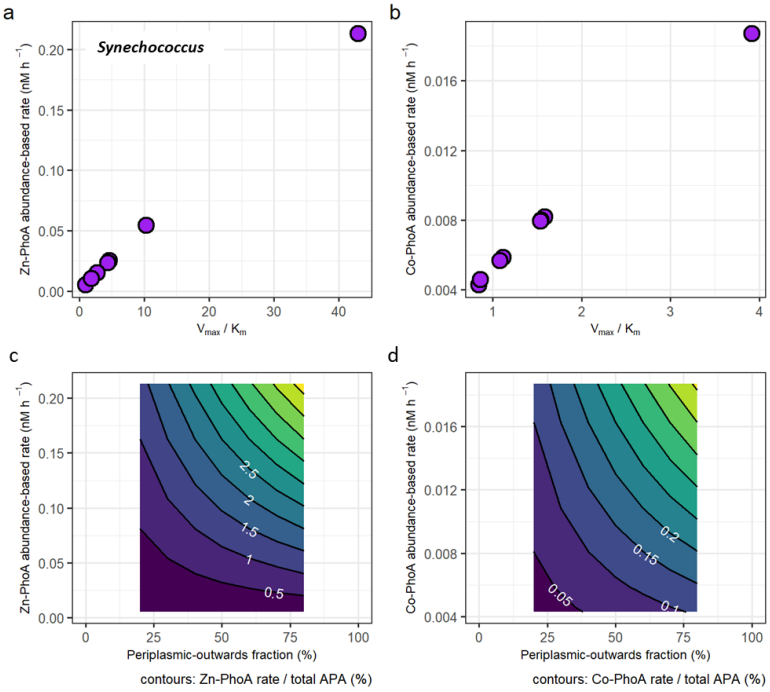

Figure 1. (a) Protein abundance-based APA estimates of Synechococcus Zn-dependent PhoA as a function of different enzyme kinetic parameters V_{max} and K_m. (b) Same as (a), but with enzyme parameters for the less efficient Co-dependent PhoA. (see Table S2). (c) The fraction of the Zn-PhoA abundance-based APA from (a) over the total APA. (d) Same as (c) but using the Co-PhoA rates from (b). Note the scale difference between (a,c) and (b,d).

The enzyme-based rates calculated here may be below that of bulk activity (APA assay) because our protein analysis focused on few species and only in the particulate phase. Alkaline phosphatase is known to be more abundant in the dissolved phase, for example as much as 72% of the APA was observed in Red Sea samples to be in the dissolved phase (Li et al., 1998). Moreover, the

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periplasmic location of alkaline phosphatase has been observed to result in loss during preservation. In a preservation study, PhoA was notably the protein with the lowest recovery in *Synechococcus* WH8102 after a month in storage compared to 101% ± 27% for the fifty most abundant proteins(Saito et al., 2011a). The combination of multiple abundant and rare microbial sources of alkaline phosphatases together contribute to the particulate, and when secreted or lost, dissolved reservoirs that make up the bulk APA.

Conclusions

This study performs taxon-specific alkaline phosphatase isoform analysis via absolute quantitative proteomics on *Prochlorococcus* and *Synechococcus,* coupled to enzyme bioassays. This approach supports the use of Zn, Co and Fe in alkaline phosphatases in the natural oceanic environment, but also adds complexity to our understanding of how these enzymes are regulated in a biogeochemical context. Our mechanistic perspective revealed that these two highly abundant microbes are only minor contributors to bulk APA, which carries important implications for the interpretation of the widely used fluorescent APA assay. Additionally, within this picocyanobacterial class, we observed heterogeneous responses of the alkaline phosphatase enzymes depending on the protein, taxonomy, biogeochemical context, and treatment. This indicates that there is significant biological diversity in the responses of individual marine organisms to experimental treatments that can be resolved by combining enzyme assay measurements with quantitative proteomics. Our results indicate a need for biochemical characterisation of key marine alkaline phosphatases, particularly with regards to their kinetics and metal co-factors as highlighted by the potential importance of Co as a metal co-factor in PhoA and possibly PhoX. Future efforts to understand the biochemical properties of marine microbes will benefit the connected interpretation of molecular, enzymatic, and biogeochemical assays, and in turn our understanding of nutrient cycling in the ocean system.

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Author contributions: NAH and KK wrote the initial manuscript draft. NAH and KK performed the proteomics analysis with help from MM and MAS. KK, NAH, NJW, CD, CM, MCL performed the experiments at sea. BST and ELM conducted the cell quota measurements. MCL, CM, AT and MAS led the research campaign. All authors commented on the manuscript.

Competing interests: The authors declare no competing interests.

Data Availability: Source data for all main and supplementary figures are provided in the supplement. The mass spectrometry proteomics data have been deposited to the ProteomeXchange Consortium via the PRIDE [1] partner repository with the dataset identifier PXD053717.

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