



Seasonal cycling of zinc and cobalt in the south-eastern Atlantic along the GEOTRACES GA10 section

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Abstract. We report the distributions and stoichiometry of dissolved zinc (dZn) and cobalt (dCo) in sub-tropical and sub-Antarctic waters of the south-eastern Atlantic Ocean during austral spring 2010 and summer 2011/2012. In sub-tropical surface waters, mixed-layer dZn and dCo concentrations during early spring were 1.60 ± 2.58 nM and 30 ± 11 pM, respectively, compared with summer values of 0.14 ± 0.08 nM and 24 ± 6 pM. The elevated spring dZn concentrations resulted from an apparent offshore transport of elevated dZn at depths between 20–55 m, derived from the Agulhas Bank. In contrast, open-ocean sub-Antarctic surface waters displayed largely consistent inter-seasonal mixed-layer dZn and dCo concentrations of 0.10 ± 0.07 nM and 11 ± 5 pM, respectively. Trace metal stoichiometry, calculated from concentration inventories, suggests a greater overall removal for dZn relative to dCo in the upper water column of the south-eastern Atlantic, with inter-seasonally decreasing dZn/dCo inventory ratios of 19–5 and 13–7 mol mol⁻¹ for sub-tropical surface water and sub-Antarctic surface water, respectively. In this paper, we investigate how the seasonal influences of external input and phytoplankton succession may relate to the distribution of dZn and dCo and variation in dZn/dCo stoichiometry across these two distinct ecological regimes in the south-eastern Atlantic.

1 Introduction

The trace metal micronutrients zinc (Zn) and cobalt (Co) play an important role in the productivity of the oceans as key requirements in marine phytoplankton metabolism (Morel, 2008; Twining and Baines, 2013). Zinc is required for the acquisition of inorganic carbon and organic phosphorus via the carbonic anhydrase and alkaline phosphatase metalloenzymes, respectively (Morel et al., 1994; Shaked et al., 2006; Cox and Saito, 2013). The requirement for Co stems from its obligation in the biosynthesis of vitamin B₁₂ (Raux et al., 2000; Rodionov et al., 2003) and, like Zn, its potential roles as a metal cofactor in carbonic anhydrase and alkaline phosphatase (Morel et al., 1994; Jakuba et al., 2008; Saito et al., 2017). Significantly, both dissolved Zn (dZn) and Co (dCo) are often scarce in surface seawater, with mean concentrations that are often similar to or relatively depleted compared with typical cellular requirements of phytoplankton (Moore et al., 2013; Moore, 2016). Hence, dZn and dCo availability have the potential to regulate phytoplankton metabolism and growth rates in some ocean regions (Sunda and Huntsman, 1992; Saito et al., 2002; Franck et al., 2003; Shaked et al., 2006; Bertrand et al., 2007; Jakuba et al., 2012; Mahaffey et al., 2014; Chappell et al., 2016; Browning et al., 2017).

The role for Zn and Co in carbonic anhydrase establishes an interaction between their ocean cycles, whereby biochemical substitutions between the enzyme-bound metals enable a stoichiometric plasticity in their cellular requirements that

can negate the effect of limited availability. For example, a number of eukaryotic algae can substitute Zn for Co as well as cadmium (Cd) in carbonic anhydrase when seawater dZn concentrations are low (Price and Morel, 1990; Sunda and Huntsman, 1995; Lane and Morel, 2000; Xu et al., 2007; Saito and Goepfert, 2008; Kellogg et al., 2020). In contrast, the prokaryotic picocyanobacteria *Synechococcus* and *Prochlorococcus* appear to have an absolute Co requirement (Sunda and Huntsman, 1995; Saito et al., 2002; Hawco and Saito, 2018). The availability and stoichiometry of dZn and dCo may therefore also exert a key control on phytoplankton community structure in some ocean regions (Leblanc et al., 2005; Saito et al., 2010; Chappell et al., 2016).

With the arrival of GEOTRACES research cruises, a number of studies have provided comprehensive data on the basin-scale distributions of Zn and Co in the Atlantic Ocean (e.g. Bown et al., 2011; Noble et al., 2012, 2017; Wyatt et al., 2014; Roshan et al., 2015; Middag et al., 2019). Such efforts have transformed our understanding of the biogeochemical processes associated with Zn and Co cycling (Saito et al., 2017; Vance et al., 2017; Weber et al., 2018; Tagliabue et al., 2018; Roshan et al., 2018), yet there are still geographically important regions of the Atlantic that remain largely understudied, including the south-eastern Atlantic.

The sub-tropical front (STF) of the south-eastern Atlantic represents the convergence of warm, predominately macronutrient-limited sub-tropical surface water (STSW) and cold, iron-limited but macronutrient enriched sub-Antarctic surface water (SASW), creating one of the most dynamic nutrient regimes in the oceans (Ito et al., 2005; Browning et al., 2014; Moore, 2016). Here, the relative supply and availability of macronutrients and iron (Fe) exert an important control in maintaining the elevated phytoplankton stock and productivity that are typical of this frontal region, particularly during austral spring and summer (Moore and Abbott, 2000; Ito et al., 2005; Browning et al., 2014). Dissolved Zn is also depleted in SASW that flows northwards to converge with STSW at the STF (Wyatt et al., 2014). However, the potential role for Zn in the mediation of phytoplankton distribution and community structure in this region is currently unclear.

Using data from two UK-GEOTRACES cruises (transect GA10) this study examines the seasonal availability and ecological stoichiometry of dZn and dCo, by analysis of their relationships with phosphate, in upper ocean waters of the south-eastern Atlantic. These data, together with measurements of phytoplankton pigment biomass and community structure, offer improved knowledge of the seasonal influences of external input and phytoplankton succession on the distribution and cycling of Zn and Co in these dynamic waters.

2 Methods

2.1 Sampling methods

Seawater samples were collected during two UK-GEOTRACES cruises in the South Atlantic Ocean (GA10; Fig. 1). The first cruise (D357) took place during austral spring 2010 (18 October to 22 November 2010), sampling the south-eastern Atlantic on board the RRS *Discovery*. During D357, two transects were completed between Cape Town and the zero meridian that represent early austral spring (D357-1) and late austral spring (D357-2), respectively. The second cruise (JC068) took place during austral summer 2011/2012 (24 December 2011 to 27 January 2012), along the same transect of the first cruise and continuing along 40° S between Cape Town and Montevideo, Uruguay, on board the RRS *James Cook*. For JC068, we present here only the repeat transect data between Cape Town and 13° W that represent the south-eastern Atlantic aspect of this transect. The stations occupied during the three transects were not identical but rather represent a coverage of the Southern Ocean and sub-tropical waters present. Where stations were reoccupied during one or more transects, they have the same station number.

All sampling bottles were cleaned according to the procedures detailed in the GEOTRACES sample handling protocols (Cutter et al., 2010). Seawater and particulate samples below 15 m depth were collected using a titanium-frame CTD with 24 trace metal clean, 10 L, Teflon-coated OTE (ocean test equipment) Niskin bottles deployed on a plasma rope. Sub-samples for dissolved trace metal analysis were filtered through 0.8/0.2 µm cartridge filters (AcroPak™ 500, Pall) into 125 mL low-density polyethylene bottles inside a class 1000 clean-air container. Each sub-sample was acidified to pH 1.7 (0.024 M) by addition of 12 M hydrochloric acid (HCl, UpA, Romil) under a class 100 laminar-flow hood. Vertical sampling for dissolved trace metals was augmented by surface samples collected at each station using a towed “fish” positioned at approximately 3–5 m depth. Fish samples were filtered in-line and acidified as described for samples collected from the titanium sampling system. Particulate samples were collected onto acid-clean 25 mm, 0.45 µm, polyethersulfone membrane disc filters (Supor®, Pall) and stored frozen (−20 °C) until shore-based analysis.

