



Particulate Rare Earth Element behavior in the North Atlantic (GEOVIDE cruise)

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10 **Abstract.** Particulate concentrations of the fourteen Rare Earth Elements (PREE), yttrium and 232-thorium have been measured in two hundred samples collected in the epipelagic (ca 0-200 m) and the mesopelagic (ca 200-1000 m) zones of the North Atlantic, during the GEOVIDE cruise (May/June 2014, R/V Pourquoi Pas ?, GEOTRACES GA01). Particulate cerium (PCe) concentrations vary from 0.2 pmol.L⁻¹ to 16 pmol.L⁻¹, particulate neodymium (PNd) ones from 0.09 pmol.L⁻¹ to 6.1 pmol.L⁻¹ and particulate ytterbium (PYb) ones from 0.01 pmol.L⁻¹ to 0.5 pmol.L⁻¹. PREE concentrations are higher close to
15 the Iberian margin and on the Greenland shelf, where PREE concentrations normalized to Post Archean Australian Shale (PAAS) display a positive Ce anomaly between 0.3 and 3, and a light REE (LREE) enrichment compared to heavy REE (HREE) illustrated by high PNd_N/PYb_N ratios (normalized to PAAS). The lithogenic fraction of the particulate REE concentration is closely related to the margin morphology and the hydrodynamic context: off the Iberian margin, up to 100%
20 of the PREEs are lithogenic and this lithogenic input spreads westward along isopycnals as intermediate nepheloid layers (INL) up to 1700 km away. Lithogenic inputs are also observed along the Greenland and Newfoundland margins, although the circulation stacks them along the coasts. PREE distributions are also controlled by the biological uptake in the surface layers and remineralization processes deeper. Low surface concentrations and some normalized REE patterns displaying a negative Ce anomaly and HREE enrichment indicate freshly formed biogenic particles. A significant relationship between biogenic silica (BSi) and PHREE is also observed in the diatom blooms occurring in the Labrador and Irminger seas. PHo/PY ratio was
25 calculated in order to identify processes independent of the ionic radius. However, we could not firmly assess the role of the iron hydroxides in the scavenging phases of these elements.

1 Introduction

Marine particles are the main way to transfer chemical species to the deep ocean together with the convection of water masses. Particles are abundant in the upper ocean, where dust inputs or massive blooms occur (up to 1000 µg.L⁻¹), but their
30 concentration decrease with depth (5 to 60 µg.L⁻¹ on average in the subsurface and deep ocean, (McCave and Hall, 2002; Stemann et al., 2002). However, particles are up to 1000 times more concentrated in elements than the dissolved phase (Lam



et al., 2015), and especially in trace elements. For example, in the subpolar North Atlantic (GEOTRACES GA01 cruise, the section studied here), particulate Fe (PFe) concentration can reach 50 nmol.L^{-1} when dissolved Fe (DFe) does not exceed 2.5 nmol.L^{-1} (Gourain et al., 2019; Menzel Barraqueta et al., 2018; Tonnard et al., 2018). The size spectra between the particulate and the dissolved phase is continuous and the separation between both pools is operational, depending on the porosity of the filters used to discriminate the two phases, usually $0.4 \mu\text{m}$ (Planquette and Sherrell, 2012). Concentrations then depend on the choice of this limit, even if the vertical flux is mostly due to the large, dense, sinking particles, in opposition to the smaller and less dense particles that are in suspension in the water column. However, these small suspended particles represent over 80% of the total particle mass (Lam et al., 2015 and references therein). In addition, their higher surface to volume ratios make suspended particles the main drivers of dissolved-particulate exchanges.

In the ocean, three main sources of particles are distinguishable. The first one is lithogenic, with inputs from the rivers, dust deposits, ice melting and resuspension of deposited sediments. The second one is biogenic, and related to the production of fresh organic matter by photosynthetic activity followed by zooplankton grazing, and the following life cycle. The last one consists in authigenic processes such as red clay, oxides and hydroxides precipitation and formation. All these sources and processes lead to a very heterogeneous pool, in time, space and composition, evolving throughout their stay in the ocean and control the density of particles and consequently their fate in the water column. Then, exchanges between the particle and dissolved phases determine the chemistry of seawater and the residence time of the chemical species. They also determine the transfer rate of elements such as carbon and micro-nutrients between the upper layers and the deep ocean, where they are stored for times going from the global circulation scale (500 – 1000 years) to geological ones.

Oceanic tracers such as rare earth elements (REE) are truly adapted to the study of these exchanges (Jeandel et al., 1995; Kuss et al., 2001; Tachikawa et al., 1999). Physical and geochemical processes such as aggregation-disaggregation, dissolution, complexation, sorption, mineralization and scavenging lead to a fractionation along the REE series, depending on their origin and intensity. Thus, measuring the distribution of REEs between the solid and dissolved phases can help tracing and quantifying these processes.

The North Atlantic is a key region of the global ocean, as it is the highest oceanic sink of anthropogenic CO_2 (Khatiwala et al., 2013). Indeed, it is together i) a major place of deep water formation, mainly by convection, which drives the Atlantic meridional overturning circulation (AMOC), and ii) a productive area, representing up to 18% of the global oceanic primary production (Sanders et al., 2014).

In this context, we present the first basin scale section of PREE concentrations and fractionation patterns obtained for suspended particles collected in the North Atlantic (SPNA), along the GEOVIDE section (GA01 GEOTRACES cruise), from the surface to 1500 m. We will specifically discuss processes affecting the PREE distribution such as lithogenic inputs from the margins, influence of biological activity and the role of ionic radius on their fate in the water column.



2 Methods

2.1 Study area: hydrographical and biogeochemical context

70 Samples have been collected in the epipelagic and mesopelagic zones (0 m – 1500 m) during the GEOVIDE cruise (16th of May 2014 to 30th of June 2014, R/V Pourquoi Pas ?) along the transect presented in Fig. 1. This figure also presents the main surface currents, as described in details (Zunino et al., 2017) and (García-Ibáñez et al., 2018), together with the three main biogeochemical provinces identified by Longhurst (1995) and described in details by Lemaitre et al. (2018). The position of the stations where PREEs were sampled (Fig. 1) were chosen to be representative of the diversity of biogeochemical provinces and water masses (Fig. 2).

75 Warm and salty waters coming from the tropical Atlantic are advected toward the Arctic by the North Atlantic Current (NAC, see Table 1 for abbreviations list). In response to air-sea exchanges and mixing with polar waters, surface waters become colder and fresher, but more importantly, denser. They tend thus to mix with underlying waters, particularly during convecting events triggered by storms. In the Nordic Seas (between 65°N and 80°N), the water column can be ventilated down to the bottom, while convection never exceeds 2000 m in the subpolar gyre. The freshly formed deep water then returns southward mainly via western boundary currents.

80 The North Atlantic Subtropical (NAST) province is characterized by warm and salty waters (García-Ibáñez et al., 2018; Longhurst, 1995; Reygondeau et al., 2018; Zunino et al., 2017). This province is depleted in nutrients, yet under influence of margin inputs, displayed a declining bloom of cyanobacteria during the cruise (Lemaitre et al., 2018). Stations #1 and #13 were sampled in the NAST. The North Atlantic Drift region (NADR) is located between the NAST and the Reykjanes ridge, with higher nutrient concentrations than in the NAST (Longhurst, 1995). A strong bloom of coccolithophorids, with a
85 maximum intensity in the Icelandic basin, occurred during the cruise, generating the highest primary production rate observed on GEOVIDE ($1740 \text{ molC.m}^{-2}.\text{d}^{-1}$, station #26, Fonseca-Batista et al., 2019) and high carbon export (up to $80 \text{ molC.m}^{-2}.\text{d}^{-1}$, station #32, Lemaitre et al., 2018). Four open ocean stations were sampled in this province: in the northern branch of the NAC (station #21), at the Subpolar front (station #26), in the northern branch of the NAC (station #32) and on the Reykjanes Ridge (station #38).

