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Elevated trace metal content of prokaryotic communities associated with marine oxygen deficient zones

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Abstract

Little is known about the trace metal content of marine prokaryotes, in part due to their co-occurrence with more abundant particulate phases in the upper ocean, such as phytoplankton and biogenic detritus, lithogenic minerals, and authigenic Mn and Fe oxyhydroxides. We attempt to isolate these biomass signals in particulate data from the US GEOTRACES Eastern Pacific Zonal Transect (cruise GP16) in the Eastern Tropical South Pacific (ETSP), which exhibited consistent maxima in P and other bioactive trace metals, and minima in particulate Mn, in the oxygen deficient zones (ODZs) of 13 stations. Nitrite maxima and nitrate deficits indicated the presence of denitrifying prokaryotic biomass within ETSP ODZs, and deep secondary fluorescence maxima at the upper ODZ boundaries of 10 stations also suggested the presence of low-light, autotrophic communities. ODZs were observed as far west as 99°W, more than 2300 km from the South American coast, where eolian lithogenic and lateral/resuspended sedimentary inputs were negligible, presenting a unique opportunity to examine prokaryotic metal stoichiometries. ODZ particulate P maxima can rival gyre mixed layer biomass concentrations, are highly sensitive to oxygen, and are in excess of amounts scavengable by local Fe oxyhydroxides and acid-volatile sulfides. Even after correction for lithogenic and ferruginous-scavenged metals, ODZ P-maxima are often enriched in Cd, Co, Cu, Ni, V, and Zn, exhibiting particulate trace metal ratios to P that exceed mixed layer biomass ratios by factors of 2-9. ODZ prokaryotic communities may be largely hidden, TM-rich pools involved in the marine cycles of these bioactive trace metals.

Oxygen deficient zones (ODZs) of the Arabian Sea and the Eastern Tropical North and South Pacific (ETNP, ETSP) are sites of diverse microbial productivity that mediate redox cycling of N, S, and C in the virtual absence of oxygen (Ulloa et al. 2012). These zones consist of functionally anoxic cores and associated oxic–suboxic transition zones at the bottom of the upper oxycline, hosting a complex cascade of aerobic and anaerobic microbially mediated biogeo-chemical processes (Tiano et al. 2014). Significant attention has been paid to ODZs in recent decades because of their importance to the global cycles of C and N, their production of the greenhouse gas N_2O , and their ongoing expansion and shoaling in response to climate change (Whitney et al. 2007; Stramma et al. 2010, 2012; Wright et al. 2012).

Low-oxygen marine environments have long been known to contain particle maxima that were originally termed "intermediate nepheloid layers" (Pak et al. 1980; Spinrad et al. 1989), and several recent studies have highlighted the likely biotic nature of these maxima within the ETSP ODZ. Whitmire et al. (2009) used drifter-collected transects of the Peruvian upwelling region to demonstrate consistent particle maxima in association with the upper edge of the ODZ and also noted a low-light, secondary deep chlorophyll maximum at stations where the ODZ is shallow enough to impinge on the euphotic zone. Using ultra-low detection limit oxygen sensors, Tiano et al. (2014) demonstrated that ODZ-associated secondary fluorescence maxima maintain oxygen concentrations in the low nanomolar range, suggesting that the autotrophic upper ODZ layer may diurnally cycle between anoxic and vanishingly oxic conditions, indicating a tight coupling between anaerobic and aerobic

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processes. Flow cytometry and genetic phylogeny conducted at ODZ sites have identified populations (10³ to 10⁴ cells mL⁻¹) of low-light-adapted cyanobacteria in this fluorescent zone, primarily Prochlorococcus (Goericke et al. 2000; Lavin et al. 2010). Acoustic backscatter of larger, oxycline-associated biota (zooplankton and larger eukaryotes) can be a reliable means to observe the base of the upper oxycline (Bertrand et al. 2010) though vertically migrating zooplankton, which feed on settling particulate organic material (POM) (Wishner et al. 1995; Williams et al. 2014), are typically excluded from the ODZ itself by the lack of oxygen (Wishner et al. 2013). Increased cell counts and particle loads in the anoxic ODZ core, by comparison, are correlated with the intensity of its denitrifying activity (Naqvi et al. 1993). Taken together, the upper ODZ appears to be a maximum for autotrophic cyanobacteria and annamox activity (Galán et al. 2009), while heterotrophic denitrification dominates within the ODZ core (Ulloa et al. 2012).

Regardless of the absolute identities of the underlying communities, ODZs are sites of intense and uniquely prokaryotic productivity in the near-micron size fraction (Lavin et al. 2010; Close et al. 2014). Furthermore, the absence of an underlying sulfidic zone, as is consistently present beneath the suboxic zones of the Black Sea and Cariaco Basin, precludes the intense and persistent diffusive metal fluxes associated with euxinic environments (Lewis and Landing 1991, 1992; Ho et al. 2004; Yiğiterhan et al. 2011). This makes the ETSP ODZ, especially offshore away from continental shelf influences, a good environment for examining the elemental composition of prokaryotic communities in the relative absence of lithogenic particles, eukaryotic phytoplankton, and detrital mixed layer biomass. These interfering phases typically mask prokaryotic biomass signatures in the surface mixed layer and near-shore environments. Particulate interferences unique to the suboxic water column, such as authigenic Fe and Mn oxyhydroxides that form and cycle there (Senn et al. 2015), do require special consideration, however.

Examinations of bioactive particulate trace metals (pTMs) in prokaryotes have typically focused on trace metal toxicity, e.g., in human-influenced coastal or sedimentary environments and wastewater bioreactors. While extremely well studied in regards to their macronutrient cycling, ODZ-representative cultured organisms such as denitrifiers, ammoniaoxidizing archaea and bacteria (AOAs and AOBs), have not to our knowledge been rigorously examined in regards to their trace metal content. Vogel and Fisher (2010) investigated uptake and utilization rates of Fe, Mn, Cd, and Zn, among other elements, in five bacterioplankton species in surface seawater cultures and found that bacterial accumulation of Fe may be significant in the surface ocean. Other pTM-prokaryote associations have been reported for Fe (Tortell et al. 1996; Sunda and Huntsman 1997; Schmidt and Hutchins 1999), Cu, and Pb (Dixon et al. 2006; Rossi and

Jamet 2008), Cd and Zn (Sunda and Huntsman 1995; Binet et al. 2003; Iyer et al. 2005), Mn (Francis et al. 2001), and V and Ni (Shirdam et al. 2006), though only a few examined prokaryotic strains relevant to the open ocean or ODZs. With prokaryotic biomass accounting for potentially half of oceanic POM (Cho and Azam 1988; Fuhrman et al. 1989) and despite these organisms' crucial global biogeochemical roles, there remains a surprising dearth of information on prokaryotic metal utilization in natural marine environments.

Redox-active trace metals, many of which are ultimately delivered to sediments via settling organic matter, also have importance as sedimentary paleotracers. Bioactive pTMs have been used to record the past extent of ODZs via sedimentary deposition of Ba, Cu, and Cd (Tribovillard et al. 2006). Sedimentary redox state, which can be influenced by ODZ extent due to its impingement along the shelf, is thought to control sedimentary retention of the paleoredox tracers Mo, Re, U, and V (Nameroff et al. 2004). Tribovillard et al. (2006) indicated that Ni and Cu may be paleoproxies for periods of high organic carbon delivery to sediments, as these pTMs can be preserved within authigenic sedimentary sulfides. In an extensive sediment core survey beneath the ETSP ODZ, Böning et al. (2004) indicated a biotic source, presumed to be the overlying euphotic zone, for Cd, Cu, Ni, Zn, and other pTMs that have delivery rates dominated by the downward vertical flux of organic matter. Metal stoichiometries and bulk TM associations for local biomass, including eukaryotic phytoplankton and ODZ prokaryotes, are thus relevant to interpretation of sedimentary paleoproxies, in addition to understanding the communities themselves.

Here we report pTM enrichments found in bulk particles associated with heterotrophic and autotrophic prokaryotic communities of the ETSP ODZ, sampled during the US GEO-TRACES Eastern Pacific Zonal Transect (EPZT) in Oct–Dec 2013 (Fig. 1). The pTM composition of these communities is often significantly different, and pTM–enriched, compared to mixed layer communities. Prokaryotic biomass associated with the ODZ may influence the marine cycles and sedimentary imprint of trace metals in the ETSP and other oxygen– deficient regions.