2.2 Trace metal analysis

Dissolved Co was determined in the ISO-accredited clean room facility (ISO 9001) at the University of Plymouth (UK) using flow injection with chemiluminescence detection, modified from the method of Cannizzaro et al. (2000) as described by Shelley et al. (2010). Briefly, dCo was determined in UV-irradiated samples using the reaction between pyrogallol (1,2,3-trihydrobenzene) and hydrogen peroxide formed in the presence of Co. Standards (20–120 pM

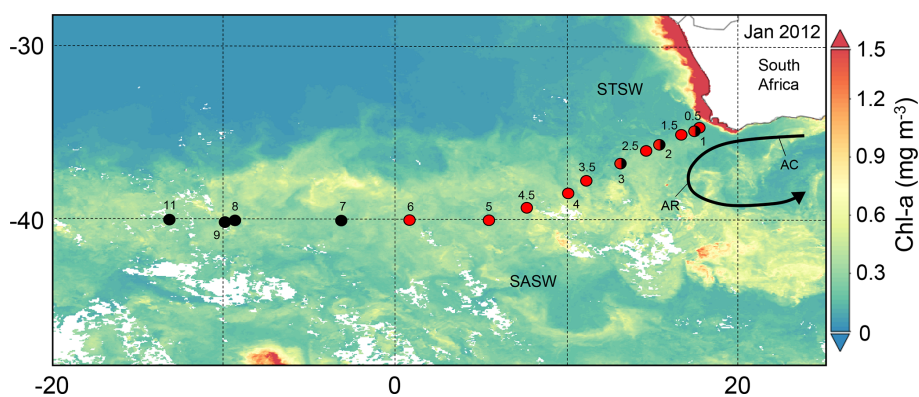


Figure 1. The south-eastern Atlantic stations sampled for dissolved Zn and Co along the GA10 section during UK-GEOTRACES cruises D357 (red circles) and JC068 (black circles), overlying a Visible Infrared Imaging Radiometer Suite (VIIRS) monthly composite image of chlorophyll *a* concentrations for January 2012 (<https://oceancolor.gsfc.nasa.gov/>, last access: 7 July 2021). Two transects were completed during D357 between Cape Town and the zero meridian that represent early austral spring 2010 (D357-1; Stations 1, 2, 3, 4, 5 and 6) and late austral spring 2010 (D357-2; Stations 0.5, 1, 1.5, 2.5, 3.5, 4.5), respectively. JC068 took place during austral summer 2011/2012, and we present here only the repeat transect data between Cape Town and 13° W (Stations 1, 2, 3, 7, 8, 9, 11). STSW: sub-tropical surface water; SASW: sub-Antarctic surface water; AC: Agulhas Current; AR: Agulhas Retroflection.

Co) were prepared in 0.2 μm filtered low-dCo seawater (16.5 ± 5.2 pM, $n = 15$) by serial dilution of a 1000 ppm Co ICP-MS standard (Romil, UK). The accuracy of the analytical method was validated by quantification of dCo in SAFe (S and D2) and GEOTRACES (GD) reference seawater (Table 1). There was no detectable analytical dCo blank, and the limit of detection (3σ of the lowest concentration standard) was 1.98 ± 0.87 pM ($n = 15$).

Dissolved Zn was determined using flow injection coupled with fluorescence detection, modified from the method of Nowicki et al. (1994) and described previously for this GEOTRACES section by Wyatt et al. (2014). The accuracy of the analytical method was validated by quantification of dZn in SAFe (S and D2) reference seawater (Table 1). The blank for dZn determination was 0.14 ± 0.13 nM, and the limit of detection (3σ of the lowest concentration standard) was 0.01 ± 0.01 nM ($n = 15$).

Measurement uncertainties were estimated after the Nordtest approach (Worsfold et al., 2019), where a combined uncertainty (u_c) is computed from day-to-day within-lab reproducibility and uncertainties associated with the determination of reference materials (Table 1). This approach creates higher uncertainties than those previously published for dZn and dCo analyses but provides a more realistic estimation of analytical uncertainty. During this study, the u_c for dZn and dCo analysis was 22 % and 19 %, respectively, similar to the 13 %–25 % reported by Rapp et al. (2017) for the determination of trace metals, including dZn and dCo, by online pre-concentration and high-resolution sector field ICP-MS detection. The elevated u_c within our data results from the greater uncertainty surrounding the very low dZn and dCo concentrations in the SAFe reference sample (Table 1), whereas the dZn and dCo u_c using only the Safe D2

are < 5 %. Hereafter, when presenting low dZn and dCo concentrations for comparison with phytoplankton biological requirements (Sect. 3.5), we apply a fixed u_c of 20 % to our data.

Total particulate trace metals (i.e. pZn, pCo, pTi) were determined using inductively coupled plasma mass spectrometry (Thermo Fisher XSeries-2) following a sequential acid digestion modified from Ohnemus et al. (2014). Potential interferences (e.g. $^{40}\text{Ar}^{16}\text{O}$ on ^{56}Fe) were minimized through the use of a collision and reaction cell utilizing 7 % H in He and evaluation of efficiency and accuracy assessed using certified reference material (CRM). Full details of the method and CRM results can be found in Milne et al. (2017).

2.3 Nutrients, phytoplankton, temperature and salinity

The dissolved macronutrients phosphate (PO_4^{3-}), silicic acid ($\text{Si}(\text{OH})_4$ but referred to as Si hereafter) and nitrate (determined as nitrate + nitrite, NO_3^-) were determined in all samples for which trace metals were determined, in addition to samples collected from a stainless-steel rosette. Macronutrients were determined using an AA III segmented-flow AutoAnalyzer (Bran and Luebbe) following colorimetric procedures (Woodward and Rees, 2001). Salinity, temperature and depth were measured using a CTD system (Seabird 911+), whilst dissolved O_2 was determined using a Seabird SBE 43 O_2 sensor. Salinity was calibrated on board using discrete samples taken from the OTE bottles and an Autosal 8400B salinometer (Guildline), whilst dissolved O_2 was calibrated using a photometric automated Winkler titration system (Carritt and Carpenter, 1966). Mixed-layer depths (MLDs) were calculated using the threshold method of de Boyer Montégut et al. (2004), where MLD is identified from a linear interpolation between near-surface density and

Table 1. Analytical validation results for open-ocean surface seawater (SAFe S), 1000 m seawater (SAFe D2) and 2000 m seawater (GEOTRACES GD). All concentrations are in nanomolar (± 1 std. dev.). Consensus value conversion = 1.025 kg L^{-1} . FIA: flow injection analysis. ND indicates sample not determined.

| | SAFe S | SAFe D2 | GEOTRACES GD |
|--------------------|-----------------------|------------------------|-----------------------|
| Zn (FIA) | 0.060 (0.020) $n = 7$ | 7.723 (0.091) $n = 12$ | ND |
| Zn consensus value | 0.071 (0.010) | 7.616 (0.256) | 1.753 (0.123) |
| Co (FIA) | 0.004 (0.001) $n = 3$ | 0.049 (0.001) $n = 2$ | 0.073 (0.004) $n = 5$ |
| Co consensus value | 0.005 (0.001) | 0.047 (0.003) | 0.067 (0.001) |

the depth at which density changes by a threshold value (0.125 kg m^{-3}).

Measurements of phytoplankton pigment biomass and community structure were made on discrete samples collected using a 24-position stainless-steel CTD rosette equipped with 20 L OTE Niskin bottles. For chlorophyll *a* analysis, samples were filtered ($0.7 \mu\text{m}$ Whatman GF/F), and then the filters were extracted overnight in 90 % acetone (Holm-Hansen et al., 1965). The chlorophyll *a* extract was measured on a pre-calibrated (spinach chlorophyll *a* standard, Sigma) Turner Designs Trilogy fluorometer. High-performance liquid chromatography (HPLC) samples ($0.5\text{--}2 \text{ L}$) for accessory pigment analyses were filtered ($0.7 \mu\text{m}$ Whatman GF/F), flash-frozen in liquid nitrogen and stored at -80°C prior to analysis using a Thermo HPLC system. The matrix factorization program CHEMTAX was used to estimate the contribution of taxonomic groups to total chlorophyll *a* (Mackey et al., 1996). Concentrations of nanophytoplankton, *Synechococcus*, *Prochlorococcus* and photosynthetic picoeukaryotes were analysed by analytical flow cytometry (AFC) using a FACSort flow cytometer (Becton Dickenson, Oxford, UK) according to the methods described in Davey et al. (2008) and Zubkov et al. (2003).