90 West of the Reykjanes Ridge, the Irminger and Labrador Seas (Fig. 1) are rich in nutrients, and belong to the Arctic region (ARCT). Large blooms of diatoms occurred in this area, which a maximum intensity at the end of May, in other words three weeks before GEOVIDE sampling in the Labrador Sea and one month before the sampling in the Irminger Sea. The western part of the ARCT region is under the influence of the Newfoundland margin. In this province, station #44 was sampled in the middle of the gyre of the Irminger Sea, station #51 in the EGCC and station #53 on the Greenland shelf. In the Labrador Sea,
95 station #64 is in the West Greenland Current (the continuity of the EGCC after it passed Cape Farewell) and station #69 is in the area of formation of LSW, where strong convection events occurred the winter before GEOVIDE (García-Ibáñez et al., 2018; de Jong and de Steur, 2016). Westward, the station #77 is close to the Newfoundland margin (ca 300 km).



2.2 Sampling at sea

Two sampling systems have been deployed during GEOVIDE to collect suspended particles: a standard CTD rosette equipped with 12 L Niskin bottles and a clean CTD rosette equipped with 12 L GO-FLO bottles. The standard rosette was used to collect samples dedicated to the concentration analyses of dissolved and particulate barium in excess (Ba_{xs}), dissolved and particulate REEs (including Nd isotopic composition) as well as ancillary parameter analyses. Ba_{xs} and PREE chemical treatment and analyses were conducted on the same samples: Ba_{xs} was first measured at the Royal Museum for Central Africa (Tervuren, Belgium), then PREEs were later analyzed at LEGOS (Toulouse, France; this work). Ba and ^{232}Th concentrations were measured at both places, allowing us to compare our procedures. Regarding the samples collected with the clean rosette, Ba, ^{232}Th and Y (with the latter belonging to REEs, named YREEs when Y is included) were also measured. Ba and ^{232}Th were used to compare the data obtained with the standard and clean rosette procedures (as done for Ba_{xs} by Lemaitre et al., 2018, with comparable results: $r^2= 0.61$, $p < 0.01$, Fig. S1 in supplementary) distinguishable by the sampling systems, filtration method, chemistry performed on filters and analysis. The comparison of Y concentrations from the two procedures validated the use of our standard rosette to sample YREEs, less prone to contamination than trace metals, as shown by van de Flierdt et al. (2012).

Sampling method and sample preparation for water collection with the standard rosette for Ba_{xs} and PYREE analyses described here follow those of Lemaitre et al. (2018). Sampling was focused on the epipelagic and mesopelagic zones (0 m – 1500 m). On board, four to eight liters were filtered using clean slightly pressurized containers. Bottles were shaken three times as recommended in the GEOTRACES cookbook, to avoid the loss of particles by sticking to the walls or settling at the bottom of the bottle. Seawater was then poured in the Perspex containers at the base of which polycarbonate filters of 0.4 μm porosity (Nuclepore®, 47 mm or 90 mm of diameter) were mounted. After sample filtration, the container was rinsed with ≤ 5 mL of ultra-pure water (Milli-Q; 18.2 $\text{M}\Omega\cdot\text{cm}$) to remove the maximum of sea salt deposited on the membranes. Finally, filters were removed using plastic tweezers and were dried under a laminar flow hood at ambient temperature before being stored in clean petri dishes.

Regarding samples collected with the clean rosette, sampling method and samples preparation are described in Gourain et al. (2019).

2.3 Sample preparation and analysis

Filters were first cut in two parts using a ceramic blade. One half was archived, while the other half was placed in a clean Teflon vial (Savillex®). The filter was then digested with a strong acid solution made of 1.5 mL HCl, 1 mL HNO₃ and 0.5 mL HF, all concentrated (Merck® Suprapur Grades). Vials were subsequently left on hot plates at 90°C overnight. The PREE concentrations were measured on 2 mL of the archived solutions, which were placed in clean 5 mL polypropylene tubes and doped with a solution containing In and Re (ca 100 ppt of both tracers) in order to correct matrix effects and sensitivity shifts during analysis. In addition to REEs, Y, Ba and ^{232}Th concentrations were measured in the same leaching



130 solution. Analyses were performed at the Observatoire Midi Pyrénées (Toulouse, France) using a high-resolution inductively
coupled plasma mass spectrometer (SF-ICPMS, Element XR, Thermo Fischer Scientific®) coupled to a desolvating nebulizer
(Aridus II, CETAC Technologies®) to minimize oxides and hydroxides production rates and thus (hydr)oxides interferences
(Aries et al., 2000). Oxide production rates were determined at the beginning and the end of every session using a mono-
elementary Ce solution ($\text{CeO} < 0.03\%$). Other REE (hydr)oxides rates were then determined using the constant proportionality
135 factor between them (Aries et al., 2000), previously determined with the same analytical configuration. Interferences represent
0.001% to 1% of the signal except for Eu (0.3% to 10%).

A five points calibration curve was established with a multi elemental standard at the beginning, the middle and the end of the
analysis, while a standard solution concentrated with $20 \cdot 10^{-12} \text{ g} \cdot \text{g}^{-1}$ of REE was measured every 5 samples. The certified
reference material SLRS-5 (NRC Canada) was systematically analyzed with the samples and their concentrations are within
140 the error bar of the consensual values published by Yeghisheyan et al. 2013, with a smaller error (see Figure S1 in
supplementary). Reproducibility was assessed by measuring two or three times several samples from the same leaching
solution, and varied from 0% to 20%, like the error.

Procedural blanks have been estimated by conducting the chemistry on clean, unused filters. The average chemical blank ($n=$
8) represents 0.01% to 5% of the sample concentrations, except for Y and Lu for which the contribution of the blank was
145 generally higher (between 1% and 30%).

Uncertainty of each concentration, estimated from error propagation was between 20% and 30% (and can be up to 40% for
Eu) of the concentration. In addition to the mass spectrometry standard deviation the other sources contributing to the final
error bar of the concentrations are: the proportion of filter analyzed, the volume of leachate and the volume taken in the archive
for the analysis. For details on errors, see Fig. S2 in supplementary.

150 Thus the hypothesis of homogeneity is assumed, in the light of the apparent consistency of suspended particles on filters.
The same digest solutions were also analysed at the Royal Museum for Central Africa (Tervuren, Belgium) mainly for
determining Ba and some other element concentrations, including ^{232}Th . It was conducted using an inductively coupled plasma
quadrupole mass spectrometer (ICP-QMS; X Series 2 Thermo Fischer®). Ba, Y and ^{232}Th were also measured in clean rosette
samples. The digestion procedure and analytical method are detailed in Gourain et al. (2019). Concentrations of Ba and ^{232}Th
155 are consistent between the analysis conducted in Brest and the PREE analysis in Toulouse. “Toulouse” Ba concentrations vs.
“Tervuren” Ba concentrations give a regression slope of 0.87 ($r^2=0.90$, $n=198$). For ^{232}Th , “Toulouse” concentrations vs.
“Tervuren” concentrations give a slope of 1.05 ($r^2=0.98$, $n=198$), which is also very comparable. Ba and Y analyses were used
to compare the clean rosette and standard rosette procedures. Consistent Ba concentrations were found when compared to the
results of Lemaitre et al. (2018; $r^2=0.61$, $n=66$). Y concentrations measured in Toulouse on samples collected with the standard
160 rosette match the concentrations obtained in Brest on the clean samples, with a regression slope of 0.93 ($r^2=0.82$, $n=78$ points
at same depths). Thus both procedures are suitable for PREE analysis. In addition, these validations allow us to discuss the
PREE concentrations with the trace metal ones from Gourain et al. (2019).



3 Results

The data set of PYREE, PBa and P²³²Th concentrations is compiled in Table 2. For sake of clarity, we only displayed PCe, PNd and PYb concentrations (Fig. 2 and 3), these three REEs representing the light REEs (Nd), heavy REEs (Yb) and a specific behavior (Ce). LREEs and HREEs are supposed to react to different processes during dissolved-particulate exchanges. Ce has a IV oxidation state in addition to the III oxidation state common to all REEs and its oxidation onto particles prevent desorption. The Ce (III)/Ce(IV) distribution is therefore a proxy of redox and desorption processes.

PCe (Fig. 2) concentrations are higher than PNd (Fig. 3 and B) concentrations, which are higher than PYb concentrations (Fig. 3 C and D), in agreement with their respective natural abundance and reactivity.