Materials and methods

Sample collection, analyses, and data access

All particulate samples were collected on the US GEOTRA-CES EPZT cruise GP16 from Lima, Peru to Tahiti in Oct–Dec 2013 (Fig. 1). Seawater was collected with the non–contaminating GEOTRACES rosette and filtered onto acid–cleaned, 25-mm diameter, 0.45 μ m pore–size SuporTM (polyethersulfone) filters in the GEOTRACES sampling van (Cutter et al. 2010; Planquette and Sherrell 2012). Typical filtration volumes through the analyzed filter halves were 4.4 L±1.4 L seawater (median±one SD). Digestion and analytical



Fig. 1. Key stations from *R/V Thompson* cruise TT303 (US GEOTRACES Eastern Pacific Zonal Transect GP16; Oct–Dec 2013), including all stations directly referenced in the text. Inset: "shelf" stations near the Peruvian coast, including Sta. 1 which is plotted as Sta. 5.5 in all other figures for longitudinal consistency. Sta. 6–13 are "ODZ" stations; Sta. 14–36 "gyre" stations. Shading indicates bathymetry.

procedures were as previously reported by our laboratory (Twining et al. 2015). Briefly, in a land-based analytical clean lab, filters were halved, digested in a mixture of 4 mol L⁻¹ each HCl/HNO₃/HF optima-grade acids at 110°C in sealed Teflon vials, dried, and brought up in 0.32 mol L^{-1} HNO₃ for analysis via ICP-MS (Thermo Element2). External multi-elemental standard curves and internal spikes of In (for ICP-MS instrumental drift) and Cs (for sample recovery purposes) were used to standardize and monitor elemental recoveries. Recoveries of the certified reference materials (CRMs) PACS-2 (marine sediment) and BCR-414 (plankton) and multi-elemental standard addition spikes into random samples were also used to monitor reported values. Recoveries of CRMs averaged $95\% \pm 8\%$ for PACS-2 and $102\% \pm$ 7% for BCR-414 across all elements. Full CRM recoveries, blank information, and methodological information including the details of hydrothermal sample analyses and laboratory intercalibration within the particle dataset, are available in the dataset metadata available via the NSF data archive at the Biological & Chemical Oceanography Data Management Office (BCO-DMO), dataset #639857.

Samples analyzed for acid-volatile sulfides (AVS) were collected using in situ pumps (McLane) suspended from a trace metal wire (Hytrel-jacketed Vectran), at the same stations and target depths as the GEOTRACES carousel, further described in Ohnemus and Lam (2015). Pumps filtered for 4 h, with the relevant flow path passing through a 51-µm polyester pre-filter, then through paired 0.8-µm pore-size, 142-mm SuporTM polyethersulfone filters. A small portion of the top Supor filter, representing $\sim 2\%$ of the filter area (0.8– 51-µm size-fraction material), was sub-sectioned into polyethylene vials and frozen at -85°C until shipboard AVS analysis, described below. Filter blanks were held within identical filter housings and submerged on one of the deeper pumps on each cast but were not actively filtered through. Blanks were handled identically to samples during processing, which was conducted in a HEPA-filtered environment

using trace metal clean procedures. The slight pore–size mismatch between pump–collected AVS samples and bottle– filtered pTMs (0.8 μ m vs. 0.45 μ m cutoff, respectively) is not expected to influence our results significantly, as used here. AVS data are available through the NSF data archive at BCO-DMO, dataset #646143.

Acid-volatile sulfide determination

Determinations of acid-volatile sulfide (AVS) were made onboard ship and used the method of Cutter and Oatts (1987) that includes acidification, gas stripping, and cryogenic trapping/preconcentration of the generated hydrogen sulfide. In this case, the filter slice was placed in a glass stripping vessel, 10 mL of deionized water added, and the stripper purged with He. The cryogenic trap was then immersed in liquid nitrogen, 10 mL of 1 mol L^{-1} HCl added through a septum in the stripping vessel and the evolved and trapped hydrogen sulfide was then quantified using the gas chromatography/flame photometric detection method of Radford-Knoery and Cutter (1993). The system was calibrated with hydrogen sulfide from a permeation tube (calibrated diffusion rate) and assuming a filtration volume of at least 5 liter (median sample volume: 7.9 liter), the detection limit was 0.2 pmol/L. It should be noted that AVS determined by this method includes monosulfides of Fe, Zn, and Ni, but not those with Cu or Hg, or mixed oxidation state sulfides such as greigite or pyrite (Cutter and Oatts 1987; Radford-Knoery and Cutter 1993).

Profiling CTD data

Reported values for transmissometer voltage (WET Labs, 25-cm path length, 660 nm), fluorescence (Seapoint), and oxygen (Seabird SBE 43) were collected via the GEOTRACES carousel Seacat 9+ (Seabird) CTD after 1 m binning of higher–frequency profiling data; upcast traces are used herein, except when otherwise noted. Station occupations were often longer than 24 h and consisted of multiple casts on three sampling systems (GEOTRACES carousel, McLane pumps, and Oceanographic Data Facility [ODF] rosette) each

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equipped with a profiling CTD. We calculated median mixed layer depths across all profiling downcasts (using a 0.125 sigma-theta density change from the surface [Bishop and Wood 2009]) as well as median depths for the primary deep chlorophyll maximum (DCM) and 10% fluorescence depths (the "particle production zone" or PPZ [Marra et al. 2014; Owens et al. 2015]). These values are reported in Table 1, along with the cast-specific ODZ start depth (upcast, defined and discussed below; Sta. 1–13) or depth of the local oxygen minimum (Sta. 14–36) from the cast on which ODZ bottle particle samples were taken.

Use of oxygen saturation and ODZ/oxycline definitions

We use oxygen saturation expressed as percent, rather than absolute oxygen concentrations, to examine the oxygen sensitivities and distribution patterns within this dataset. Our rationale is twofold and derives from our desire to best utilize the high-resolution oxygen data collected from a moving sensor (CTD-mounted Seabird SBE 43). First, the oxygen gradients of greatest interest-in the upper 10-30 m of the ODZ-are often very steep and narrow. The myriad biotic processes occurring in this region are known to be highly sensitive to nanomolar-level oxygen concentrations, so even slight instrumental hysteresis or response-time lag could lead to oxygen profiles that are offset from reality. We also noted a dataset–wide offset of \approx +7 μM in the SBE 43 sensor calibration, which was the mean oxygen concentration reported by the sensor within the ODZ. We have subtracted this offset given that oxygen concentrations in the ODZ have been shown by more sensitive switchable trace oxygen (STOX) sensors to be functionally nil (<1-10 nM) throughout the region (Tiano et al. 2014). Our use of oxygen saturation in percent purposefully avoids over-interpretation of the absolute oxygen concentrations reported at the lowerend of our sensor's capabilities. Secondly, surface mixed layer temperatures, and thus oxygen saturation values, varied widely across the >8000 km cruise track (Fig. 2A). Surface features from a wide range of mixed layer temperatures can be conveniently coalesced into a narrower range of near-saturation values (i.e., 95-110%) by using oxygen saturation rather than absolute oxygen concentration.

Oxygen saturation was calculated using the equations of Benson and Krause (1984) and the 1-m-binned GEOTRACES carousel SBE 43 oxygen sensor values. Future versions of the cruise dataset may correct the reported sensor concentrations to better align with ODF carousel Winkler oxygen titrations, which report oxygen concentrations below the method's detection limit within the ODZ ($\leq \approx 3 \mu M$).

We determined the upper boundary of the ODZ at each station using the first sustained appearance of curvature in the raw (uncorrected) SBE 43 oxygen sensor data on the upcast during which ODZ particulate samples were collected. This determination was guided, in part, by where the downcast trace indicated oxygen approaches zero. Our definition of the ODZ minimizes temporal separation from bottle sample collection and obviates uncertainty due to sensor hysteresis (vertical lag before reading zero) evident in the downcast data through the same depths. It is also consistent with the results of Tiano et al. (2014) who demonstrated that oxygen sensors are more stable on the upcast, and that ODZ-associated secondary fluorescence maxima typically exhibit ≤ 0.4 µM oxygen which, by our definition, fall almost entirely within the ODZ. The region above the ODZ where oxygen increases towards the surface is referred to as the upper oxycline (up to 90% oxygen saturation); the increase in oxygen with depth below the ODZ will be referred to as the lower oxycline. We lack the sensor capability to directly differentiate the oxic-suboxic ODZ transition zone (where aerobic respiration can occur at the top of the ODZ) from the functionally anoxic ODZ core (where anaerobic respirations dominate). We do, however, use the appearance of fluorescence in the upper ODZ of many stations to delineate autotrophic (fluorescent) ODZ samples from heterotrophic (nonfluorescent) ODZ bottle samples.