3 Results and discussion

3.1 Hydrographic setting and macronutrient distributions

The prominent water masses along the D357 and JC068 transects (Fig. 2) were identified by their characteristic thermohaline and macronutrient properties (Sarmiento et al., 2004; Ansorge et al., 2005; Browning et al., 2014). Wyatt et al. (2014) provide a more detailed description of the JC068 hydrography along the entire GA10 section. Whilst we aim to compare the near-shore versus offshore distributions of micro- and macronutrients, note that sub-Antarctic mode water was not sampled for trace metals during the D357-2 late-spring transect, and therefore only the early-spring and the summer values are discussed for SASW hereafter.

3.1.1 Surface mixed layer

During all three transects the STF was identified by a sharp potential temperature (θ) gradient in the upper 200 m, with the $\theta 15^\circ \text{C}$ isotherm corresponding well to changes in macronutrient concentrations between STSW and SASW. North of the STF, mixed-layer macronutrient concentrations (Table 2) decreased in STSW between the three occupations of the transect. The largest relative depletion observed was for NO_3^- , with a ~ 2.7 -fold reduction in mean inventory concentration from $870\text{--}326 \mu\text{mol m}^{-3}$ between early spring and summer, whilst PO_4^{3-} and Si concentrations were reduced 1.5- and 1.4-fold, respectively. The largest absolute depletion was observed for Si, with a reduction of $848 \mu\text{mol m}^{-3}$ between early spring and summer. Conversely, summer SASW mixed-layer mean concentrations of NO_3^- , PO_4^{3-} and Si were relatively 1.6-, 1.4- and 2.1-fold lower than early spring, respectively, whilst the largest absolute depletion of $1912 \mu\text{mol m}^{-3}$ was observed for NO_3^- . SASW mixed-layer concentrations of NO_3^- and PO_4^{3-} were at least 2.1-fold higher than for STSW during the study, whilst the Si concentration was at least 1.5-fold lower, highlighting the relative deficiencies in major nutrients between high- and low-latitude-derived surface waters (Sarmiento et al., 2004; Moore, 2016).

3.1.2 Sub-surface waters

The Southern-Ocean-derived sub-Antarctic mode water (SAMW) and underlying Antarctic intermediate water (AAIW) were identified using their characteristic core-potential-density ($\sigma_\theta = 26.8 \text{ kg m}^{-3}$) (Sarmiento et al., 2004; Palter et al., 2010) and thermohaline ($S < 34.4$, $\theta > 2.8^\circ \text{C}$) properties (Fig. 2). Wyatt et al. (2014) have identified these water masses along this section between 200 and 500 m. During all three transects, low sub-surface (50–500 m) macronutrient concentrations were observed between 13 and 16°E , associated with salinity maxima. The feature conforms to the mean locality and depth range of Agulhas water (Duncombe Rae, 1991), clearly highlighting the penetration of Indian Ocean water into northward-flowing SAMW.

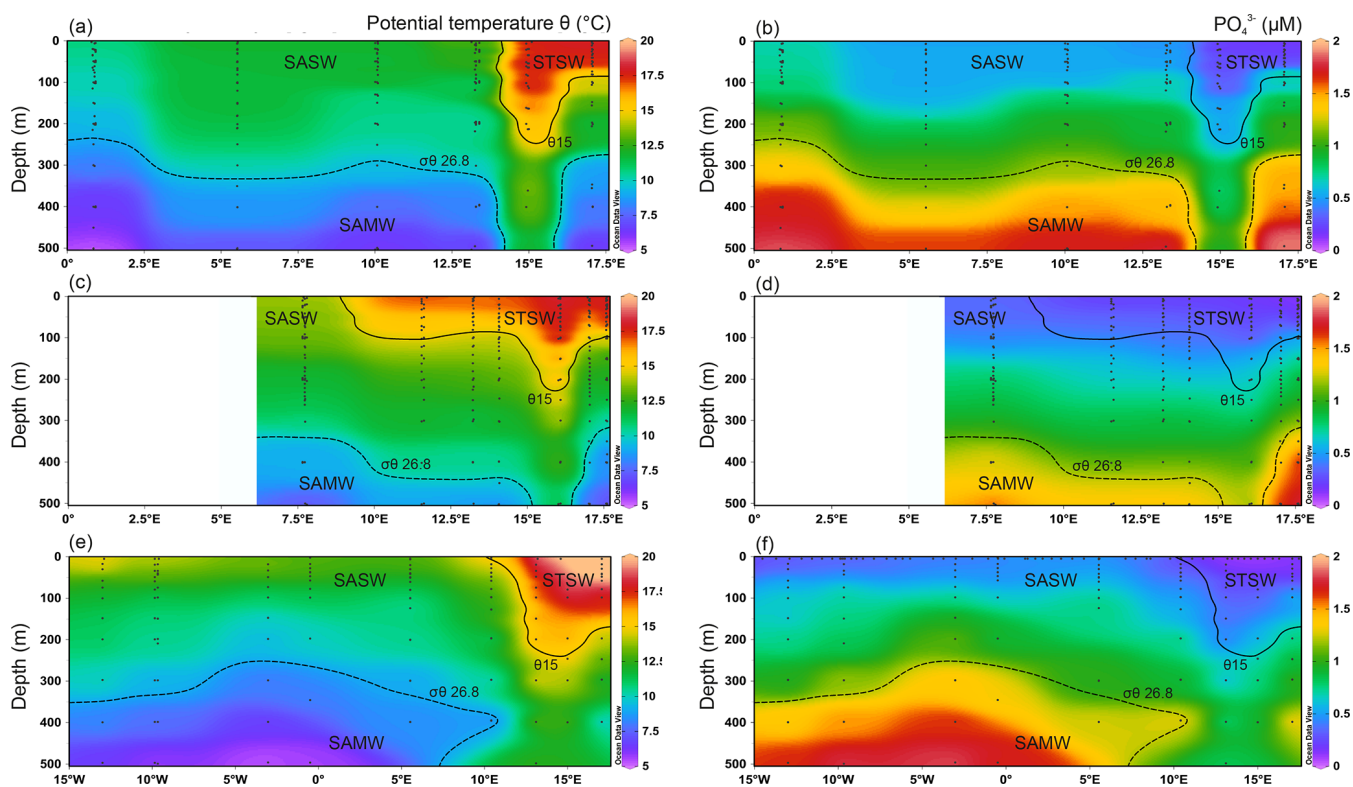


Figure 2. Upper 500 m potential temperature (θ) and dissolved PO_4^{3-} distributions for the south-eastern Atlantic along early-spring (a, b; D357-1), late-spring (c, d; D357-2) and summer (e, f; JC068) transects. The dominant Southern Ocean (SASW and SAMW) and South Atlantic (STSW) water masses that influence the distribution of nutrients are shown. The θ 15 °C isotherm (solid contour) represents a practical definition of the STF location, whilst SAMW is identified by the median-potential-density (σ_θ) isopycnal (26.8 kg m⁻³; dashed contour; see Sect. 4.1.). STSW: sub-tropical surface water; SAMW: sub-Antarctic mode water; AAIW: Antarctic intermediate water.