3.1 Cerium

Particulate Ce concentrations (PCe, Fig. 2) vary between 0.2 pmol.L⁻¹ (station #64) and 16.3 pmol.L⁻¹ (station #32; Fig. 2). They are higher close to the Iberian margin (station #1: 1 pmol.L⁻¹ < PCe < 9.4 pmol.L⁻¹) and on the Greenland shelf (station #53: 5.7 pmol.L⁻¹ < PCe < 14.6 pmol.L⁻¹). In the NAST (for station #13) and the NADR regions, vertical profiles present a surface or subsurface maximum at all stations. Below 200 m depth, PCe concentrations decrease and reach a value of 2 pmol.L⁻¹ in the mesopelagic zone. PCe concentrations are higher to the east of the polar front (stations #13 and #21) compared to the west (stations #26, #32 and #38). A second maximum is observed at greater depth at station #13 and in the NADR region (except close to the subarctic front, at station #26). In the ARCT region, surface PCe concentrations are lower and increase at the bottom of the epipelagic zone for all open-sea stations (PCe > 1 pmol.L⁻¹). Maximum concentrations are observed at the bottom of the epipelagic zone at stations #44, #64 and #69. PCe concentrations are more variable in the mesopelagic zone of the ARCT region than in the NADR region, and higher than the ones observed at the surface except at station #69 where they are about within the same range (1 pmol.L⁻¹ < PCe < 2 pmol.L⁻¹). PCe profiles differ from that of PNd and PYb at two stations only: station #38, where peaks of PCe are observed at 100 m and 800 m, that are not observed for the other PREE profiles; station #44, where PCe concentrations are more variable in the epipelagic zone than PNd and PYb, with maxima at 120 m and 160 m that are not observed for other PREEs.

3.2 Neodymium

As for PCe (and other PREEs, see supplementary information), PNd concentrations are the highest close to the Portugal and Greenland margins with values up to 4.5 pmol.L⁻¹ in the upper 100 m (Fig. 3 a and B). Concentrations decrease as the distance to margins increases, as seen at stations #13 where PNd hardly reaches 1 pmol.L⁻¹. Low PNd values were also measured at station #77, although this station is relatively close to the Newfoundland margin, but located out of the continental shelf.



3.3 Ytterbium

Distributions of PNd and PYb (Fig. 3) differ on several points: three stations (#13, #44 and #69) display a maximum in subsurface for PYb that is not observed for PNd, whereas a local maximum in PNd is identified at 160 m at stations #64 and #69, but not for PYb. In the open ocean, at stations #21, #26, #32 and #38, concentrations are higher in the surface layer (from 0 m to 200 m). The highest concentrations were determined in the NADR region, which was the most productive during the cruise (Fonseca-Batista, 2018). Concentrations then decrease with depth to become constant, except at station #38 where they increase again in the mesopelagic zone. In the ARCT region, surface concentrations of PNd are lower at 100 m compared to 250 m, similar to station #1 while the contrary is observed in the NADR region.

3.4 PNd_N/PYb_N ratios

To highlight a possible fractionation between LREE and HREE, the PNd_N/PYb_N ratio is calculated from concentrations normalized to PAAS, in order to get rid of the natural abundance effect of the REEs. Results are presented in Fig. 4, interpolated along the section. This ratio presents a high variability, changing from 0.01 to 4.3. Higher ratios (> 1) are observed along the margins, decreasing as the distance to the coast increases. PNd_N/PYb_N is lower at the surface (< 0.7) except at stations #1, #38 and #53, and increases in the subsurface layers ($0.7 < \text{PNd}_N/\text{PYb}_N < 1$) in the open ocean. The strongest ratio value is observed in the core of the epipelagic zone at station #21 (Fig. 4C), where high concentrations of PLa, PCe, PPr and PNd are also observed. However, for other stations with a similar enrichment, no high PNd_N/PYb_N ratio are observed.

4 Discussion

4.1 Comparison with other studies

PREE data in suspended particles are very scarce in the literature. To our knowledge, for the North Atlantic, only one other set of concentrations was published by Kuss et al. (2001), who measured PREEs in samples centrifuged from several m³ of water at a depth of 7 m, collected along the 20°W meridian between 30°N and 60°N. These authors observed PCe concentrations ranging between 0.2 pmol.L⁻¹ and 4.9 pmol.L⁻¹ with higher concentrations close to the margins especially near the European one, consistent with our data. Their PNd concentrations of about 0.5 pmol.L⁻¹ to the east of the NADR are also consistent with ours. PNd and PYb concentrations reported by Tachikawa et al. (1999) at a station located in a mesotrophic zone of the north-east tropical Atlantic, directly influenced by Saharan dust (6 g.m⁻².yr⁻¹ to 15 g.m⁻².yr⁻¹, Rea, 1994), are higher than those reported here (PNd = 2.6 pmol.L⁻¹ and PYb = 0.94 pmol.L⁻¹ at 10 m). Contrastingly, these authors observed lower concentrations than ours at the oligotrophic site of their study, where the dust flux is lower than at the mesotrophic site (4-5 g.m⁻².yr⁻¹, Rea, 1994) but higher than that found during the GEOVIDE cruise (2 ng.m⁻³ to 500 ng.m⁻³, Shelley et al., 2017). However, these author's PCe concentrations are similar to those reported in this study, and that for both the mesotrophic and



220 oligotrophic sites. The difference of concentrations observed for the other PREEs can be explained by the high particle
concentrations characterizing the compared to the tropical one, even if dust inputs are higher on the later (Gehlen et al., 2006).
PREE are found both in the lithogenic and authigenic phases of the particles. Schematically, particles are often represented
with a “lithogenic core” coated by authigenic material (Bayon et al., 2004; Sholkovitz et al., 1994). The lithogenic has an
external origin, product of the continental weathering transported by the winds or discharged by the rivers on the margins. The
225 authigenic phase is of internal origin, major phases being biogenic matter (particulate organic matter POM, biogenic silica
BSi, calcium carbonate CaCO₃) and metal oxides and hydroxides such as MnO₂ and Fe(OH)₃. REEs in the authigenic phase
are scavenged by organic coatings and/or iron and manganese oxides and hydroxides that are known to be the main carrier of
REEs, scavenged by adsorption during their precipitation (Bau, 1999; Bau and Koschinsky, 2009). REEs could also be
absorbed in inorganic planktonic tests (CaCO₃, Palmer, 1985 and BSi, Akagi, 2013) or biogenic byproducts as barite
230 (Ba_{xs}, Guichard et al., 1979). LREEs would be more sensitive to oxide phases of Fe and Mn, while HREEs, more soluble, could
react preferentially with biogenic phases (Akagi, 2013; Bertram and Elderfield, 1992; Grenier, 2018; Pham et al., 2019). Their
distribution coefficients are also varying with depth and the nature of the particle phases (Schijf et al., 2015).
Thus, differentiating the distribution of the REEs in the two phases allows us to estimate the fraction implied in scavenging
processes by the authigenic phase, while the lithogenic one enables to picture continental inputs. The high PREE concentrations
235 observed in Fig. 3 close to the Portugal margin and on the Greenland shelf suggest that particulate material is released by the
margins to the water column. Lithogenic REE fraction can be quantified using conservative lithogenic tracers such as Al, Th
or Ti (Gourain et al., 2019; Tachikawa et al., 1997). These authors used Al as a lithogenic tracer, but here we chose to use
²³²Th. Indeed, the lithogenic fractions calculated from particulate Al (PAI) concentrations were often higher than 100% in
surface waters close to the margins, revealing that a fraction of the total PAI is likely in the authigenic phase (Lerner et al.,
240 2018; Van Beueskom et al., 1997). In addition, Al being more prone to contamination was sampled with the clean rosette
(Gourain et al., 2019), while ²³²Th used for calculation was measured in the same samples as PREEs, collected with the standard
rosette. The concentration of the lithogenic PREE fraction in particles is calculated by multiplying the ²³²Th concentration in
a given sample by the ratio of the considered REE on ²³²Th in the upper crust (Rudnick and Gao, 2014, Eq. (1)).

$$[REE]_{litho} = [^{232}\text{Th}] \times \left(\frac{[REE]}{[^{232}\text{Th}]} \right)_{UCC} \quad (1)$$

245

$$\%REE_{litho} = \frac{[REE]_{litho}}{[REE]} \quad (2)$$

$$REE_{authi} = REE_{total} - REE_{litho} \quad (3)$$

These PREE lithogenic concentrations are then divided by the total PREE concentrations to obtain the percentage of particulate
250 REE with a lithogenic origin (Eq. (2)). The authigenic concentrations are then obtained by subtraction of the lithogenic
concentrations to the total concentrations (Eq. (3)).