Background correction for vertically sourced particles in Table 3

All ODZ particulate element maxima are present among some background of particles sourced from the overlying euphotic zone and oxycline, each likely with distinct remineralization rates and behaviors across the elemental suite. To isolate ODZ biomass elemental signals from particles sourced vertically above the ODZ, specifically for the Sta. 11– 13 stoichiometries reported in Table 3, a means to constrain the profile background was needed. We used a simple linear approach: at each station, reference depths above and below the ODZ were chosen for each elemental profile. A linear fit between the reference depths was used as a baseline in each profile, which was then subtracted from the local ODZ maxima. These values are referred to as "background–corrected."

Pigment analyses

Phytoplankton pigment concentrations were collected from the surface mixed layer and analyzed via HPLC at the University of Hawai'i. Pigments have been grouped according to broad size/function categories—microplankton, nanoplankton, and picoplankton—following Hooker et al. (2005).

Results and discussion

Mixed layer biogeography

The US GEOTRACES EPZT cruise comprised 36 stations across more than 8000 km of the ETSP (Fig. 1). The surface transect spanned large biogeochemical gradients associated with boundary and equatorial upwelling, so key surface mixed layer characteristics (Fig. 2) were used to group stations into three regions. The "Shelf ODZ" Sta. 1–5 (with Sta. 1 plotted as Sta. 5.5 in most figures, based on its longitude) overlie the Peruvian shelf and shelf break. These stations **Table 1.** Left four columns: median depth in meters, across all profiling casts at each station, for the surface mixed layer ($\Delta \sigma_{\Theta}$ from surface = 0.125), primary deep chlorophyll maximum, particle production zone (depth where 10% of primary DCM fluorescence remains), and minimum oxygen concentration (Sta. 14–36 only). Right two columns: on the cast when ODZ bottle particles were collected: ODZ start depth (described in methods), and thickness of the upper oxycline (calculated as the ODZ start depth minus ML depth).

		Mixed layer depth ($\Delta \sigma_{\Theta} = 0.125$)	Primary deep chlorophyll max (DCM) depth	Particle production zone depth	Oxygen minimum depth	ODZ Start depth	Upper oxycline thickness			
Stations		All depths and thicknesses in m								
Shelf ODZ	1 (5.5)	46	34	66		84	37			
stations	2	8	6	30		59	51			
	3	8	6	25		35	27			
	4	16	9	43		50	34			
	5	25	20	50		59	33			
Offshore	6	31	38	80		100	69			
ODZ stations	7	50	36	94		98	48			
	8	46	53	105		114	67			
	9	53	53	109		111	58			
	10	78	60	121		153	75			
	11	51	55	130		190	140			
	12	41	47	124		215	174			
	13	78	62	139		231	153			
Gyre stations	14	89	75	164	322					
	15	74	82	168	375					
	16	56	59	150	365					
	17	57	59	163	372					
	18	36	69	174	359					
	19*	30	72	174	362					
	20	49	62	191	393					
	21	59	74	167	373					
	22	56	82	179	376					
	23	45	74	169	350					
	24	54	68	163	359					
	25	76	92	163	350					
	26	57	84	169	334					
	27	60	65	184	320					
	28	67	78	162	343					
	29	66	74	175	351					
	30	74	81	168	347					
	31	65	67	146	337					
	32	84	84	153	335					
	33	65	84	186	336					
	34	58	52	175	310					
	35	67	57	170	328					
	36	60	74	179	376					

*Data from ODF carousel only.

were characterized by comparatively cool, nutrient–rich, upwelling–influenced water (surface T = 13.9-17.4°C; PO₄ > 1 μ M; NO₃ > 7.5 μ M), shallow mixed layer depths (8–46 m vs.

30–89 m elsewhere), macroplankton pigment dominance (diatoms and dinoflagellates), and strong particle (transmissometer) and particulate P enrichments (particulate P: 169–



Fig. 2. Key surface mixed layer parameters (*y*-axes) from west to east across the transect (station number on *x*-axes, in reverse scale for longitudinal consistency): (**A**) temperature, (**B**) nitrate, (**C**) phosphate, (**D**) particulate P, (**E**) bulk particle concentration (via transmissometer voltage), (**F**) bulk pigment fractionation. Median values are plotted in panel A–E, and Sta. 1 is plotted as Sta. 5.5 for longitudinal consistency. Vertical dotted lines separate the three station groupings, from right to left (east to west): shelf Sta. 1–5 (cool, nutrient–rich water; high particle loads, elevated macroplankton concentrations); ODZ Sta. 6–13 (lower particulate P; declining nutrients; lower particle loads, mixed nano/picoplankton dominance); gyre stations \geq 14 (lowest particulate P and nutrients; warmest temperatures, picoplankton dominance).

235 nM). In contrast, "Offshore ODZ" Sta. 6–13 were characterized by an ecosystem shift to smaller nanoplankton cells, coincident with rising temperatures (18–23°C), declining PO₄ (0.5–1.5 μ M) and NO₃ (0.8–14.7 μ M), and significantly lower particulate P (25.6–45.6 nM). The "gyre" Sta. 14–36 were picoplankton–dominated, exhibiting the warmest surface temperatures (22.7–29.1°C) and the lowest and most narrowly constrained concentrations of particulate P (21.7–32.0 nM), PO₄ (0.20–0.47 μ M), and NO₃ (0.3–3.9 μ M).

Hydrography

Upper waters of the shelf stations are influenced by Equatorial Subsurface Waters (ESSW) moving poleward in the ODZ via the Peru–Chile Undercurrent, while the offshore ODZ stations are dominated by Subantarctic Water (SAAW) moving equator-ward via the Humboldt Current (Silva et al. 2009; Llanillo et al. 2013). Offshore stations are dominated by Subtropical Surface Water (STSW) formed under the high evaporation in the gyre, moving poleward (Fiedler and Talley 2006). More detailed watermass analyses will be presented in forthcoming publications by other GEOTRACES GP16 cruise participants.

Subsurface ODZ

The subsurface ODZ was apparent in a suite of sensor and chemical data available from the GEOTRACES dataset (Fig. 3). As demonstrated by the section oxygen contour lines, all shelf and ODZ stations display strong oxygen consumption in an upper oxycline that begins near the base of the mixed layer. At Sta. 1–13, where oxygen approaches zero at the bottom of the upper oxycline, the appearance of dissolved nitrite, particulate P, bulk particles (measured by transmissometry), a secondary deep fluorescence maximum (at Sta. 1–10), and loss of particulate Mn all indicate a transition into the ODZ, which is approximated in Fig. 3 sections by the 1% oxygen saturation contour. From east to west the ODZ start depth deepens from a minimum of 35 m at Sta. 3 to 231 m at Sta. 13, the westernmost station to achieve suboxic conditions (Table 1). This westward **Table 2.** Upper sections: Ratios used in lithogenic (left) and ferruginous (right) corrections. Upper and bulk continental crust ratios to Ti from (Taylor and McLennan 1995) and are compared to lithogenic–rich benthic nepheloid layer samples near South America (Sta. 3–9; n = 38 samples). Values used in the lithogenic correction are bolded. For elements that lacked good correlation with Ti in eastern boundary nepheloid layers (italicized r^2 values), the larger of upper or bulk continental crust values was used, except for Mn where we use the lower, upper continental crust value to avoid over–correction of a potential carrier phase (Mn oxyhydroxides). Ferruginous corrections are based on metal:Fe ratios in Fe-oxyhydroxide–rich particles sampled near the East Pacific Rise hydrothermal plume at Sta. 18, 2450–2550 m (n = 10 samples). Lower sections: median lithogenic (left) and ferruginous (right) corrections (percent of pTM concentration subtracted ± 1 SD) in the mixed layer and ODZ depth regions. Ferruginous corrections are shown for ODZ samples only, as lithogenic corrections can account for nearly all Fe in mixed layer samples.