Table 2. South-eastern Atlantic dissolved micro- and macronutrient mean concentration inventories for the upper water column during early-spring (D357-1), late-spring (D357-2) and summer (JC068) transects. STSW and SASW waters were defined using the θ 15 °C isotherm (Sect. 3.4) and are compared with total inventories calculated for the shallower mixed layer (in parentheses) that include continental inputs of dissolved Zn and Co. Zn / PO_4^{3-} , Co / PO_4^{3-} and Zn / Co represent the concentration inventory ratios for STSW and SASW, respectively. STSW: sub-tropical surface water; SASW: sub-Antarctic surface water.

| Oceanographic regime | Transect | Zn (nmol m ⁻³) | Co (nmol m ⁻³) | NO_3^- (μmol m ⁻³) | PO_4^{3-} (μmol m ⁻³) | Si(OH)_4 (μmol m ⁻³) | Zn / PO_4^{3-} (μmol mol ⁻¹) | Co / PO_4^{3-} (μmol mol ⁻¹) | Zn / Co (mol mol ⁻¹) |
|----------------------|--------------|----------------------------|----------------------------|---|--|---|---|---|----------------------------------|
| STSW | Early spring | 624 (1597) | 32 (30) | 2694 (870) | 333 (203) | 3735 (2790) | 1876 | 97 | 19 |
| | Late spring | 384 (592) | 23 (17) | 1846 (763) | 276 (191) | 2781 (2326) | 1387 | 82 | 17 |
| | Summer | 158 (139) | 29 (24) | 1557 (326) | 226 (139) | 2711 (1942) | 699 | 129 | 5 |
| SASW | Early spring | 182 (112) | 14 (13) | 6035 (5300) | 615 (566) | 1875 (1847) | 296 | 22 | 13 |
| | Summer | 83 (94) | 12 (10) | 4143 (3388) | 439 (400) | 1027 (886) | 188 | 26 | 7 |

3.2 Zn and Co distributions of the south-eastern Atlantic Ocean

3.2.1 Surface mixed layer

Figure 3 shows the dZn and dCo distributions for the upper 500 m of the south-eastern Atlantic for the D357 and JCO68 transects. For full-depth dZn distributions along JC068 refer to Wyatt et al. (2014). In the surface mixed layer, dZn

and dCo concentrations ranged from 0.01–4.57 nM and 1–50 pM, respectively. The large range in dZn concentrations resulted from an apparent offshore transport of elevated dZn within STSW between 20–50 m during early spring (1.48–4.57 nM; Stations 1–2) that was reduced by late spring (0.48–1.76 nM; Stations 0.5–1.5) and was absent during summer (0.01–0.13 nM; Stations 1–2). Similarly, but to a lesser extent, elevated dCo concentrations were observed in STSW between 10 and 50 m during early and late spring (15–50 pM)

compared with summer (18–33 pM). Our findings are consistent with previous observations of elevated dissolved and particulate trace metals over the same depth range in waters close to South Africa, including Co, Fe, Mn and Pb (Chever et al., 2010; Bown et al., 2011; Boye et al., 2012; Paul et al., 2015). We postulate that these trace metal enrichments can arise from atmospheric inputs and/or the lateral advection of metal-enriched waters from the Agulhas Current (AC) and/or South African continental shelf, and we discuss this further in Sect. 3.3. In SASW, mixed-layer dZn and dCo concentrations ranged from 0.01–0.25 nM and 3–18 pM, respectively, during the study, significantly lower than STSW values, with the lowest concentrations observed during the summer transect (Table 2).

3.2.2 Sub-surface waters

During the early-spring D357-1 transect, elevated dZn and dCo concentrations were observed between the surface mixed layer and 500 m (1.48–3.85 nM and 39–62 pM, respectively) at the station closest to the South African continent (Station 1). Here, the highest dZn concentrations were associated with the dZn-enriched waters (20–55 m) described above for the surface mixed layer. During the late-spring D357-2 transect, the near-shore (Stations 0.5–1) dZn concentrations were lower (0.31–1.76 nM), whilst dCo remained similar to early-spring values (27–57 pM). During summer, near-shore (Station 1) sub-surface dZn concentrations were markedly lower (0.03–0.50 nM) than spring values, whilst dCo concentrations (17–52 pM) were only marginally lower. In offshore waters, sub-surface dZn concentrations ranged from 0.01–1.01 nM across all three transects, with extremely low values in the upper 400 m (0.22 ± 0.21 nM) and the highest values between 400 and 500 m. The absence of a significant return path for dZn with SAMW to waters above 400 m at this latitude (Wyatt et al., 2014; Vance et al., 2017) is likely an important control on dZn distributions across all three transects. In contrast, dCo concentrations were depleted in the upper 200 m (1–35 pM) and elevated in SAMW (23–56 pM), suggesting that these Southern-Ocean-derived waters also play an important role in upper-water-column dCo distributions of the South Atlantic.

To assess whether seasonal changes in sub-surface supply could influence dissolved Zn and Co concentrations in the upper water column of the south-eastern Atlantic, we examined the metal versus PO_4^{3-} distributions of underlying SAMW and AAIW. Throughout this paper, metal : PO_4^{3-} is used to indicate an uptake remineralization ratio derived from a regression slope, whilst metal / PO_4^{3-} denotes a concentration ratio. Figure 4 and Supplement Table S1 show how the dZn : PO_4^{3-} regression slope for SAMW and AAIW varied little between the three transects. These slopes are a function of the pre-formed micro- and macronutrient concentrations and the uptake/remineralization ratio of the source waters as well as mixing during advection between the Southern

Ocean and South Atlantic (Vance et al., 2017; Middag et al., 2019). The dZn : PO_4^{3-} slopes steepen with the introduction of AAIW with higher dZn / PO_4^{3-} concentration ratios, yet it is the relatively shallow slopes of overlying SAMW that imply a low and relatively consistent sub-surface supply of dZn to STSW and SASW of the South Atlantic (Wyatt et al., 2014). The shallower waters overlying SAMW clearly show elevated dZn concentration, specifically during the spring transects, compared with what could be delivered if sub-surface supply was the dominant source governing dZn availability in surface waters (Fig. 4). It is therefore unlikely that a change in sub-surface supply from underlying SAMW is responsible for the change in dZn inventories of STSW and SASW between the three transects.

Similarly, the dCo : PO_4^{3-} regression slope varied little between the three transects (Fig. 4 and Supplement Table S1). In dCo : PO_4^{3-} space, a single slope can be fit to SAMW and AAIW with no net scavenging effect on dCo distribution over the upper 1000 m. Like dZn, the waters overlying SAMW displayed spring dCo concentrations elevated above that potentially delivered via the SAMW supply. During summer, however, SAMW may provide a sub-surface source of dCo (Fig. 4c) to overlying waters, highlighting how Southern-Ocean-derived waters may play important, yet different, roles in upper-water-column metal distributions of the south-eastern Atlantic.

3.3 Shelf-derived sources of Zn and Co

Potential sources of trace metals to surface waters of the south-eastern Atlantic include atmospheric inputs from South Africa and Patagonia (Chance et al., 2015; Menzel Barraqueta et al., 2019) as well as interactions with shelf and slope waters of the Agulhas Bank (Bown et al., 2011; Boye et al., 2012; Paul et al., 2015). During the D357 spring transects, elevated mixed-layer dZn and dCo concentrations (up to 4.57 nM and 50 pM, respectively; Sect. 3.2) were observed at stations closest to the Agulhas Bank shelf and slope (Stations 0.5, 1, 1.5 and 2). Here, we compare these metal elevations with respect to the aforementioned sources. Firstly, we encountered only brief, light rain during the study and thus minimal wet deposition of atmospheric aerosol. By combining the median flux of atmospheric dry deposition for soluble Zn and Co for the south-eastern Atlantic (Zn: $6.0 \text{ nmol m}^{-2} \text{ d}^{-1}$; Co: $0.05 \text{ nmol m}^{-2} \text{ d}^{-1}$; Chance et al., 2015) with the mean mixed-layer depth (34 m) for STSW during D357, dust dissolution is estimated to add approximately 5.5 and 0.05 nmol m^{-3} dZn and dCo, respectively, over a 1-month period. These inputs are low compared with the mixed-layer metal inventories, representing $< 1\%$ of dZn and dCo concentration in STSW during the D357 transects (Table 2), and would not be sufficient to generate distinct mixed-layer maxima. It is likely, therefore, that the dZn and dCo elevations originated from the advection of metal-enriched waters from the western Agulhas Bank, a region