The percentage of lithogenic PNd along the section is represented in Fig. 5. Then we chose to represent the average value of the lithogenic fractions of the remaining PREEs for the LREE, excepted for PCe on one hand and for HREE on the other hand. Profiles of five selected stations (#1, #26, #51, #53 and #77) are shown in Fig. 5. These stations are representative of the three
255 different distributions observed along the section. Error bars represent the standard deviation of the resulting averages, the contribution of the error on the concentrations being negligible compared to the later. For some points at station #1 and at 160 m at stations #13 and #32, the calculated lithogenic proportion exceeds 100%, suggesting an excess of ^{232}Th in the particles, likely authigenic, or a difference between the adsorption kinetics of ^{232}Th and REE, as reported by Hayes et al. (2015). In these cases, we capped the lithogenic proportion to 100%.

260 PREE concentrations are normalized to Post-Archean Australian shale, PAAS (Rudnick and Gao, 2014). This normalization allows i) a better diagnostic of the fractionation between PREEs and ii) comparison with patterns in the literature. As shown by the flat PAAS-normalized patterns of the lithogenic fractions (Fig. S3 in supp. mat.), PAAS is a valuable reference to represent the lithogenic material. In addition, it does not present any significant difference in REE composition with shales and loess from Europe, North America and China (Rudnick and Gao, 2014). Normalization to atmospheric depositions has
265 been put aside as these inputs were low during the cruise (Shelley et al., 2017) and normalization to dusts led to patterns depleted in Eu and Gd, and enriched in Tb, Dy, Ho and Er (data from Patey et al., 2015, on dusts collected close to Cape Verde; Fig. S3), less representative of lithogenic inputs than PAAS. Patterns normalized to PAAS are presented in Fig.5 for selected stations. To facilitate readability, patterns of each sample are averaged by layers displaying similar values. Error bars represent the standard deviation of the concentration series, the errors on PAAS concentrations being negligible compared to it. A REE
270 pattern obtained in the Atlantic seawater at 12°S (Zheng et al., 2016) is also represented together with station #26 patterns, for comparison.

4.2 Lithogenic supply by the margins

The high concentrations of PREEs (Fig. 3) at stations #1 and #53 reflect lithogenic inputs from the margins. At these stations, the lithogenic PREE fractions range between 50% and 100%, the highest ones being observed at station #1 (Fig. 5). The
275 relatively flat patterns displayed at these stations for total PREE indicate a weak fractionation of PREEs, with a little enrichment in LREEs due to their lower solubility compared to the HREEs. These maxima can be seen beyond the Subpolar Front until station #32, spreading along the isopycnals 27.05 and 27.4 over 2500 km from the Iberian margin (Fig. 6). Similar maxima have been reported by (Gourain et al., 2019, Fig. 6 B) for lithogenic PFe and PMn), lithogenic PMn being taken by these authors as tracer of sediment resuspension.

280 Above the Greenland shelf, at station #53, the proportion of lithogenic PREE is also high, only slightly lower than at station #1 (median contributions of 59% for PLREE and 83% for PHREE; Fig. 5). Unlike what is observed to the eastern end of the section, these lithogenic particles remain on the shelf and do not spread offshore. Except at the surface for LREE, the lithogenic proportion are lower than 50% at stations #51 and #64 in the Irminger Sea and in the Labrador Sea respectively. This containment of particles along the shelf is explained by the circulation. Indeed, the East Greenland Irminger Current (EGIC)



285 is a strong narrow current bypassing Greenland along its shelf (23.4 ± 1.9 Sv, Daniault et al., 2016), likely preventing exchanges
between the Irminger Subpolar Mode Water (IrSBPMW) and waters of the Greenland shelf, transported by the EGCC current
which flows parallel to the coast (green and orange arrows around the Greenland southeastern tip in Fig. 1). Our observations
are consistent with that of Lacan and Jeandel, 2005, who showed that the Nd isotopic signatures (ϵNd) of SPMW transported
by the EGIC do not vary significantly along the Greenland shelf. In the same way, the lithogenic influence is moderate at
290 station #77, land-ocean exchanges being reduced due to the EGCC again (1.5 ± 0.2 Sv, Daniault et al., 2016). While the
lithogenic fraction is still relatively high at this station ($50\% < \text{REE}_{\text{litho}} < 80\%$ below 150m), the fractionated patterns indicate
that other processes are at play.

High lithogenic proportions are also observed at station #69 but not to the east of the Labrador Sea (station #64; Fig. 5). In
contrast with the Iberian margin, no intermediate enriched layers are observed (Figs. 2 and 3) and the lithogenic fraction is less
295 important and remains roughly constant below 200 m at stations #51 (about 45%) and #77 (around 60%, Fig. 5). Normalized
total concentrations display fractionated PREE patterns, underlining authigenic processes likely at play in this area.

Comparable lithogenic percentages have been reported by Garcia-Solsona et al. (2014) between South Africa and Antarctic,
from 0% at the surface to 80% deeper, with higher lithogenic proportions for HREE than for LREE. Using PAI as a lithogenic
tracer, Tachikawa et al. (1999) evaluated the lithogenic proportion to be between 50% and 80% at the different Eumeli sites
300 in the east tropical Atlantic.

Gourain et al. (2019) reported similar results than ours for PFe and PMn at the same stations along GEOVIDE. These authors
observed strong lithogenic contribution from the Iberian margin spreading until station #32, lower contribution along the
Newfoundland margin and no particular lithogenic contribution along the Greenland margin, in agreement with our
observations. Using lithogenic PMn as a tracer of sediment resuspension, they observe that 100% of PMn is originating from
305 sediment resuspension at station #1 between 250 m and 1000 m (their Fig. 4). Interestingly, Le Roy et al. (comm. pers.)
observed an unexpected maximum of ^{227}Ac activity at 500 m at stations #1 and #21, indicating a strong sediment source, again
consistent with the PREE data. At station #13 at 200 m, however, no lithogenic maximum is identified. This could result from
the compression of isopycnals, leading to the merging of the two maxima observed eastward (Fig. 4). Unfortunately, the
different sampling resolutions for PREE and ^{227}Ac do not permit to further compare data between these tracers except at the
310 surface of station #1, where a maximum of ^{227}Ac is consistent from the lithogenic PREE signal.

These layers highly enriched in lithogenic particles could be attributed to the formation of intermediate nepheloid layers (INL)
at 250 m and 500 m along the Iberian margin, similarly to those revealed slightly north by McCave and Hall (2002). A
contribution of the Mediterranean Water (MW) to these high concentrations and lithogenic proportions cannot be excluded
too, but the lack of data in the core of the MW (1000 m to 1500 m, García-Ibáñez et al., 2018) prevents us from further
315 investigations.

A highly energetic process is needed to enhance such strong resuspension of lithogenic matter. It may be due to the friction
and energetic excitation of internal waves along the continental slope (Cacchione, 2002). Another source can be the erosion of
the coast by the strong current (from $0.05 \text{ m}\cdot\text{s}^{-1}$ to $0.1 \text{ m}\cdot\text{s}^{-1}$) coming out from Gibraltar and flowing northward along the Iberian



margin (Gourain et al., 2019; McCave and Hall, 2002; Zunino et al., 2017). Our observations could also result from a
320 combination of both, with generation of internal waves south of station #1 generating sediment resuspension, those particles
being advected northward by the current.

To sum up, margins can provide significant amounts of particulate lithogenic material to the ocean; nevertheless, occurrence
and magnitude of these inputs are depending on the morphology of the margin and the hydrodynamical forcing, leading (or
not) to nepheloid layer formations.