Lithogenic end-members (mol:mol Ti) (bold is value used)					Ferruginous end-member (mmol:mol Fe) (Sta. 18 hydrothermal plume samples; <i>n</i> = 10)					
Element	Sta. 3–9 benthic particles	Upper con- tinental crust	Bulk conti- nental crust	r ² vs. Ti in benthic particles	Element	Median	Max	SD	RSD	
Al	42.8	47.5	27.6	0.76	Al	6.88	18.5	4.12	60%	
Ва	_	6.40E-02	1.62E-02	0.08	Ва	1.84	8.38	2.14	116%	
Cd	_	1.39E-05	7.73E-06	0.04	Cd	4.94E-03	7.80E-03	1.30E-03	26%	
Со	_	2.71E-03	4.37E-03	0.11	Co	6.59E-02	8.10E-02	9.20E-03	14%	
Cu	_	6.29E-03	1.05E-02	0.17	Cu	2.40	3.20	0.318	13%	
Fe	10.9	10.0	11.2	0.98	Fe	_		_	_	
La	2.25E-03	3.45E-03	1.02E-03	0.72	La	3.87E-02	5.06E-02	5.10E-03	13%	
Mn	_	1.74E-01	2.26E-01	0.03	Mn	97.3	134	26.7	27%	
Ni	_	5.44E-03	1.59E-02	0.00	Ni	0.116	0.320	0.069	59%	
Р	_	3.61E-01	2.03E-01	0.06	Р	216	250	12	5%	
Pb	1.39E-03	1.54E-03	3.42E-04	0.41	Pb	7.89E-02	2.39E-01	5.66E-02	72%	
Th	6.22E-04	7.36E-04	1.34E-04	0.96	Th	1.33E-04	9.00E-04	3.00E-04	226%	
Ti	_	_	_	_	Ti	0.730	1.67	0.325	45%	
V	4.07E-02	1.88E-02	4.00E-02	0.67	V	4.15	4.42	0.143	3%	
Y	4.01E-03	3.95E-03	2.00E-03	0.70	Y	0.121	0.541	0.135	112%	
Zn	_	1.73E-02	1.09E-02	0.22	Zn	0.845	17.58	5.25	621%	
	_	_	_	_	_	_		_	_	
	Median lithogenic corrections				Median ferruginous corrections					
	Mixed layer	SD	ODZ	SD		ODZ	SD			
Al	_	_	_	_	Al	1%	0.4%			
Ва	3%	6%	1%	1%	Ва	1%	2%			
Cd	0%	0%	0%	0%	Cd	0%	0.2%			
Co	1%	3%	8%	8%	Co	5%	6%			
Cu	1%	2%	2%	1%	Cu	5%	8%			
Fe	100%	25%	24%	20%	Fe	_				
La	6%	9%	11%	9%	La	5%	9%			
Mn	7%	12%	47%	31%	Mn	100%	21%			
Ni	1%	1%	1%	1%	Ni	0%	0.4%			
Р	0%	0%	0%	0%	Р	2%	2%			
Pb	4%	4%	8%	4%	Pb	15%	12%			
Ті	_	_	_	—	Ti	3%	2%			
V	3%	7%	6%	7%	V	20%	22%			
Y	8%	13%	14%	10%	Y	11%	17%			
Zn	1%	1%	2%	1%	Zn	3%	3%			

deepening of the ODZ corresponds to a thickening of the upper oxycline and a deepening of the "particle production zone" (the depth where 10% of primary DCM fluorescence remains [Marra et al. 2014]): the upper oxycline is only 27-m thick at Sta. 3 but more than 150-m thick at Sta. 12 and 13.

Table 3. P concentrations (nM) and element:P stoichiometries (mmol:mol) for several station groupings, comparing mixed layer (ML) and ODZ values (median \pm SD). All values are after lithogenic and ferruginous corrections. Station-by-station data are shown in graphical form in Fig. 8. Station groupings: Sta. 1–5 (shelf ODZ), 6–10 (autotrophic offshore ODZ), and 11–13 (heterotrophic offshore ODZ). Bold: ODZ stoichiometries that differ significantly from ML in the same region (*t*-test, p < 0.05). For Sta. 11–13, back-ground-corrected (ODZ local maxima) P concentrations and heterotrophic stoichiometries are also provided. Right four columns: ratio of median ODZ/ML stoichiometry for each region.

	Sta. 11–13 (heterotrophic ODZ)			Sta.	Sta. 6–10 Sta. 1–5 (she			f) ODZ/ML stoichiometry ratios				
	Mixed layer	ODZ	ODZ Bkgr. Corr.	Mixed layer	ODZ	Mixed layer	ODZ	Sta. 11–13 Bkgr. Corr.	Sta. 11–13 (Het. ODZ)	Sta. 6–10	Sta. 1–5 (shelf)	
P (nM)	30 ± 4	7.6 ± 1.9	3.7 ± 1.8	32 ± 8	18 ± 4	192 ± 79	28 ± 5					
(mmol:mol)												
Cd:P	0.41 ± 0.20	0.95 ± 0.25	1.4 ± 0.4	1.1 ± 0.13	$\textbf{0.57} \pm \textbf{0.13}$	0.54 ± 0.24	$\textbf{0.40} \pm \textbf{0.05}$	3.4	2.3	0.53	0.73	
Co:P	$\textbf{0.27} \pm \textbf{0.06}$	0.43 ± 0.16	$\textbf{0.58} \pm \textbf{0.33}$	$\textbf{0.27} \pm \textbf{0.04}$	$\textbf{0.22} \pm \textbf{0.06}$	0.08 ± 0.05	0.11 ± 0.03	2.1	1.6	0.83	1.4	
Cu:P	1.1 ± 0.20	7.6 ± 1.6	9.4 ± 5.4	$\textbf{0.86} \pm \textbf{0.15}$	5.3 ± 1.1	$\textbf{0.40} \pm \textbf{0.37}$	3.6 ± 1.2	8.7	7.0	6.1	9.0	
Ni:P	3.6 ± 1.7	4.6 ± 1.2	$\textbf{5.2} \pm \textbf{1.9}$	1.0 ± 0.3	$\textbf{4.8} \pm \textbf{1.0}$	0.54 ± 0.39	4.6 ± 0.51	1.4	1.3	5.0	8.5	
V:P	$\textbf{2.6} \pm \textbf{3.3}$	$\textbf{2.9} \pm \textbf{1.0}$	$\textbf{4.6} \pm \textbf{2.8}$	1.0 ± 1.7	$\textbf{2.3} \pm \textbf{0.70}$	0.17 ± 0.16	1.0 ± 0.77	1.8	1.1	2.4	5.9	
Zn:P	$\textbf{0.80} \pm \textbf{0.20}$	$\textbf{3.1} \pm \textbf{0.54}$	$\textbf{3.0} \pm \textbf{1.4}$	$\textbf{0.89} \pm \textbf{0.33}$	$\textbf{2.8} \pm \textbf{0.72}$	1.0 ± 0.88	$\textbf{2.6} \pm \textbf{0.24}$	3.8	3.8	3.1	2.5	

Particulate P maxima in the ODZ

Initial observations of the particulate dataset indicated significant subsurface maxima in particulate P at low oxygen that were distinct from surface mixed layer maxima and suggested in situ production of biogenic material within the ODZ. Particulate P concentrations from the upper 400 m of offshore ODZ stations plotted vs. depth (Fig. 4A) and oxygen saturation (Fig. 4B) exhibit three distinct particle regimes: a mixed layer P maximum associated with oxic mixed layer biomass; the upper oxycline, where P rapidly declines and where Mn oxyhydroxides appear (Figs. 3E, 4C,D); and a distinct, local P maximum associated with the ODZ. When plotted vs. depth, particulate P data collapse into a rather consistent "Martin curve" power law decay below the euphotic zone (Fig. 4B; b-value -1.07, trendline not shown), but concentrations of P in ODZ samples are distinctly enriched.

The ODZ P maxima were coincident with near-complete, vertically sharp loss of particulate Mn, which drops from typical oxycline concentrations of 100–300 pM to < 100 pM in the ODZ (median particulate Mn concentration in the ODZ: 17 pM). This distribution is consistent with Mn oxy-hydroxides precipitated by Mn–oxidizing bacteria just beneath the mixed layer (Cowen and Bruland 1985) being reductively dissolved as they settle into the suboxic ODZ (Fig. 4C,D). In addition to particulate Mn minima, biological utilization of secondary electron acceptors beginning at or near the 1% saturation horizon indicated a distinct shift in water column redox state. Secondary nitrite maxima (Fig. 3A) and nitrate deficits (not shown) indicate that loss of fixed N is occurring within the ODZ.