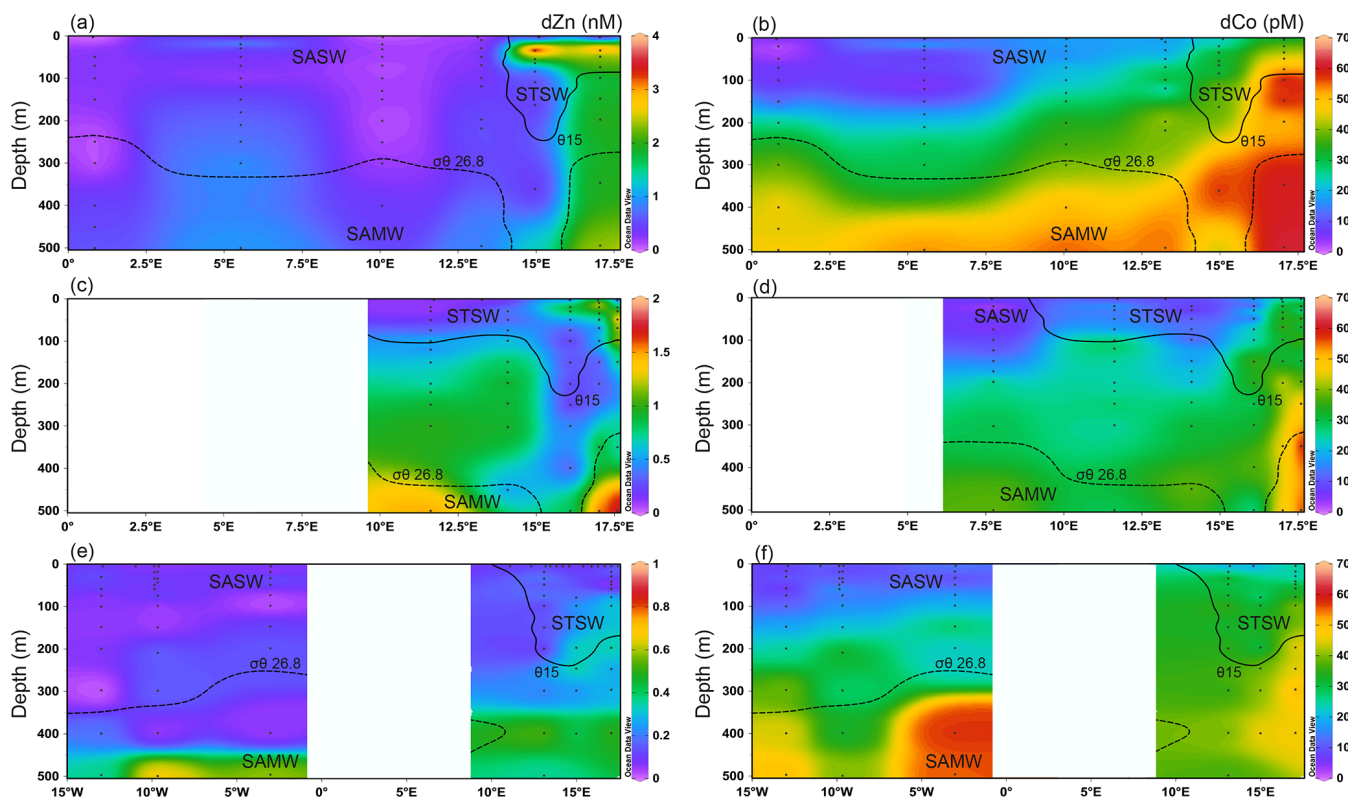


Figure 3. Upper 500 m dissolved Zn and Co distributions for the south-eastern Atlantic along early-spring (a, b; D357-1), late-spring (c, d; D357-2) and summer (e, f; JC068) transects. The STF is delineated by θ 15°C (solid contour), whilst the influence of SAMW is evident by the median-potential-density ($\sigma\theta$) isopycnal (26.8 kg m^{-3} ; dashed contour; see Sect. 4.1.). STSW: sub-tropical surface water; SAMW: sub-Antarctic mode water; AAIW: Antarctic intermediate water. Note the changing y-axis scales for dZn distribution.

identified as a distinct source of both dissolved and particulate trace metals to the south-eastern Atlantic (Chever et al., 2010; Bown et al., 2011; Boye et al., 2012; Paul et al., 2015) and/or from the leakage of Indian Ocean water into the south-eastern Atlantic via the AC.

The detachment of Agulhas rings and filaments from the AC during its retroflection back towards the Indian Ocean constitutes a source of Pb to the surface south-eastern Atlantic along the D357 transects (Paul et al., 2015). Whilst we observed elevated mixed-layer dZn and dCo at $\sim 15^\circ \text{E}$ during both D357 transects, the absence of metal enrichment across the depth of the AC salinity maxima (Figs. 2 and 3) suggests that the signal must be entrained from elsewhere. Furthermore, dZn concentrations from the AC along the east coast of South Africa do not exceed 0.5 nM in the upper 200 m (Gosnell et al., 2012). It is therefore likely that the dZn and dCo enrichment was derived from the Agulhas Bank. The AC has been shown to meander over and interact with the Agulhas Bank, forming eddies and filaments on the shoreward edge of the AC proper that tend to move northwards along the western shelf edge and into the south-eastern Atlantic (Lutjeharms and Cooper, 1996; Lutjeharms, 2007), potentially delivering shelf-derived sedimen-

tary material. We found no evidence of a fluvial signature in our data, and no significant fluvial source for trace elements to our study region has been reported in the literature. We focus here on the more likely scenario of sedimentary inputs as the driver of mixed-layer dZn and dCo elevations at the shelf and slope stations during D357. Despite no available particulate trace metal data for the D357-1 early-spring transect for direct comparison with the highest dZn and dCo elevations, we observed elevated mixed-layer particulate Zn (pZn; 0.08–1.40 nM) and Co (pCo; 8–49 pM) at stations closest to South Africa during the D357-2 late-spring transect (Stations 0.5, 1 and 1.5; Fig. S1), coincident with elevated dZn (0.05–1.82 nM) and dCo (1–43 pM). Furthermore, for the upper 500 m at Stations 0.5 and 1, we found strong positive correlations between particulate aluminium and titanium (pAl / pTi; slope = $41.7 \text{ mol mol}^{-1}$, Pearson's $r = 0.99$, $n = 15$), as well as particulate Fe and titanium (pFe / pTi; slope = $10.2 \text{ mol mol}^{-1}$, Pearson's $r = 0.99$, $n = 15$), indicative of a strong lithogenic source. Whilst there are presently no South African sedimentary data against which we can compare our water column values, our pAl / pTi and pFe / pTi slope ratios are in excess of upper crustal mole ratios (34.1 and 7.3 mol mol^{-1} , respectively; McLen-