325 **4.3 REE fractionation: Ce anomalies**

Ce is the only REE having a (IV) oxidation state in the water column. When adsorbed onto particles together with other REEs,
oxidation (biotic or abiotic) makes it less prone to desorption than other REEs, leading to Ce enrichment of the particulate
phase (Byrne and Kim, 1990; Elderfield, 1988; Moffett, 1990, 1994; Tachikawa et al., 1999). This oxidation is thought to
occur for authigenic Ce adsorbed on Fe(OH)₃ and MnO₂ (Bau, 1999; Bau et al., 1996). This P_{Ce} enrichment is commonly
330 quantified by the ratio of the P_{Ce} concentration on the theoretical P_{Ce} concentration calculated using its neighbors P_{Nd} and
P_{Pr} and expressed as Ce*, following Bolhar et al. (2004):

$$\frac{Ce}{Ce^*} = \frac{[Ce]}{2 * [Pr] - [Nd]} \quad (4)$$

In the present set of data, this ratio is always larger than one (positive anomaly) except at stations #26, #32, #51 and #77
335 between the surface and ca. 100 m, where P_{Ce} is depleted compared to other P_{REEs}, as already discussed above. This surface
minimum is followed by a pronounced (Ce/Ce*>3) positive anomaly down to 200 m. At greater depth, the anomaly is relatively
higher in the NADR region compared to the NAST and ARCT regions.

These results indicate that Ce oxidation occurs after particles left the surface, leading to a subsurface maximum. At greater
depths, remineralization rates are high in the ARCT region, moderate in the NAST region and low in the NADR region
340 (Lemaitre et al., 2018). It is likely that lower remineralization rates conduct to higher net fluxes from the dissolved phase
toward the particulate phase, associated with a strong and irreversible scavenging of Ce by adsorption and oxidation, while a
fraction of the other trivalent REEs are released by desorption from the particles (Bau, 1999; Tachikawa et al., 1999). A
stronger anomaly could also reflect higher particle concentrations offering higher surface reactive areas, but particle mass
remains unknown at those depths and beam transmission does not allow identifying higher particle concentrations.

To compare P_{Ce} to Mn and Fe (hydr)oxides, particulate Fe(OH)₃ and MnO₂ concentrations are calculated with the formula of
Lam et al. (2017, Figure S6) using P_{Mn}, P_{Fe} and P_{Al} data from Gourain et al. (2019). The observed decoupling between Ce
anomaly and MnO₂ distributions indicates that more processes are at play than the biologically mediated oxidation that would
lead to similar distributions of the two tracers (Moffett, 1990). Different complexation conditions between these two elements
likely prevent the occurrence of a good correlation. The positive Ce anomaly is not observed when the residence time of
350 particles is short, which is the case in the NADR and the ARCT regions where the export is strong (stations #51 and #77,
Lemaitre et al., 2018). CaCO₃ formation can explain the observed negative anomalies at station #26 and #32 (Garcia-Solsona



et al., 2014; Haley et al., 2005; Sholkovitz and Shen, 1995). However, this hypothesis does not hold at station #21 where CaCO_3 concentration is high while the PCe anomaly is positive (Fig. S4 in supplementary). A surface photoreduction could explain the surface weak anomalies by reducing Ce(IV) in Ce(III) that can be desorbed from particles. The MnO_2 depletion in the surface waters of NADR region would support this hypothesis, although this MnO_2 depletion reaches 500 m (ie below the euphotic layer, Fig S6A in supplementary). In the NAST region, a MnO_2 surface minimum occurs too, but does not correspond to a PCe/Ce* minimum.

4.4 The influence of biological activity on the REE distributions

4.2 The influence of biological activity on the REE distributions

360 The surface of open-sea stations (all of them except #1 and #53) are characterized by a higher lithogenic proportion for LREEs than for HREEs, meaning that the expected relative enrichment of the authigenic phase in LREE -due to their lower solubility- is not observed (see the vertical profiles reported in Fig. 5). Thus, an uncommon enrichment of the authigenic material in HREEs is observed in these samples, also shown in the total PREE patterns (Fig. 5 and Fig. S4 in supplementary). Indeed, these patterns display a negative Ce anomaly on the first hundred meters and enrichment in HREEs characterized by high
365 PYbN/PNdN ratios (>1 , Fig. 4) that can reach 1000 m. Such kind of pattern is classically observed for dissolved REEs, more rarely for PREEs. Suspended PREEs displaying such “dissolved-type” pattern suggest that they have been likely formed through absorption than adsorption processes, the latter leading to fractionation between the REEs, which is not the case during the uptake of REEs in the carbonate planktonic shells (Palmer and Elderfield, 1986). The negative PCe anomaly (Fig. 7) suggests recently formed particles on which Ce oxidation leading to positive PCe anomaly have not occurred yet. All these
370 stations are subject to a strong primary production (Fonseca-Batista et al., 2019), so the preferential transfer of HREEs from the dissolved phase to the authigenic particulate one likely occurs when the biological stripping is active. However, this transfer is more important in the ARCT region than in the NADR region, leading to stronger HREE enrichments, while the highest bloom activity was observed in the NADR region with a maximum at station #26. The high prevalence of coccolithophorids characterizing this bloom (Lemaitre et al., 2018) could explain the relatively low HREE enrichment, except at station #26. In
375 the NADR region, the patterns flatten with depth to present a quasi-lithogenic signature under 60 m, suggesting that particles with a strong organic signature have not reach this depth yet. In the ARCT region, the bloom was dominated by diatoms, still active at station #51 and declining at the others (Fonseca-Batista et al., 2019; Lemaitre et al., 2018). This declining bloom leads to a strong export, but high remineralization rates decrease the biological signature proportion in favor of the lithogenic one at depth (Fig. 5). Even if the characterization of the authigenic phase leans on ^{232}Th and the assumption that REEs and ^{232}Th
380 behave similarly from the original lithogenic source, it is thus very likely that biological uptake appears to have a strong effect on the total REE patterns observed.

A relationship between HREEs and biogenic matter, mostly BSi, have been suggested by Akagi (2013) following thermodynamic calculations. According to this work, 10% to 20% of REEs are forming a $\text{REE}(\text{H}_3\text{SiO}_4)_2^+$ complex with silicic acid, this proportion being more important as the atomic number is low and with depth. Complexation of REE with



385 silicates was further confirmed by Patten and Byrne (2017), although these authors estimated lower complexation constant, thus a less important fraction of REEs complexed by silica. In addition, significant correlations were observed between dissolved Si and dissolved HREE by Bertram and Elderfield (1992; western Indian Ocean), Akagi et al. (2011, North Pacific Ocean); Stichel et al. (2012) and Garcia-Solsona et al. (2014), both in the Atlantic sector of Southern Ocean), Grenier et al. (2018; Kerguelen Islands) and Pham et al. (2019; Solomon Sea). Contrastingly in other areas, the correlation between SiOH₄ and REEs present a curvature or is absent (Patten and Byrne, 2017, their Fig. 7; Zheng et al., 2016, their Fig. 11). Even if the causes of such relationships are not clear, the PYbN/PNdN ratio in the authigenic phase is the highest between the surface and 50 m in the Irminger Sea and the Labrador Sea, where BSi concentrations are also the highest (Sarhou et al., 2018), reflecting a bloom dominated by diatom species (Fig. S5A and B in supplementary material). Although the correlation between BSi and PHREEs stays weak (from R²=0.06 for Ho to R²=0.4 for Lu), this correlation coefficient rises with the atomic mass number (Fig. S5C), showing that BSi has a significant influence on authigenic PREE distributions from Tb to Lu that does not appear for lighter REEs. These correlations may show that in some areas the HREE distributions are linked to the biogeochemistry of silicate, and not only to a conservative mixing as shown by Zheng et al. (2016) and de Baar et al. (2018). This relationship would depend on the abundance and the nature of particles (the occurrence of diatoms), and on the speciation of REEs in the dissolved phase as shown by de Baar et al. (2018). Akagi, (2013) and Akagi et al. (2011) proposed an incorporation of the silica-REEs complexes during the frustules construction, but the mechanism under this enrichment during diatom blooms remains to be clarified. Linking it to usual complexation and adsorption processes is not straightforward since this would imply a sharp break between LREE and HREE affinities with BSi, which has still to be demonstrated. In addition, an effective relationship between BSi and PHREE can be blurred by other scavenging processes implying particulate Mn and Fe (hydr)oxides, also known to influence the slope between LREE and HREE.