The P maxima associated with the ODZ are not slight: even after correction for lithogenic phases (crustally derived, refractory particles) and ferruginous phases (authigenic Fe oxyhydroxides; Table 2, discussed later), ODZ P concentrations have 25% and 75% quartiles of 11 nM and 26 nM, respectively, compared to < 5 nM in similar depths of the oxic water column. Linear interpolation between points above and below the ODZ can be used to estimate P concentrations in excess of profile background concentrations. Local P maxima in the ODZ estimated in this manner range between 1 nM and 17 nM. By comparison, surface mixed layer biomass at offshore stations ranges between 22 nM and 46 nM P. Particulate Fe enrichments in the ODZ (shown relative to Ti; local crustal value of 10.9 mol:mol indicated) are also apparent, especially in shelf-associated stations and in the upper ODZ at Sta. 6–9.

To allow side–by–side comparison of ODZ features across stations, parameters were plotted on a depth scale with the upper oxycline/ODZ boundary defined as 0 m (Fig. 5). For stations lacking an ODZ, west of Sta. 13 at 100°W, the depth scale was set to zero at the oxygen minimum (Table 1). ODZ–associated secondary fluorescence maxima are notably confined to a depth range between \approx 10 m above and \approx 40 m below the ODZ start, and fluorescence is absent at westernmost ODZ Sta. 11–13. Particulate P and bulk particle (transmissometry) maxima generally coincide with one another and appear sharply constrained by the upper ODZ boundary, though we note that discrete bottle sampling did not always exactly align with transmissometry particle maxima. Particulate P and bulk particles notably extend more deeply into the ODZ than does fluorescence, suggesting a



Fig. 3. Sections of (**A**) nitrite, (**B**) particulate P, (**C**) bulk particles (transmissometer voltage), (**D**) fluorescence, (**E**) particulate Mn, and (**F**) particulate Fe:Ti, overlaid by contours of oxygen saturation (%; dashed lines and labels) between gyre Sta. 18 in the west and shelf Sta. 2 in the east. Sta. 1 is plotted as Sta. 5.5 for longitudinal consistency. Horizontal scale (station number) allows visual comparison of shelf to offshore stations and is not meant to approximate true spatial distance, though offshore stations west of Sta. 6 are evenly spaced (Fig. 1). The oxycline/ODZ boundary is approximated by the 1% oxygen saturation contour.

shift from autotrophic to heterotrophic biomass as depth within the ODZ increases.

Secondary fluorescence maxima found in the upper ODZ at Sta. 1–10 are likely associated with the pigments of lowlight adapted *Prochlorococcus* (Goericke et al. 2000; Lavin et al. 2010) and perhaps anoxygenic photosynthetic prokaryotes (Repeta et al. 1989; Béjà et al. 2002; Allgaier et al. 2003) that are known to inhabit these low-predation, lowlight niches. Secondary fluorescence maxima are notably absent at the westernmost stations with the deepest ODZs (Sta. 11–13), perhaps indicating insufficient light for autotrophic processes (though photosynthetically active radiation [PAR] was not measured on the CTD). The absence of fluorescence yet continued appearance of local P maxima at Sta.



Fig. 4. Particulate P in the upper 400 m of ODZ and gyre stations (stations \geq 6) plotted (**A**) vs. depth, and (**B**) vs. sample oxygen saturation. For clarity, only samples with \leq 50 nM particulate P are shown. In panel A, shallow (2 m) towed–fish surface samples that lack paired oxygen data are plotted at 105% oxygen saturation. Sample shading and shape indicate general oxygen state: white circles: oxic euphotic zone samples with oxygen saturation \geq 90%; gray squares: upper oxycline (oxygen saturation between 1% and 90%); black diamonds: ODZ (oxygen saturation < 1%). (**C**, **D**) Same as A and B, but for particulate Mn.

11–13 suggest the presence of completely heterotrophic communities in these ODZs, where different (typically enriched) trace metal stoichiometries were also observed (see *Particulate trace metals (pTMs) in the ODZ* section).

An especially clear indicator of the oxygen sensitivity of the deeper ODZ particle assemblages occurred at Sta. 8 where two maxima in bulk particles (transmissometry) were observed (Fig. 5D). A steep drop in particles at \approx 50 m into the ODZ was associated with the appearance of a local oxygen maximum there, a phenomenon also occasionally observed in drifter transects of the region (Whitmire et al. 2009). Particle load increases again below this oxygen signal, once oxygen has dropped to negligible values. Despite the capability for many upper ODZ organisms to rapidly switch between aerobic and anaerobic metabolisms (Tiano et al. 2014), disruptions to the anoxic environment deeper in the ODZ appear to be associated with strong, vertically rapid decreases in the local particle load. We take this result to indicate the strong oxygen sensitivity, and likely predominantly biotic nature, of much of the ODZ particle regime.

Correction for lithogenic and ferruginous phases

To isolate the elemental composition of biogenic phases from other particle types, we performed sequential corrections for lithogenic (crustally sourced) and ferruginous (in situ produced Fe–rich, e.g., Fe oxyhydroxide) phases.



Fig. 5. Overlain profiles at Sta. 4 through 13 (colors, annotations) for (**A**) Chl-*a* fluorescence, (**B**) particulate P (after correction for lithogenic and Feoxides-see below), and (**C**) bulk particle concentration (transmissometer voltage), within the upper portion of the ODZ. Depth scale is relative to the upper boundary of the ODZ (dashed horizontal line). ODZ-associated secondary fluorescence maxima are generally confined to the upper 40 m of the ODZ and are absent at Sta. 11–13. ODZ P-maxima and bulk particle maxima (transmissometry) exhibit deeper penetration into the ODZ than fluorescence, especially at stations nearer shore. Profiles are truncated where oxygen rises above 40% oxygen saturation, to provide vertical context above the ODZ. (**D**) At Sta. 8 an intervening local oxygen maximum (gray shaded area) separates a shallower, intense particle maximum from a deeper, less intense particle maximum. A steep drop in particle abundance is seen at the upper edge of the oxygen maximum, while below, particles only increase again once oxygen has dropped.

Lithogenic correction

As lithogenic tracers, both particulate Al and Ti have merits (Ohnemus and Lam, 2015), with Al more consistent in abundance across crustal sources (typically $\approx 8\%$ by weight) but having demonstrable in situ scavenging, while Ti is more variable across crustal sources but has lower scavenging potential. We opted to use Ti as a lithogenic tracer because we were able to estimate the composition of local lithogenic material using eastern margin (Sta. 1–9), near-bottom benthic nepheloid layer (BNL) samples. These samples showed the presence of resuspended BNLs via transmissometry and Ti and Al concentrations that were well correlated with many pTMs. The element:Ti lithogenic ratios in BNL particles are compared to prior estimates for upper and bulk continental crust in Table 2. Some bioactive metals with low crustal abundances exhibited poor correlations with Al and Ti. For these elements, except Mn, the larger of either the bulk or upper continental crustal ratio to Ti was used to provide a conservative (upper-bound) estimate of lithogenic contributions. For Mn, we used the smaller, upper continental crust value to prevent over-correction of the authigenic Mn-oxyhydroxide carrier phase. This decision proved rather insignificant, however, as particulate Mn is already very low within the ODZ, even prior to lithogenic correction (Fig. 3, Mn).

The transect region is notable for its extremely low aeolian dust inputs (Mahowald et al. 2008; Wagener et al. 2008); resultant lithogenic corrections were generally less than 8% for most bioactive TMs in mixed layer samples (Table 2). In the case of Fe, however, mixed layer abundances were low and variable enough to be statistically indistinguishable from lithogenic ratios to Ti, except in the mixed layers of shelf Sta. 2-5 that retained 24-60% of their Fe after lithogenic correction. In the ODZ, lithogenic corrections for most bioactive elements were minimal, suggesting predominantly non-lithogenic particulate carrier phases there. A non-negligible portion $(24 \pm 20\%)$ of the particulate Fe load within the ODZ, however, was found to be associated with lithogenic material. One possibility is that despite low regional aeolian lithogenic inputs, long residence times for fine lithogenic particles settling into the upper ODZ may allow for some refractory particle accumulation there (Ohnemus and Lam, 2015).