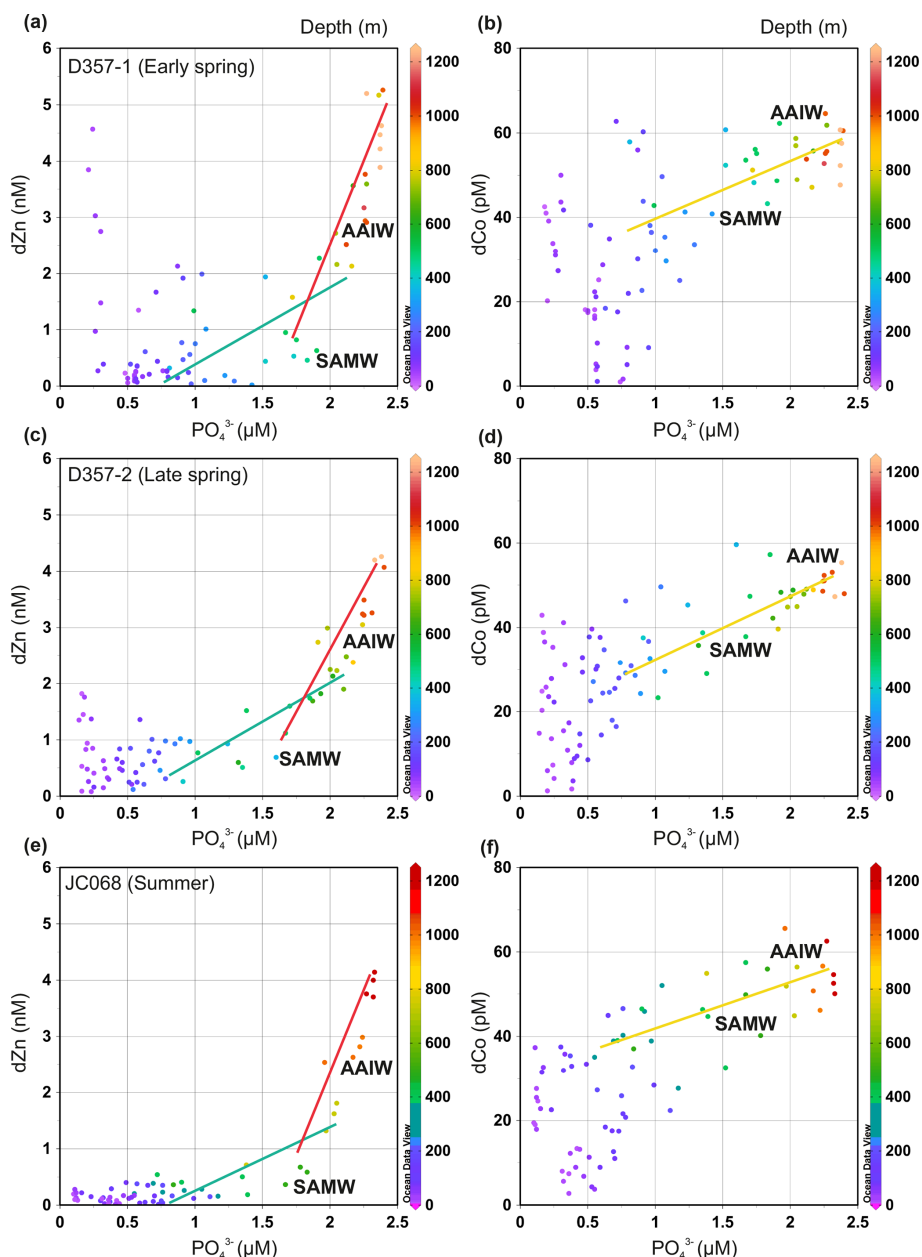


Figure 4. The dissolved Zn and Co versus PO_4^{3-} distribution for the south-eastern Atlantic during early-spring (a, b; D357-1), late-spring (c, d; D357-2) and summer (e, f; JC068) transects. The green and red lines indicate the $\text{dZn} : \text{PO}_4^{3-}$ regression slopes for SAMW and AAIW, respectively. The yellow line indicates the $\text{dCo} : \text{PO}_4^{3-}$ regression slope for SAMW and AAIW combined. The equations for regression lines are detailed in Supplement Table S1. SAMW: sub-Antarctic mode water; AAIW: Antarctic intermediate water. The full depth $\text{dZn} : \text{PO}_4^{3-}$ relationship along JC068 can be found in Wyatt et al. (2014).

nan, 2001). These 500 m ratios are also steeper than the aggregate slopes for the full depth of the Atlantic Ocean away from hydrothermal sources (32.1 and 7.4 mol mol^{-1} , Pearson's $r > 0.97$, $n = 593$; Schlitzer et al., 2018). Given the refractory nature of lithogenic pTi across diverse oceanic environments (Ohnemus and Lam, 2015), this may suggest the resuspension and dissolution of Agulhas Bank sediments en-

riched in dAl and dFe, followed by westward offshore transport, a common feature of the Bank's physical circulation during spring and summer (Largier et al., 1992). Such processes may in turn provide an additional source of dZn and dCo to STSW of the south-eastern Atlantic. For example, Little et al. (2016) proposed that oxygen-deficient, organic-rich, continental-margin sediments may constitute a signifi-

cant global sink within the marine Zn cycle. These sediments could additionally provide a local source of dZn following remineralization. Recent model outputs have likewise highlighted oxygen-deficient boundary sediments as a dominant external source of Co to the oceans (Tagliabue et al., 2018). Given that oxygen-depleted ($< 45 \mu\text{M}$) bottom waters are prevalent across the western Agulhas Bank (Chapman and Shannon, 1987; Chapman, 1988), considered to arise from high organic-matter input to sediments and its bacterial decomposition, a sedimentary source of dZn and dCo appears likely.

3.4 Trace metal stoichiometry of the upper south-eastern Atlantic

In addition to seasonal variations in the lateral advection of continentally derived trace metals, the lower dZn and dCo concentrations in STSW during summer compared with spring likely reflect differences in biological utilization. Here, we examine the micro- and macronutrient concentration inventories to assess the trace metal stoichiometry of the south-eastern Atlantic over seasonal timescales. The data were grouped into STSW and SASW regimes, with STSW defined by $\theta \geq 15^\circ\text{C}$. This isotherm was located at a mean depth of 144 ± 96 m across the study compared with a mean mixed-layer depth of 39 ± 10 m, and thus the inventories for SASW were determined over this depth for comparison with STSW (Table 2). Early- and late-spring STSW samples in the depth range 20–55 m that clearly exhibited continentally derived elevated dZn and dCo were removed from the analysis in order to compare stoichiometry with respect to biological processes. For SASW, micronutrient sampling did not occur during late spring and therefore only early-spring and summer values are compared.

Distinct temporal trends in the stoichiometric relationship with PO_4^{3-} were evident for both dZn and dCo (Fig. 4). Within STSW, the dZn / PO_4^{3-} inventory ratio ranged from 699–1876 $\mu\text{mol mol}^{-1}$ (Table 2), with the highest value observed during early spring and the lowest during summer. Combined with summer dZn concentrations 4 times lower than early spring, this suggests strong biological uptake of dZn alongside PO_4^{3-} between seasons. In contrast, lower dZn / PO_4^{3-} ratios of 372 and 188 $\mu\text{mol mol}^{-1}$ were observed in SASW during early spring and summer, respectively. Here, the absolute change in dZn concentration between spring and summer was lower than for STSW but was greater for PO_4^{3-} , likely reflecting the increased availability of PO_4^{3-} in these Southern-Ocean-derived waters (Table 2) and an open-ocean phytoplankton community that has lower trace metal requirements than its counterparts north of the STF. Such dZn / PO_4^{3-} ratios sit at the lower end of cellular Zn / P reported for the diatom-type and haptophyte-type phytoplankton typical of this region (~ 100 – $1100 \mu\text{mol mol}^{-1}$; Twining and Baines, 2013, and references therein), highlight-

ing the importance of micronutrient processes with respect to Zn availability.

In contrast to dZn, the spatiotemporal variation observed for STSW dCo / PO_4^{3-} was small, with ratios ranging from 82–129 $\mu\text{mol mol}^{-1}$ (Table 2), likely reflecting external inputs to the oceans and biological Co requirements that are typically 4 times less than for Zn (Ho et al., 2003; Roshan et al., 2016; Hawco et al., 2018). The STSW dCo / PO_4^{3-} ratio decreased between early- and late-spring transects, potentially in part due to the westward expansion of STSW during late spring (Fig. 2) and subsequent mixing with SASW depleted in dCo relative to PO_4^{3-} (Fig. 3). This dilution is likely also true of dZn and Si, yet their STSW concentration inventories may be sufficiently high so as to mask this effect. Unfortunately, an insufficient quantity of late-spring SASW data are available with which to affirm this postulation. The highest dCo / PO_4^{3-} ratio was observed during summer due to the preferential biological removal of PO_4^{3-} relative to dCo.

In SASW, dCo / PO_4^{3-} was consistently low, with ratios of 23 and 26 $\mu\text{mol mol}^{-1}$ for early spring and summer, respectively. Much higher inventory ratios of $\sim 580 \mu\text{mol mol}^{-1}$ can be calculated over similar depths for open-ocean North Atlantic waters (GA03 Stations 11–20; Schlitzer et al., 2018), likely reflecting an elevated atmospheric Co input and/or an extremely low surface PO_4^{3-} inventory (Wu et al., 2000; Martiny et al., 2019).

Our results provide evidence for the greater availability and preferential removal of dZn relative to dCo in the upper water column the south-eastern Atlantic based on STSW dZn / dCo inventory ratios of 19, 17 and 5 mol mol^{-1} for the three transects and SASW ratios of 13 and 7 mol mol^{-1} for early spring and summer, respectively (Table 2). With relatively consistent inter-seasonal dCo inventories for STSW and SASW, indicating a more balanced ecophysiological regime with regard to dCo organization, the change in dZn / dCo inventory stoichiometries principally reflects changes in dZn concentration. We postulate that the inter-seasonal variations in dZn and dCo availability and stoichiometry of the south-eastern Atlantic reflect changes in the relative nutritional requirement of resident phytoplankton and/or biochemical substitution of Zn and Co to meet nutritional demand.

3.5 Phytoplankton controls on trace metal ecological stoichiometry

Here we discuss the principle phenomena that together likely explain our observations of seasonally decreasing dZn / dCo inventory stoichiometries in STSW and SASW of the south-eastern Atlantic, i.e. the preferential removal of dZn relative to dCo, leading to low dZn availability, and differences in phytoplankton assemblages with different cellular-metal requirements.