405 **4.5 The PAAS-normalized particulate Ho/Y ratio: a proxy of processes independent of the ionic radius**

Yttrium (Y) and the lanthanide holmium (Ho) are characterized by about the same ionic radius and identical charge, making them “geochemical twins” (Bau, 1999). The PAAS-normalized particulate ratio (PHoN/PYN) highlights differences in their distributions, and therefore allows identifying radius-independent fractionation processes affecting YREE in seawater. We choose to normalize PHo/PY measured in our particulate samples to the PAAS ratio to reveal any relative loss or enrichment compared to continental material (Fig. 8). Due to differences of electron configuration, Ho is more prone to establish ionic bounds, and then to be preferentially scavenged by adsorption onto (hydr)oxides as FeOH₃ and MnO₂. In comparison, Y is preferentially scavenged when covalent bounds are established (Censi et al., 2007; Bau, 1999; Bau et al., 1995). Along the GEOVIDE section, PHoN/PYN ratio varies between 0.4 and 1.5, with most of the values less than 1 (i.e. depleted compared to PAAS), which does not support the expected preferential scavenging of Ho. Moreover, PHoN/PYN relationship with FeOH₃ and MnO₂ doesn't fit to any pattern (Fig. 9). PHoN/PYN is higher when [Fe(OH)₃] > 10⁻² μg.L⁻¹, while there is no evidence of a higher HoN/YN ratio when MnO₂ content increases. However, while there is a pronounced east-west gradient in the



Fe(OH)₃ distribution, the PHoN/PYN ratio (Fig. S6 in supplementary) is low (<0.6) in Labrador Sea surface waters (station #69), the Irminger Sea (stations #44 and #51) and from the surface to 750 m in the NADR region (stations #21, #26 and #32). These locations are depleted in both MnO₂ and Fe(OH)₃ (Fig. S6 in supplementary), leading to a weak adsorption of Ho. All along the section, low ratios are observed at the surface and until 800 m in the areas of marked biological productivity (stations #26, #32, #69), although they are directly linked to primary production intensity. This suggests a preferential scavenging of Y during the formation of biogenic matter, as reported by Censi et al. (2007), and not through simple adsorption which would support preferential scavenging of Ho. In the NADR region, at depths between 200 m and 600 m that are characterized by strong and positive PCe anomaly and a persistent PHREE enrichment, a PHo depletion is observed at stations #26 and #21. The low remineralization rates observed in this area could make persistent the PY enrichment formed at the surface, when the positive PCe anomaly suggests that a dynamic scavenging occurs. The difference between those two elements can be explained by a preferential scavenging of LREE compared to HREE during adsorption processes. The PHo enrichment at station #32 between 350 m and 600 m goes with the most important Ce positive anomaly, indicating intensive exchanges by adsorption with the dissolved phase, and then a stronger scavenging of REEs. In the ARCT region, slightly lower ratios are observed at station #69 than for others, and this station is characterized by a lower primary production and the higher remineralization rates along the section (Lemaitre et al., 2018). Ho, more adsorbed than Y is then more prone to be released, leading to a lower signal due to the Y enrichment of the covalent part of the particles, less easily remineralized. The higher ratios at other ARCT stations indicate scavenging by particles, although the Ce anomaly is lower than in the NADR region. Thus, our results show that high PHoN/PYN ratio not only reflects sorption processes driven by the occurrence of Fe(OH)₃ and to a lesser extent MnO₂ but all processes favoring outer-sphere complex formation, which would promote Ho and other REEs scavenging compared to Y. The relationship with particle production, particle residence time and remineralization rates is not clear indicating the influence of other parameters not identified yet.

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Table 1: List of region, water masse and current abbreviations.

Regions	
SPNA	Subpolar North Atlantic
NAST	North Atlantic Subtropical
NADR	North Atlantic drift
ARCT	Arctic
Water masses	
ENACW	East North Atlantic Central Water
MW	Mediterranean Water
SPMW	Subpolar Mode Water
IrSPMW	Irminger Subpolar Mode Water
LSW	Irminger Subpolar Mode Water
Currents	
NAC	North Atlantic Current
ERRC	East Reykjanes Ridge Current
IC	Irminger Current
EGIC	East Greenland Irminger Current
EGCC	East Greenland Coastal Current

620 **Table 2: Particulate REE, Y, Ba and ²³²Th concentrations with the corresponding 2σ error.**