Ferruginous correction

Corrections for ferruginous phases in the ODZ are less constrained than for lithogenic phases, as less is known about the scavenging of a broad suite of trace metals in seawater by authigenically precipitating Fe phases. To address this, we examined two types of Fe-rich particulate samples: those found in association with the shelf ODZ stations (Fig. 3F, particulate Fe:Ti) and in the uniquely Fe-rich East Pacific Rise hydrothermal plume, where hydrothermal fluids rich in Fe²⁺ are rapidly oxidized and precipitate as Fe oxyhydroxides upon mixing with oxic seawater. These hydrothermal particles, along with a small fraction of dissolved Fe, remain suspended during a decades-long transit thousands of kilometers to the west (Resing et al. 2015). At the sampling point nearest to the ridge axis, Sta. 18, the suspended plume was concentrated (lower and upper particulate Fe quartiles for n = 10 plume samples were 48 nM and 79 nM, respectively). Fe speciation in these samples was dominated by Fe(III) oxyhydroxides, while Fe sulfides were undetectable (Toner et al. 2014). In Fig. 6, we compare the particulate TM:Fe ratios in hydrothermal plume samples to those in Ferich shelf ODZ samples, the latter of which are best exemplified by the benthic nepheloid layer sample at Sta. 2 which had 130 nM Fe. Ultimately, we use the particulate TM:Fe signature of hydrothermal plume particles to correct for the ferruginous end-member for the following reasons.

An important consideration is the potential for authigenic Fe phases to scavenge dissolved phosphate from seawater, potentially leading to erroneous interpretations of the presence of biomass. Importantly, however, we note that the scavenging of P by hydrothermal Fe oxyhydroxides yields only ≈ 0.2 mol P per mole Fe precipitated, consistent with previous measurements at the EPR (0.17 mol:mol P/Fe) (Feely et al. 1998) and estimates of an upper limit of 0.5 mol P per mol Fe during Fe²⁺ precipitation in seawater (Gunnars and Blomqvist 1997; Gunnars et al. 2002; Senn et al. 2015). In the ODZ, ratios of P/Fe in particles were much higher median 9.3 mol P/mol Fe, ranging from 2.3 to 140 mol P/mol Fe. This suggests the dominance of biogenic phases, not ferruginous phases, in carrying particulate P, even in shelf ODZ particles where the Fe content of particles was high.

Vanadium has an oxyanion–dominated speciation in seawater with a known affinity for positively charged Fe oxyhydroxide surfaces (Trefry and Metz 1989). Indicative of its potential to demonstrate Fe oxyhydroxide scavenging, ratios of V to Fe in Fe–rich shelf ODZ particles were similar to those in hydrothermal plume particles (Fig. 6, V), as were ratios of Cu/Fe (Fig. 6, Cu), La/Fe, and Y/Fe (not shown). This suggests Fe-oxyhydroxide–dominated carrier phases for these elements in Fe–rich shelf ODZ samples, though corrections for these elements in the majority of ODZ samples, where Fe contents are lower, nevertheless remain rather moderate (medians corrections; Table 2).

Ratios of other pTMs to Fe in shelf ODZ samples were much higher than in plume particles (e.g., Fig. 6, Cd, P). This is potentially explained by the substantial concentrations of biomass (P) over the shelf, with more than 1000 nM particulate P (vs. 130 nM Fe) in the Sta. 2 BNL sample alone. The benthic/shelf sources of particulate Fe to shelf ODZ stations also apparently supply significant concentrations of biomass or biodetritus, which could be ultimately sourced from the overlying water column and/or the sediments. These inputs may also have undergone multiple sedimentation/resuspension cycles, further altering their composition. Given that the composition of this organic matter remains unknown, and that particulate P (biomass), not Fe, may be the dominant carrier phase for many pTMs, we based our ferruginous correction on the abiotically dominated hydrothermal plume particles. In doing so, we assume that all non-lithogenic Fe within the ODZ is composed of this ferruginous end-member, and correct the other pTMs using the median Sta. 18 pTM:Fe ratios (Ferruginous end-member; Table 2).

For most bioactive trace metals and P, ferruginous corrections based on hydrothermal plume pTM:Fe ratios were small in the ODZ (typically less than 10%; Table 2), with higher median corrections for V, Pb, and Y. Already vanishingly low Mn concentrations in the ODZ can be completely accounted for by ferruginous phases. Elements corrected for both lithogenic and ferruginous phases (Cd, Co, Cu, Ni, V, and Zn) are referred to with the suffix "_NoLithOrFe," while



Fig. 6. Scatterplots of select bioactive pTMs (Cd, P, Cu, and V; *y*-axes) vs. particulate Fe (*x*-axes) after correction for only lithogenic abundances ("_NoLith") in shelf ODZ samples (gray asterisks) and samples from the East Pacific Rise hydrothermal plume (Sta. 18–26, 2000–2800 m depth, pFe > 1 nM; black diamonds). Robust linear regressions (MATLAB, which iteratively re-weights samples to remove the influence of outliers [Holland and Welsch 1987]) and r^2 values for the fits are shown (color coded lines, text). Insets: close-up views of the lower ends of the distributions.

elements corrected only for lithogenic phases (Fe and Mn, as needed) are referred to as "_NoLith."

Particulate trace metals (pTMs) in the ODZ

Local maxima in many bioactive pTMs (Cd, Co, Cu, Ni, V, and Zn) were coincident with ODZ P/biomass enrichments, suggesting elevated biotic accumulation of trace metals by ODZ organisms. These enrichments remain even after correction for lithogenic and ferruginous phases (Fig. 7). Median stoichiometries for the heterotrophic, non-fluorescent ODZ biomass sampled at Sta. 11–13 (solid lines, Fig. 7) are between 2.8 and 8.6 times higher than those of transectmedian mixed layer biomass (dashed lines, Fig. 7), with these two end-members constraining most particulate samples.

Our ferruginous correction using Fe–rich hydrothermal scavenging of particulate TMs is a best effort (high–end) constraint in that we have assumed that all Fe within the ODZ is ferruginous in nature. We can, however, examine the Fe abundances in ODZ particles corrected only for lithogenic



Fig. 7. Scatterplots of bioactive pTMs (*y*-axes) vs. P (*x*-axes) for all stations \geq 6 after correction for both lithogenic and ferruginous phases ("_NoLithOrFe"), except Fe and Mn which have been corrected for only lithogenic phases ("_NoLith") (see next page). Shading and shapes are as in Fig. 4: white circles: oxic euphotic zone (oxygen saturation \geq 90%); grey squares: upper oxycline; black diamonds: ODZ. Trendlines and annotations (slopes) show the transect-median mixed layer stoichiometry (dashed line) and median ODZ stoichiometry observed at Sta. 11–13 (solid line), which only exhibited heterotrophic biomass.

material relative to P (Fig. 7, Fe:P) to look for possible trends in Fe usage and/or scavenging. Doing so, we find a Fe:P ratio of 39.7 mmol:mol in the heterotrophic ODZ samples of Sta. 11–13 (solid line, Fig. 7, Fe:P), while stations nearer shore exhibited much higher Fe:P ratios (points above the line, Fig. 7, Fe:P) demonstrating greater loading of ferruginous particles in the eastern ODZ. It remains to be demonstrated how (or if) this higher ferruginous content is associated with the autotrophy present in eastern ODZs. One possible link is through the stronger secondary nitrite maxima at those stations: metabolic Fe(II) oxidation, and, in some species, mineral encrustation of cells, have been observed in mixotrophic denitrifying bacteria (Klueglein et al. 2014) that may be present in such communities. Relatedly, though the ferruginous phase can account for all Mn in the ODZ (Table 2), so too can the biotic phase (Fig. 7, Mn). Prior to ferruginous correction of Mn, nearly all ODZ biomass exhibits a stoichiometry of \sim 1 mmol Mn/mol P, similar to the Mn content exhibited by surface mixed layer biota (dashed line, Fig. 7, Mn). It thus appears possible that all of the picomolar–level Mn found in the ETSP ODZ may be in association with biotic material, rather than with ferruginous phases.

Figure 8 presents the stoichiometries in upper ODZ samples from each station after correction for lithogenic and ferruginous content. Stoichiometric trends in the surface mixed layer and non–ODZ oxygen minima (west of Sta. 13) are also provided for context. Given the absence of fluorescence at



Fig. 7. (continued)

Sta. 11–13, and the elevated metal:P ratios seen in many samples there, we further delineate the offshore ODZ stations into those exhibiting fluorescence ("autotrophic ODZ", Sta. 6–10) and those without ("heterotrophic ODZ", Sta. 11–13). P–normalized stoichiometries are summarized by region in Table 3, which provides medians and standard deviations for mixed layer (ML) and ODZ samples, as well as ODZ/ML ratios.