Satellite images show elevated surface chlorophyll concentrations across the south-eastern Atlantic STF compared with waters farther north and south, with peak concentrations observed during summer in January 2012 (Fig. 1). Profiles of total chlorophyll *a* concentration (Fig. S2) also show maximum summer values in the upper water column of STSW (1.02 mg m⁻³) and SASW (0.49 mg m⁻³) compared with spring values (< 0.61 and < 0.36 mg m⁻³, respectively). This is consistent with the hypothesis that increasing irradiance, coupled with shallower mixed-layer depths (de Boyer Montégut et al., 2004), results in enhanced growth conditions across the STF between September and February (Browning et al., 2014). Diagnostic pigment analyses (Fig. 5a) indicated that eukaryotic nanophytoplankton, specifically *Phaeocystis*-type haptophytes, dominated the early-spring STSW chlorophyll *a* content (73 %) but with a reduced contribution during summer (20 %). Maximum growth rates for cultured *Phaeocystis antarctica* have been achieved under elevated Zn concentrations (Saito and Goepfert, 2008), and thus, the dominance of this haptophyte would likely contribute to the removal of dZn between spring and summer. Furthermore, an increased summer diatom contribution (13 % chlorophyll *a* compared with near-zero during spring transects) would have further reduced the dZn inventory, with diatoms having at least 4 times higher cellular-Zn/P ratios than co-occurring cell types (Twining and Baines, 2013).

The fact that both *Phaeocystis* and diatomaceous nanophytoplankton maintain a contribution to the summer STSW chlorophyll *a* complement, when dZn availability is low, is intriguing. Both *P. antarctica* and the large coastal diatoms *Thalassiosira pseudonana* and *Thalassiosira weissflogii* have been shown to be growth-limited in culture by free-Zn²⁺ concentrations ≤ 10 pM (Sunda and Huntsman, 1992; Saito and Goepfert, 2008). A simple estimate of summer STSW free-Zn²⁺ availability, based on North Atlantic organic-complexation data (> 96 %; Ellwood and Van den Berg, 2000), indicated that free Zn²⁺ averaged 6.3 ± 5.3 u_c pM, suggesting the potential for growth limitation of these phytoplankton. In addition, when comparing the south-eastern Atlantic dZn stoichiometry with the cellular requirements of phytoplankton grown under growth-rate-limiting conditions (Fig. 6), we found summer STSW dZn : PO₄³⁻ to be in deficit of the requirements of coastal *T. pseudonana* but not those of the smaller, open-ocean diatom *Thalassiosira oceanica*. The variation in cellular Zn/P between small and large phytoplankton is related to the higher surface-area-to-volume ratio of smaller cells and the limitation of diffusive uptake rates at low Zn²⁺ concentrations (Sunda and Huntsman, 1995). This would suggest that the lower dZn availability in summer STSW should influence phytoplankton species composition by selecting for smaller organisms with lower cellular-Zn requirements, confirmed by a ratio of picophytoplankton to nanophytoplankton at least 4 times higher during summer compared with spring values. The comparison further implies that the presence of *Phaeocystis* and diatoms in summer

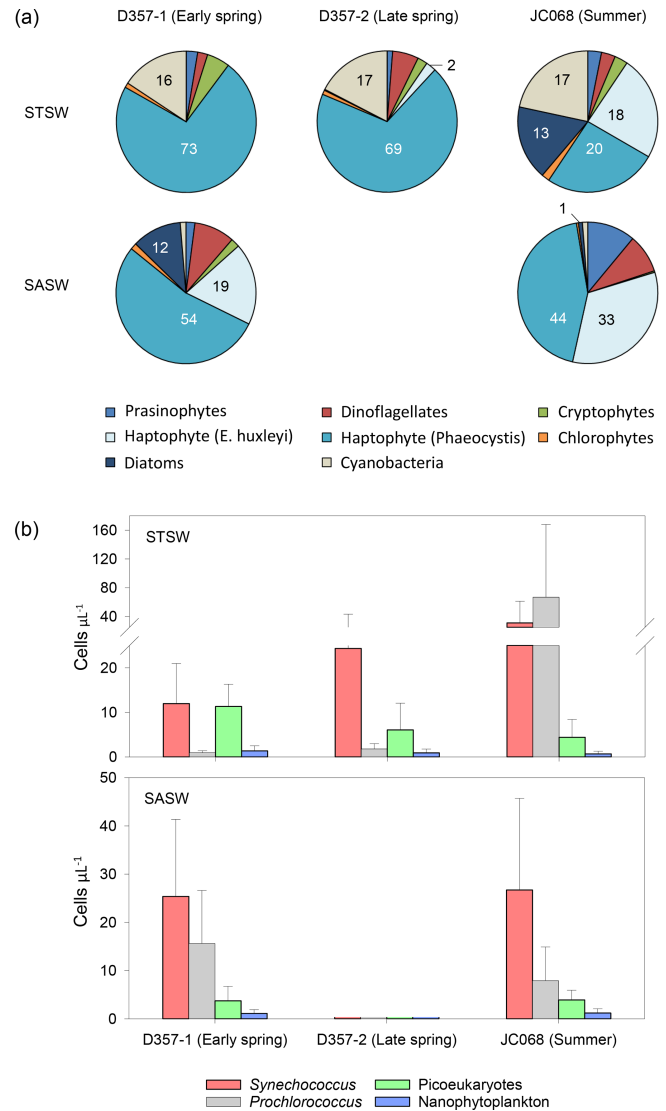


Figure 5. Seasonal differences in (a) pigment-derived taxonomic contributions to total chlorophyll *a* (percentage) and (b) AFC counts of *Synechococcus*, *Prochlorococcus*, nanophytoplankton (approx. > 2 μm) and photosynthetic picoeukaryotes (approx. < 2 μm) in the south-eastern Atlantic.

STSW may be linked with their metabolic Zn–Co–Cd substitution capability, potentially allowing them to overcome some portion of their Zn deficiency. Largely connected to carbonic anhydrase enzymes, several species of eukaryotic phytoplankton are capable of biochemical substitution of Zn, Co or Cd to maintain optimal growth rates under low-trace-metal conditions (Price and Morel, 1990; Sunda and Huntsman, 1995; Lee and Morel, 1995; Lane and Morel, 2000; Xu et al., 2007; Saito and Goepfert, 2008; Kellogg et al., 2020). For example, metabolic substitution of Co in place of Zn has been observed to support the growth of *P. antarctica*, *T. pseudonana* and *T. weissflogii* in media with Zn²⁺ < 3 pM (Sunda and Huntsman, 1995; Saito and Goepfert, 2008; Kel-

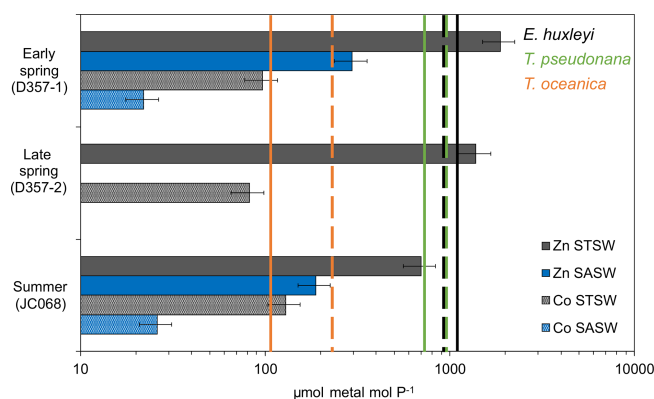


Figure 6. Metal / PO_4^{3-} inventory ratios for the upper water column of the south-eastern Atlantic (horizontal bars) compared with laboratory estimates of cellular ratios in eukaryotic phytoplankton below which growth limitation occurs (solid vertical lines represent Zn/P with no added Co to media, whilst dashed lines represent Co/P with no added Zn; phytoplankton data from Sunda and Huntsman, 1995). Error bars on inventory ratios represent 20% combined uncertainty for dZn and dCo analyses (see Sect. 2.2). This figure is adapted from that in Saito et al. (2010) and implies that inter-seasonal differences in metal : PO_4^{3-} stoichiometry could impact phytoplankton community composition in the south-eastern Atlantic.

logg et al., 2020). Thus, the lower mixed-layer dCo inventory of summer STSW relative to early spring may be in part related to enhanced dCo uptake through biochemical substitution alongside the growth of phytoplankton with distinct Co requirements.