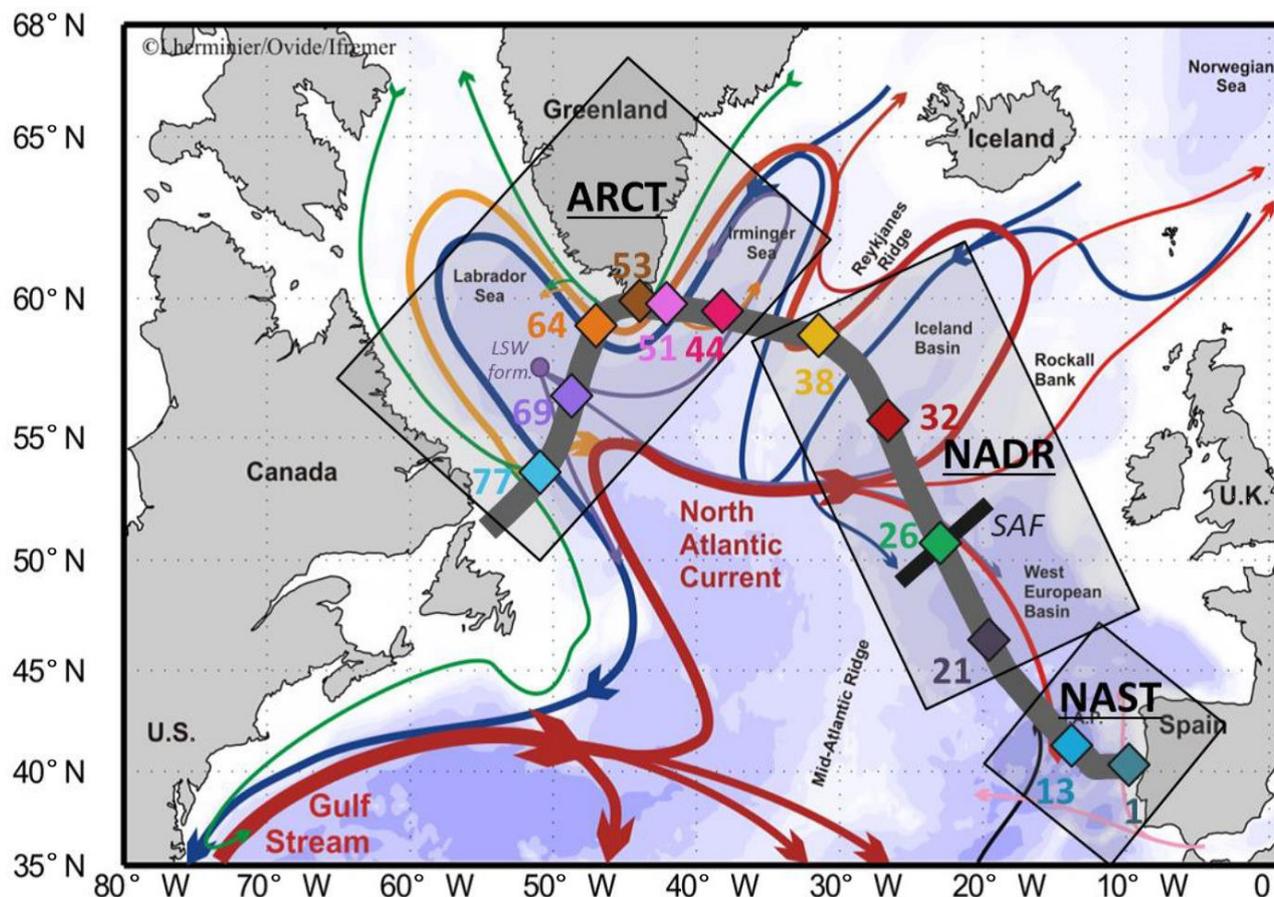
Station	Longitude	Latitude	Depth [m]	La	Ce	Co	Pr	Nd	Sm	Eu	Gd	Tb	Dy	Ho	Er	Tm	Yb	Lu	Y	Ba	Th	2σ	
1	-10.0359	40.333	20	962	324	1997	476	216	510	1231	1231	1231	1231	1231	1231	1231	1231	1231	1231	1231	1231	1231	
1	-10.0359	40.333	40	1730	377	2904	635	367	85	1377	326	333	77	60	13	238	53	35	8	192	44	36	8
1	-10.0359	40.333	60	596	133	1101	245	61	15	218	51	47	12	11	3	44	10	7	2	41	21	5	2
1	-10.0359	40.333	79	1014	214	1497	317	155	34	510	111	112	30	22	5	95	21	31	7	71	16	14	3
1	-10.0359	40.333	98	1552	365	2848	657	289	77	1061	252	222	63	44	10	153	40	20	5	116	30	22	5
1	-10.0359	40.333	119	1248	320	2649	677	273	78	955	267	230	68	37	9	147	37	20	5	185	16	8	2
1	-10.0359	40.333	139	1303	317	3179	773	260	64	963	234	214	54	36	9	147	37	20	5	185	16	8	2
1	-10.0359	40.333	160	1466	384	3938	1015	317	91	1123	299	228	73	46	12	172	48	22	6	126	34	23	6
1	-10.0359	40.333	200	2591	579	6041	1335	578	131	2090	470	362	86	77	17	282	66	38	9	104	24	14	3
1	-10.0359	40.333	249	3458	935	7935	2045	769	213	2815	724	535	158	105	27	392	100	57	16	286	74	38	8
1	-10.0359	40.333	299	3735	970	8831	2298	904	259	3001	850	445	130	109	28	401	103	53	15	314	77	53	14
1	-10.0359	40.333	309	2553	654	5692	1421	602	167	2139	644	626	161	77	19	308	76	41	11	234	68	43	11
1	-10.0359	40.333	499	2821	726	6219	1590	654	196	2426	698	483	145	90	24	338	107	41	13	253	79	46	11
1	-10.0359	40.333	597	3392	817	7103	1706	733	179	2593	629	436	131	97	24	330	88	41	10	127	32	18	5
1	-10.0359	40.333	702	5732	1400	9612	2346	1228	301	4505	1113	826	217	184	45	656	177	103	26	610	148	121	29
1	-10.0359	40.333	800	2554	610	5303	1264	558	156	1979	477	366	95	70	17	239	63	36	9	200	48	37	9
1	-10.0359	40.333	1000	2558	601	5145	1201	557	132	1994	471	378	98	90	21	276	65	41	11	222	54	41	10
1	-10.0359	40.333	1505	3644	961	7402	1916	844	240	2849	744	497	134	71	18	286	78	41	12	221	57	40	10
13	-13.8877	41.383	20	1381	462	3330	1102	290	101	1113	365	422	137	61	21	182	59	26	9	138	45	26	8
13	-13.8877	41.383	40	1219	380	1035	324	187	58	785	246	202	65	57	18	254	80	50	16	403	125	106	33
13	-13.8877	41.383	60	999	313	895	279	125	40	414	131	91	22	23	8	99	31	18	6	109	35	26	8
13	-13.8877	41.383	78	918	285	1467	456	135	42	500	157	101	34	23	7	100	31	21	7	104	33	24	7
13	-13.8877	41.383	100	612	187	1004	308	117	36	429	133	88	31	20	6	91	31	13	4	81	25	17	5
13	-13.8877	41.383	120	853	262	1375	423	122	38	460	142	82	28	20	7	88	27	12	4	79	26	16	5
13	-13.8877	41.383	140	775	237	1356	416	147	46	541	166	116	58	25	8	98	31	14	5	82	25	17	5
13	-13.8877	41.383	160	1258	395	3917	1226	225	823	259	161	52	35	11	166	58	25	8	169	53	36	11	5
13	-13.8877	41.383	200	741	255	1075	359	142	48	557	188	117	44	15	9	54	34	18	6	54	20	8	5
13	-13.8877	41.383	250	1418	482	2443	822	244	88	1029	557	214	77	67	23	178	60	52	19	145	51	30	10
13	-13.8877	41.383	300	1833	592	2871	924	279	91	895	288	102	52	32	17	110	35	18	4	78	27	10	3
13	-13.8877	41.383	400	979	299	1610	490	209	731	224	139	48	30	10	128	40	18	6	102	32	19	6	5
13	-13.8877	41.383	500	964	323	2325	769	202	71	777	255	96	43	14	127	41	18	6	96	31	19	6	5
13	-13.8877	41.383	600	774	251	1113	675	160	56	900	280	126	47	25	8	107	34	14	5	88	29	17	5
13	-13.8877	41.383	700	806	253	2070	647	165	52	597	188	119	39	26	8	113	37	16	5	90	29	18	6
13	-13.8877	41.383	800	917	280	2192	669	201	63	741	227	145	47	30	9	125	39	17	6	98	31	19	6
13	-13.8877	41.383	1000	1076	327	2356	780	205	72	765	258	149	52	43	14	129	42	18	6	97	34	18	6
13	-13.8877	41.383	1501	1026	353	2379	811	209	72	765	268	149	52	43	14	142	51	22	8	94	32	18	6
21	-19.6724	46.544	10	978	244	1188	290	153	37	555	137	108	30	7	115	28	20	5	123	31	28	7	94
21	-19.6724	46.544	20	3837	2397	15279	1896	260	68	862	224	167	51	14	214	63	43	12	6	69	18	14	3
21	-19.6724	46.544	40	756	199	812	199	89	22	325	82	68	18	17	4	69	18	14	3	42	11	6	2
21	-19.6724	46.544	60	962	241	1433	358	195	51	764	189	147	43	26	7	148	38	23	6	147	39	29	7
21	-19.6724	46.544	76	988	257	1824	471	171	46	652	174	122	32	32	8	118	31	16	5	97	27	20	5
21	-19.6724	46.544	100	9587	2397	15279	1896	260	68	862	224	167	51	14	214	63	43	12	6	69	18	14	3
21	-19.6724	46.544	120	787	191	1312	318	162	41	681	165	125	37	32	8	110	26	16	4	93	24	17	4
21	-19.6724	46.544	140	1050	256	2456	605	195	49	745	191	114	32	34	9	123	30	17	5	90	24	17	4
21	-19.6724	46.544	160	404	98	774	187	88	23	327	80	77	19	28	7	69	16	10	3	57	15	11	3
21	-19.6724	46.544	200	634	160	1071	266	133	34	522	133	111	29	28	7	99	25	14	4	78	21	15	4
21	-19.6724	46.544	300	751	182	2121	509	146	37	567	138	113	30	33	8	110	27	16	4	90	24	17	4
21	-19.6724	46.544	400	1065	211	1659	406	173	45	648	166	136	34	39	10	111	27	35	10	90	26	17	4
21	-19.6724	46.544	500	821	141	1659	406	173	45	648	166	136	34	39	10	111	27	35	10	90	26	17	4
21	-19.6724	46.544	600	1681	410	3232	773	222	54	737	182	123	23	23	6	88	22	15	4	73	19	15	4
21	-19.6724	46.544	700	1085	170	2095	514	280	71	1152	284	463	115	64	16	184	45	27	7	139	38	25	6
21	-19.6724	46.544	800	807	197	2185	527	138	35	524	127	113	31	30	8	97	24	13	4	79	20	15	4
21	-19.6724	46.544	1000	1009	251	1947	478	260	66	1029	266	215	58	59	15	171	42	25	7	129	35	23	6
21	-19.6724	46.544	1200	533	131	1359	334	120	31	446	111	89	23	20	5	81	20	11	3	37	9	5	1
26	-22.6046	50.278	20	986	301	642	194	163	50	610	185	126	43	28	9	127	40	18	6	1			