The Cd:P and Co:P ratios of heterotrophic ODZ biomass in the upper ODZ at Sta. 11–13 are notably elevated compared to stations with shallower, autotrophic ODZ biomass to the east. At the autotrophic ODZ stations, the highest Cd and Co quotas are found in fluorescent, upper ODZ samples which also roughly follow surface mixed layer trends, especially for Co, perhaps reflecting the presence of *Prochlorococcus* cells in both the upper ODZ and the mixed layer. Non– fluorescent, deeper ODZ samples, which are likely heterotroph-dominated, are generally lower in Cd and Co. Elevated Cd and Co quotas at Sta. 11–13 may be somehow linked to the absence of autotrophic biomass there, to their location at the top of the ODZ, or perhaps associated with station or depth-specific organism(s) not discernable in our dataset.

The other metals (Cu, Ni, V, and Zn) show more consistent trends across heterotrophic and autotrophic ODZ biomass and are also more consistently distinct in composition from mixed layer biota. Copper stoichiometries appear somewhat enriched in deeper, non-fluorescent ODZ samples, with Cu:P typically reaching a maximum between 30 m and 60 m into the ODZ (annotations, Fig. 8, Cu:P), and are also somewhat higher at Sta. 11–13. This may reflect elevated Cu demand, a shift in Cu speciation/bioavailability, or Cu accumulation by heterotrophic organisms living in the ODZ core compared to the autotrophic-dominated upper ODZ.



Fig. 8. Element:P ratios in upper ODZ samples (individual points, mmol:mol P) compared to surface mixed layer samples (red line) and gyre Sta. 14–20 oxygen minimum samples (black line; 250–400 m depth). All ratios plotted have been corrected for lithogenic and ferruginous phases. Error bars for surface mixed layer and gyre oxygen minima represent one standard deviation from the station median. Text annotations indicate sample depth in meters below the upper limit of the ODZ. Symbol shading indicates fluorescence intensity: ODZ samples with fluorescence are shaded green and samples without fluorescence are blue. From right to left, vertical lines separate the shelf ODZ stations (<6), fluorescent/autotrophic ODZ stations (6–10), non–fluorescent/heterotrophic ODZ stations (11–13) and gyre stations (>13).

Interestingly, recent meta–omic results predict the opposite effect, with greater Cu demand expected for upper ODZ species and greater Fe demand in the ODZ core (Glass et al. 2015).

ODZ Ni quotas are also rather consistent across the transect and with depth in the ODZ, at between 2 and 7 mmol:mol, and are enriched compared to the mixed layer (generally < 2 mmol:mol). Saito et al. (2014) attributed significant Ni utilization in this region to *Prochlorococcus* nickel superoxide dismutase (Ni–SOD), which may contribute to the constancy in autotrophic ODZ Ni stoichiometries. SOD has been shown to be constitutively expressed in some anaerobic marine organisms (Dolla et al. 2006) and may be necessary for dealing with the oxidative stress induced by free metals.

A link between decreased Zn availability and Cd substitution in carbonic anhydrase is known in phytoplankton (Cullen and Sherrell 2005) and this effect appears in our dataset as an anti–correlation between mixed layer Zn and Cd quotas (Fig. 8, Cd and Zn, red lines). The same effect does not appear obvious for the ODZ prokaryotic communities, however: the most elevated ODZ Cd quotas, seen at Sta. 11–13, do not appear to be associated with abnormally low Zn quotas, indicating a decoupling between Cd and Zn demand in the subsurface. Similar to Ni, ODZ Zn stoichiometries are consistently enriched compared to the oxic euphotic zone, suggesting that prokaryotic communities maintain higher standing stoichiometries of this metal.

Across all elements, we note that ODZ stoichiometries in the heterotrophic ODZ of Sta. 11–13 are similar to the bulk particle stoichiometries observed at the oxygen minima of non–ODZ gyre stations to the west (black lines in Fig. 8), which lack the distinct local maxima in P and pTMs seen in the ODZ. We have isolated the local maxima in P and pTMs for the heterotrophic biomass at Sta. 11–13 by subtracting linear profile backgrounds. This heterotrophic–only prokaryotic signature (Table 3, "Bkgr. Corr" values) is similar to (Co and Ni) or greater than (Cd, Cu, and V) the typical ranges for gyre oxygen minimum particles seen in Fig. 8, and are slightly lower for Zn. Whether any of the P–normalized stoichiometries we report for ODZ prokaryotic communities are descriptive of metal demands by more globally distributed oxygen minimum prokaryotic communities, however, will require more direct investigation.

Mn oxyhydroxides and acid-volatile sulfides

Authigenic Mn oxides and comparisons to the Black Sea

Particulate Mn exhibits strong depletion within the ETSP ODZ, indicating reductive dissolution of upper oxyclinesourced Mn oxyhydroxides as these particles settle into the suboxic zone (Figs. 3C,D, 7, Mn:P). Some low picomolarlevel particulate Mn may be retained internally by ODZ biota in stoichiometric trends similar to those observed in surface biomass (Fig. 7, Mn), or possibly in association with ferruginous phases. The apparent lack of exposed Mn oxyhydroxides in a way simplifies the ODZ particulate regime: we can ignore authigenic Mn, which is known to scavenge Co and Pb (Sherrell and Boyle 1992; Noble et al. 2015). The absence of Mn oxyhydroxides in the ODZ is also in stark contrast to the Mn-rich suboxic/ferruginous zones of the Black Sea and Cariaco Basin systems, where Mn oxyhydroxides and Fe oxyhydroxides coexist at nM levels (Lewis and Landing 1991, 1992; Yiğiterhan et al. 2011). In the Black Sea suboxic zone these high levels of authigenic carrier phases (24-138 nM particulate Mn, 7-27 nM particulate Fe [Yiğiterhan et al. 2011]) likely overwhelm any biological signals, and the presence of a thick euxinic layer beneath the Black Sea suboxic/ferruginous zone further complicates the dissolved and particulate trace metal fluxes there. This makes direct compositional comparisons between the systems difficult and highlights the potentially unique nature of the ETSP with respect to biogenic metal accumulation and speciation.

The near-complete ODZ-associated loss of Mn also provides a sharp boundary in basin-scale pTM distributions: west of 100°W, Mn oxides produced in the upper oxycline persist in the upper water column as there is no suboxic layer to remove them as they settle. East of 100°W, upper oxycline-sourced Mn oxides are reduced in the ODZ and can only re-form in the lower oxycline beneath the ODZ (and then, only at stations deep enough to have such a feature). This may affect pTM scavenging in the region, especially for the micronutrient Co (Hawco et al. 2016), which is scavenged by Mn. The eastward vertical compression of the upper oxycline may also affect the intensity of Mn (and thus Co) removal as the total volume available for Mn scavenging decreases. Scavenging by Mn may also differ between the upper vs. lower oxyclines, perhaps due to the presence of different dissolved TM pools in each region. Shoaling of the ODZ over time, as has already been observed in response to

climate change (Stramma et al. 2012), may further influence these effects.

Acid-volatile sulfides

We also examined the concentrations of acid-volatile sulfides (AVS), which were analyzed at sea on freshly filtered samples collected by in situ pumps on separate casts at many stations. The AVS method quantifies the sum of many mono-sulfide phases (FeS, ZnS, NiS, and CdS) but not CuS, HgS, or mixed-valence sulfides such as pyrite. AVS concentrations are plotted vs. oxygen saturation in Fig. 9a. In the upper 1000 m of the transect, AVS concentrations ranged between 0.5 pM and 264 pM (median: 3.6 pM; mean \pm 1 SD: 25 pM \pm 50 pM) which is stoichiometrically low in abundance compared to most sulfide-forming pTMs (Fig. 9B). Particulate Ni and Zn were typically in great stoichiometric excess to AVS in the euphotic zone and the ODZ, where stoichiometric ratios of \leq 1:1 would be expected if sulfides were the only, or dominant, particulate carrier for these elements. For visual clarity, and due to the ferruginous phases known to be present at most ODZ stations, Fe:AVS ratios (median: 74 mol:mol, range: 2.2-1800 mol:mol) are not plotted in Fig. 9B.