In contrast to *Phaeocystis*, *Emiliania huxleyi*-type haptophytes were near-absent in spring STSW (< 5% chlorophyll *a*; Fig. 5a) and increased in contribution during summer (18%). *E. huxleyi* appear to have a biochemical preference for Co over Zn (Xu et al., 2007), which could potentially be a contributing factor to the increased fraction of this haptophyte in summer STSW. Based on Co organic-complexation data for south-eastern Atlantic STSW (> 99%; Bown et al., 2012), however, even the maximum dCo concentration of 56 pM (estimated free $\text{Co}^{2+} = 0.56 \pm 0.11 \mu\text{c pM}$) observed for STSW during this entire study would limit the growth of cultured *E. huxleyi* in the absence of Zn or Cd (Sunda and Huntsman, 1995; Xu et al., 2007). This is supported by inter-seasonal dCo / PO_4^{3-} stoichiometries in deficit of the cellular requirements of cultured *E. huxleyi* (Fig. 6). Despite this, Xu et al. (2007) showed that *E. huxleyi* can maintain significant growth at only 0.3 pM Co^{2+} in the presence of Zn, with the limitation by and substitution of these metals reported to occur over a range of free-ion concentrations (0.2–5 pM) that are relevant to summer conditions of the south-eastern Atlantic. This assessment implies an additional need for Zn in phytoplankton nutrition due to low dCo availability throughout the south-eastern Atlantic, which may accelerate the decrease in dZn / dCo inventory ratio between seasons.

The elevated summer STSW chlorophyll *a* concentrations were accompanied by increased cell concentrations of the *Synechococcus* and *Prochlorococcus* (up to 100 and 400 cells μL^{-1} , respectively) relative to early-spring abundance (Fig. 5b). This pattern suggests an inter-seasonal community shift towards smaller picocyanobacterial cells that is coincident with decreased dZn availability. *Synechococcus* and *Prochlorococcus* are thought to have little or no Zn requirement and relatively low Co requirements (growth-limited by $\leq 0.2 \text{ pM Co}^{2+}$; Sunda and Huntsman, 1995; Saito et al., 2002). This, alongside their small cell size, hence greater capacity for acquiring fixed nitrogen under conditions where this nutrient is depleted, may allow these prokaryotes to flourish following depletion and export of Zn associated with *Phaeocystis* and diatom blooms. This supposition is supported by a persistently high abundance of *Synechococcus* and *Prochlorococcus* (> 1000 cells μL^{-1}) relative to eukaryotic nanophytoplankton in the dZn-depleted surface waters of the Costa Rica Dome (Saito et al., 2005; Ahlgren et al., 2014; Chappell et al., 2016). Here, surface dCo concentrations were maintained above those of surrounding waters by the biological production of Co-binding ligands (Saito et al., 2005). The increased abundance of these prokaryotic autotrophs in summer STSW of the south-eastern Atlantic may have also contributed to the inter-seasonal decrease in dCo inventory.

In contrast to STSW, cell counts of eukaryotic phytoplankton and prokaryotic cyanobacteria in SASW varied little between early spring and summer (Fig. 5b), indicative of a more balanced ecophysiological regime. The fractional contribution of *Phaeocystis* (Fig. 5a), the dominant contributor to the SASW chlorophyll *a* complement, was similar between transects at 54% and 44%, respectively, whilst the contribution of *E. huxleyi* increased from 19%–33% between spring and summer, respectively. Whilst it is proposed that the low Fe supply rate to these waters provides a dominant control on phytoplankton biomass and composition (Browning et al., 2014), low dZn and dCo availability may also be important drivers of such change. The summer SASW dZn inventory ($0.08 \pm 0.07 \mu\text{c nM}$) and stoichiometry with PO_4^{3-} (Fig. 6) indicate growth-limiting conditions for *Phaeocystis* and *E. huxleyi* in the absence of cambialistic metabolism (Sunda and Huntsman, 1995; Saito and Goepfert, 2008; Xu et al., 2007). The presence of these phytoplankton therefore indicates that Zn biochemical substitution occurs in oceanic waters of the south-eastern Atlantic. A lower Co half-saturation growth constant for cultured *P. antarctica* ($K_m \approx 0.2 \text{ pM Co}^{2+}$) compared with *E. huxleyi* ($K_m \approx 3.6 \text{ pM Co}^{2+}$) further suggests that *Phaeocystis* species may more effectively occupy low-dZn and low-dCo environments (Saito and Goepfert, 2008), such as SASW of the South Atlantic.

Conversely, the absence of a significant diatom contribution to summer SASW chlorophyll *a* (Fig. 5a) relative to early spring is surprising as the summer dZn / PO_4^{3-} inven-

tory ratio is in excess of the cellular-Zn/P requirements of typical oceanic diatoms such as *T. oceanica* (Fig. 6). Furthermore, whilst the $d\text{Co}/\text{PO}_4^{3-}$ ratio of summer SASW is in deficit of the cellular Co/P below which growth limitation of *T. oceanica* may occur, this species has been shown to grow effectively at $\text{Co}^{2+} < 0.1 \text{ pM}$ in culture in the presence of Zn (Sunda and Huntsman, 1995). The low diatom fractional contribution to summer SASW may be instead related to low Fe availability (Browning et al., 2014) and stress-induced Si exhaustion. In support of this, we calculate summer SASW mixed-layer Si concentrations ($0.9 \pm 0.3 \mu\text{M}$) to be 50 % of early-spring values ($1.8 \pm 0.2 \mu\text{M}$), with a dissolved NO_3^-/Si stoichiometry of 3.8 mol mol^{-1} , close to the 4 mol mol^{-1} shown to limit diatom growth in culture (Gilpin et al., 2004) and in contrast to the 2.9 mol mol^{-1} calculated for early spring.

4 Conclusions

We report the distributions of dZn and dCo in the upper water column of sub-tropical and sub-Antarctic waters of the south-eastern Atlantic during austral spring and summer periods. We identify an apparent continental source of dZn and dCo to sub-tropical waters at depths between 20–55 m, derived from sedimentary inputs from the Agulhas Bank. In contrast, open-ocean sub-Antarctic surface waters displayed largely consistent inter-seasonal mixed-layer dZn and dCo concentrations, indicating a more balanced ecophysiological regime with regard to their organization. The vertical distributions of dZn and dCo in the upper water column were similar to that of PO_4^{3-} , indicating that biological drawdown in surface waters and mixing with underlying Southern-Ocean-derived waters travelling equatorward significantly influence their distribution. Absolute trace metal concentrations alongside concentration inventory ratios suggest the preferential utilization of dZn relative to dCo in the south-eastern Atlantic, with dZn/dCo decreasing from 19–5 mol mol^{-1} between early spring and summer in STSW and from 13–7 mol mol^{-1} in SASW. This pattern is consistent with our understanding of the cellular requirement of phytoplankton (Twining and Baines, 2013). The inter-seasonal removal of dZn results in summer concentrations that are potentially growth-limiting for certain phytoplankton species estimated to be present in these waters by diagnostic pigment analyses. We therefore suggest that cambialistic metabolic substitution between Zn and Co and potentially Cd is an important factor regulating the growth, distribution and diversity of phytoplankton in the south-eastern Atlantic.

Data availability. The trace metal and macronutrient data sets used for analyses in this study are available at <https://www.bodc.ac.uk/geotraces/data/idp2017/> (Schlitzer et al., 2018, GEOTRACES GA10) and phytoplankton data at <https://www.bodc.ac.uk/> (BODC, 2021).

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Author contributions. MCL and EPA acquired the funding. NJW, MCL, AM, TJB, EMSW and HAB collected samples at sea. NJW conducted the Zn and Co measurements, EMSW the macronutrient measurements and TJB the phytoplankton measurements. NJW prepared the manuscript with significant contributions from all co-authors.

Competing interests. The authors declare that they have no conflict of interest.

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