Cadmium: AVS abundances were well above 1:1 mol:mol in the oxic euphotic zone but generally $\leq 1:1$ in the ODZ (Fig. 9C), implying the potential for a sulfide carrier phase there. Recent work (Janssen et al. 2014) has indicated a potential sulfide carrier for particulate Cd enrichments observed in Atlantic basin oxygen minima where they are thought to form in the low-oxygen microenvironments of larger sinking particles, even when ambient water column oxygen concentrations are $> 50 \mu$ M. These sulfides may then persist in the water column as fine Cd-rich particles once the larger sinking particles have remineralized. In this transect, we expected to see evidence for these processes in the ODZ particle environment, where active S cycling is thought to occur (Ulloa et al. 2012). Given the low (generally < 100 pM) concentrations of AVS in the ODZ, it appears that typically half of AVS present, and frequently all of it, would need to be bound to Cd to account for both phases' abundances (mean ODZ Cd:AVS ratio: 0.43 ± 0.27 mol:mol), at the near total exclusion of the other AVS-forming elements (Ni, Zn, and Fe) that exhibit great stoichiometric excesses to Cd and AVS at the same depths. Although stoichiometrically feasible, the \leq 1:1 relationship observed for Cd:AVS in ODZ particles is alone not convincing evidence for a dominant sulfide carrier for this single element, at least in the suboxic environment of the ETSP.

Abundances of AVS in the ODZ do appear to show a relationship with particulate P, however, exhibiting ratios of ≈ 1 mmol AVS:mol P throughout the ODZ (Fig. 9D). This indicates a potential biotic carrier phase for much of the observed AVS in the ODZ, as also indicated previously for



Fig. 9. (A) Concentrations of AVS (pM, log scale) plotted as a function of oxygen saturation (%). Surface AVS concentrations are indicative of mixed layer AVS sources associated with assimilatory sulfate reduction during photosynthesis. Elevated concentrations near the ODZ, observed through Sta. 13, may also indicate AVS accumulation due to dissimilatory sulfate reduction (the "cryptic sulfur cycle"), though autotrophic processes also occur at many upper ODZ depths. (**B**, **C**) Element–specific metal:AVS ratios (mol:mol, log scale; after litho- and ferruginous correction) as a function of oxygen saturation (%, log scale). Black squares, Cd:AVS; gray circles, Ni:AVS; white diamonds, Zn:AVS. Dashed line: 1:1 expected stoichiometric ratio for metal:sulfide phases. Nearly all euphotic and ODZ samples have Ni:S, Zn:S, and Fe:S (not shown) ratios greater than 1:1, implicating dominance of non–sulfidic particulate carrier phases for those elements. Cd:AVS ratios are consistently greater than 1:1 in the euphotic zone, but generally $\leq 1:1$ within the ODZ (C, dotted reference line). (**D**) In most ODZ samples, AVS concentrations exhibit a fairly consistent particulate AVS/P ratio (≈ 1 mmol:mol) indicating a predominantly biotic carrier. Elevated AVS:P samples occasionally seen in the upper ODZ at several stations may indicate transient free sulfides, though AVS concentrations never exceeded 265 pM.

many pTMs. Occasionally elevated "flier" AVS concentrations (>100 pM; AVS:P>4 mmol:mol, Fig. 9D), sometimes observed in the upper 50 m of the ODZ, may indeed be local free sulfide maxima related to S–cycling processes that are difficult to reliably sample with discrete bottles. Taken as a whole, however, the limited carrying capacity of observed AVS phases, from a broad sampling of ODZ depths across the transect, suggests that sulfides are minor and/or transient contributors to bulk pTM carrying capacity compared to biotic (P–rich) particles.

Community complexity and dataset limitations

Most prokaryotic ODZ organisms are known via their genetic fingerprints (Ulloa et al. 2012), and few have been assessed with regards to their trace metal associations, either in field studies or trace metal–controlled laboratory cultures. Our observations of consistent P–maxima, pTM–maxima, and elevated pTM quotas throughout the ETSP ODZ provides a first look at the bulk trace metal associations of these under–sampled ecosystems. Our measurements reveal consistent patterns of elevated trace metal associations by biogeochemically important, prokaryote–rich communities. At many stations, ODZ communities expressed elevated stoichiometries for the bioactive trace metals Cd, Co, Cu, Ni, V, and Zn compared to euphotic zone eukaryotic phytoplankton (Twining and Baines 2013).

Although we conclude that many pTM enrichments found in the ODZ are associated with microbial biomass, these communities are also known to be taxonomically and biochemically complex. ODZ ecosystems conduct vertically compressed, if not overlapping, redox cycling of O, N (nitrate/nitrite/ammonium), Mn, and "cryptic" S (Canfield et al. 2010), among other potential redox acceptors. Eastern stations display significant autotrophic fluorescence at upper ODZ depths, while deeper stations to the west exhibit biomass maxima without autotrophy, suggesting heterotrophic communities only. Our bulk digestions of filtered particles from these complex communities are thus limited to interpretation at the broad ecosystem level, rather than directly informing about metal utilization by specific organisms.

Remaining questions

Several questions naturally emerge from our observations of elevated prokaryotic pTM observations within ODZs. Are these elevated particulate TM:P ratios unique to ODZ prokaryotes, and do they occur because of access to generally larger inventories of dissolved TMs in the subsurface? Denitrifying and other N-cycling ODZ biota are expected to have unique biochemical/enzymatic demands for trace metals (Moffett et al. 2007; Glass et al. 2015), so access to stable, elevated dissolved metal pools may support niche anaerobic respirations. In essence, the unique biogeochemical environment of the ODZ may have produced unique biotic stoichiometries, perhaps even reflective of the ancient anoxic ocean (Saito et al. 2003). An additional, non-mutually exclusive possibility is that luxury uptake of micronutrient metals is more general beneath the euphotic zone, as has been shown for Cd and Zn in the Ross Sea (Saito et al. 2010). The metal requirements and stoichiometric flexibilities of prokaryotes must be more rigorously examined to determine if elevated pTM associations are unique to the ODZ and which metal enrichments are associated with which organisms. If elevated pTM stoichiometries are common to prokaryotes found more widely throughout the water column (e.g., Fig. 8; gyre oxymin stoichiometries), their stoichiometries would provide a starkly different biomass compositional end-member than those of phytoplankton that are widely used in modeling experiments. Even if pTM enrichments are limited to ODZ biomass and are not common outside the ODZ, settling and remineralization of ODZ prokaryotic biomass into the deep ocean and underlying sediments may need to be considered separately when parameterizing local metal fluxes.

Future work should help constrain corrections for scavenged, authigenic, ferruginous minerals such as ferrihydrite and lepidocrocite (Senn et al. 2015) which have higher concentrations and greater influence on particle speciation nearer the shelf. Although biomass concentrations (via P) exceeded those of ferruginous phases throughout the ODZ, ferruginous pTM scavenging remains a complicating factor in understanding this water column particle regime. Nevertheless, separation of the various particulate phases as attempted here is perhaps simpler in the ETSP than in the authigenic Mn–, Fe–, and sulfide–rich suboxic zones of the Black Sea and Cariaco Basin.

How do elevated-pTM associations in ODZ biomass relate to local and global cycling of pTMs throughout the oceans? At shallower, margin-associated sites ODZs directly interact with sediments, so deposition and preservation of pTM-rich prokaryotic biomass could influence the sedimentary record of these metals. We observed that metal:P ratios of ODZ communities are typically elevated, sometimes many fold, vs. mixed layer communities (Table 3), with metal stoichiometries highest at the heterotrophic-dominated ODZs of Sta. 11–13 (Table 3, "Bkgr. Corr." stoichiometries). Though large, sinking particle production is presumably diminished within the ODZ due to the exclusion of grazers, persistent settling fluxes of fine, metal-rich biomass (or their adherence to particles sinking through the ODZ) into the sediments may influence the sedimentary record of sites overlain by ODZs. Oxic autotrophic biomass production and vertical fluxes presumably dominate prokaryotic biomass signals in these highly productive regions, but the presence of prokaryotic end-members with elevated metal stoichiometries may still need to be considered during sedimentary interpretations of some bioactive metals, e.g., Ni (Böning et al. 2015).

The longitudinal extent of the ODZ and its microbial communities is ultimately dependent on the intensity of regional upwelling. In this transect of the ETSP along 12–13°S, the ODZ and its TM–rich biota extended well beyond the region of shelf–associated productivity (\approx 400 km off-shore)—to as far as 2300 km offshore. Redfield–like metal utilization ratios, long presumed to be set by phytoplankton biomass in the surface ocean, have well recognized links to global metal cycles, deep water mass signatures, and broad oceanic composition (Morel 2008). Though these ODZ ecosystems and stoichiometries may be constrained to a limited oceanic volume, the potential for TM–rich prokaryotes to influence biomass/water-mass relationships on larger spatial scales remains unknown.

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Conflict of Interest

None declared.

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