Station	Longitude	Latitude	Depth [m]	La	Ce	Pr	Nd	Sm	Eu	Gd	Tb	Dy	Ho	Er	Tm	Yb	Lu	Y	Ba	Th	2σ
32	-26.7030	55.515	40	1142	271	178	200	51	726	188	54	25	90	23	13	3	91	21	14	3	1632
32	-26.7030	55.515	60	593	145	398	96	112	294	108	85	42	79	21	16	4	48	13	7	2	851
32	-26.7030	55.515	80	827	113	606	153	144	40	545	111	119	38	27	4	43	12	6	2	1130	
32	-26.7030	55.515	100	627	150	362	85	119	30	426	101	90	25	21	5	85	20	14	1	860	
32	-26.7030	55.515	120	558	140	403	100	89	23	337	85	75	20	18	5	71	18	9	1	762	
32	-26.7030	55.515	140	381	92	2518	602	70	17	273	63	2	44	11	3	56	13	8	1	691	
32	-26.7030	55.515	160	649	158	896	218	89	22	335	84	70	18	31	8	57	14	5	1	574	
32	-26.7030	55.515	200	312	72	432	99	50	12	192	45	7	21	6	3	1	20	5	2	1	460
32	-26.7030	55.515	300	589	140	1143	269	101	25	375	90	47	11	3	9	1	28	7	3	1	679
32	-26.7030	55.515	380	543	131	975	235	90	23	347	87	68	19	11	3	31	9	4	1	1	556
32	-26.7030	55.515	450	11425	2812	16272	4695	140	35	428	108	91	22	24	6	99	25	10	3	1	596
32	-26.7030	55.515	500	545	132	1366	328	108	27	418	103	89	24	23	6	81	20	13	3	1	612
32	-26.7030	55.515	598	470	111	1201	287	102	25	390	93	72	21	5	70	17	10	3	4	1	674
32	-26.7030	55.515	700	531	141	1427	373	115	34	418	100	78	21	5	1	29	7	4	1	1	705
32	-26.7030	55.515	800	578	144	1565	386	115	30	423	104	80	23	7	79	19	11	3	4	1	590
32	-26.7030	55.515	1000	559	138	1469	365	114	29	442	116	72	33	5	1	84	6	7	3	1	846
38	-31.2660	58.845	10	4901	1132	6127	1419	372	87	1393	328	47	58	14	12	3	76	18	9	2	2166
38	-31.2660	58.845	20	2725	607	1704	379	504	112	2535	564	555	79	80	18	420	94	48	11	271	61
38	-31.2660	58.845	40	1667	398	1095	259	345	86	1276	303	244	64	54	13	217	55	30	8	169	44
38	-31.2660	58.845	60	609	145	358	90	125	32	477	116	99	24	21	5	84	27	11	3	68	17
38	-31.2660	58.845	80	437	103	478	115	72	17	274	67	56	15	13	3	54	13	7	2	43	12
38	-31.2660	58.845	100	378	91	3169	751	68	17	252	61	51	14	13	3	50	12	7	2	40	11
38	-31.2660	58.845	120	410	101	345	133	68	17	252	65	56	16	15	4	53	13	7	2	42	12
38	-31.2660	58.845	140	454	109	565	135	68	16	262	64	50	15	13	3	53	12	8	2	46	11
38	-31.2660	58.845	160	751	183	1144	275	108	27	417	105	79	21	23	6	77	19	11	3	62	17
38	-31.2660	58.845	180	492	117	547	129	81	19	310	74	69	17	4	75	22	10	2	55	14	11
38	-31.2660	58.845	200	468	109	599	142	85	20	334	79	67	18	17	4	66	16	16	4	57	14
38	-31.2660	58.845	300	578	142	1022	251	120	31	477	121	102	28	37	4	98	24	15	4	82	22
38	-31.2660	58.845	400	865	214	1615	398	197	50	716	198	146	41	51	13	154	39	23	6	132	34
38	-31.2660	58.845	500	834	202	1652	401	200	50	862	202	185	49	52	15	197	52	26	7	78	20
38	-31.2660	58.845	600	1145	274	2292	544	241	61	980	247	205	59	70	18	203	49	32	8	175	45
38	-31.2660	58.845	700	951	218	1782	423	196	46	1092	239	57	45	10	160	37	25	6	74	19	9
38	-31.2660	58.845	800	2191	530	3086	1249	340	89	1824	288	151	43	38	9	148	36	27	124	32	6
38	-31.2660	58.845	1000	896	216	1774	421	238	60	969	245	208	54	56	13	208	50	32	9	148	36
44	-38.954	59.62	10	719	169	592	136	121	29	445	105	89	22	25	6	115	26	25	6	129	36
44	-38.954	59.62	20	823	191	477	110	131	32	564	145	97	25	29	7	127	30	21	5	135	38
44	-38.954	59.62	40	545	128	476	111	109	26	436	101	94	22	25	6	101	23	20	5	92	22
44	-38.954	59.62	60	792	184	799	186	107	25	386	90	86	20	20	5	87	20	12	3	75	17
44	-38.954	59.62	80	394	87	911	202	80	18	293	65	64	14	15	3	60	13	8	2	49	12
44	-38.954	59.62	100	456	110	716	174	93	23	336	82	64	16	20	7	65	14	9	2	55	14
44	-38.954	59.62	120	384	84	1813	395	84	20	307	71	65	15	15	3	63	14	9	2	54	13
44	-38.954	59.62	140	402	93	608	140	85	20	307	71	65	15	15	3	63	14	9	2	54	13
44	-38.954	59.62	160	1739	409	3254	765	122	29	402	95	73	17	18	4	78	18	n.d.	n.d.	n.d.	n.d.
44	-38.954	59.62	200	665	145	1155	253	140	31	488	108	95	22	18	4	84	19	12	3	64	14
44	-38.954	59.62	300	505	112	1082	239	112	25	395	87	82	19	21	5	76	17	12	3	64	14
44	-38.954	59.62	400	504	112	918	204	111	25	401	88	88	19	21	5	83	20	11	3	62	14
44	-38.954	59.62	500	603	139	1205	276	128	30	452	104	84	20	26	6	81	18	n.d.	n.d.	n.d.	n.d.
44	-38.954	59.62	600	577	128	1147	254	127	29	452	100	90	20	19	4	84	20	14	3	62	14
44	-38.954	59.62	700	473	109	825	190	96	23	336	78	72	17	19	5	67	16	9	2	50	12
44	-38.954	59.62	800	259	60	471	109	55	13	194	45	43	10	10	2	36	9	2	28	7	5
44	-38.954	59.62	1000	573	126	1394	306	125	28	431	96	86	16	16	4	84	16	12	3	64	14
44	-38.954	59.62	1401	901	211	1848	436	142	34	511	126	108	25	46	16	98	23	13	3	78	18
51	-42.013	59.8	10	587	137	415	98	102	24	376	88	84	19	20	5	95	22	10	25	22	5
51	-42.013	59.8	19	415	90	518	110	57	13	212	46	45	10	11	3	47	10	8	2	57	14
51	-42.013	59.8	40	607	143	497	118	101	24	371	88	81	19	20	5	88	21	13	3	85	22
51	-42.013	59.8	60	636	145	490	113	100	25	497	121	84	19	20	5	81	24	12	3	82	17
51	-42.013	59.8	80	653	148	1497	340	110	29	421	115	91	61	25	6	84	19	12	3	38	10
51	-42.013	59.8	100	726	162	1113	253	131	30	460	105	91	20	22	5	88	21	17	4	72	17
51	-42.013	59.8	120	574	133	1387	320	145	34	524	104	24	31	7	103	23	14	3	86	20	16
51	-42.013	59.8	140	664	174	1444	330	156	37	555	127	113	26	26	6	106	26	13	3	81	20
51	-42.013	59.8	160	767	155	1330	312	151	35	533	123	103	24	25	6	105	25	14	3	75	18
51	-42.013	59.8	180	925	209	1833	411	206	47	728	163	143	32	34	8	126	28	17	4	101	24
51	-42.013	59.8	200	1105	259	2306	539	247	59	888	223	172	45	45	12	201	61	25	7	131	37
51	-42.013	59.8	300	1358	295	2470	539	142	32	539	189	22	29	9	101	22	15	3	86	19	7
51	-42.013	59.8	400	1045	246	2107	494	231	54	826	193	165	38	41	10	165	39	21	5	131	31
51	-42.013	59.8	500	1061	229	2598	562	279	62	1029	223	208	45	54	12	192	46	27	6	139	35

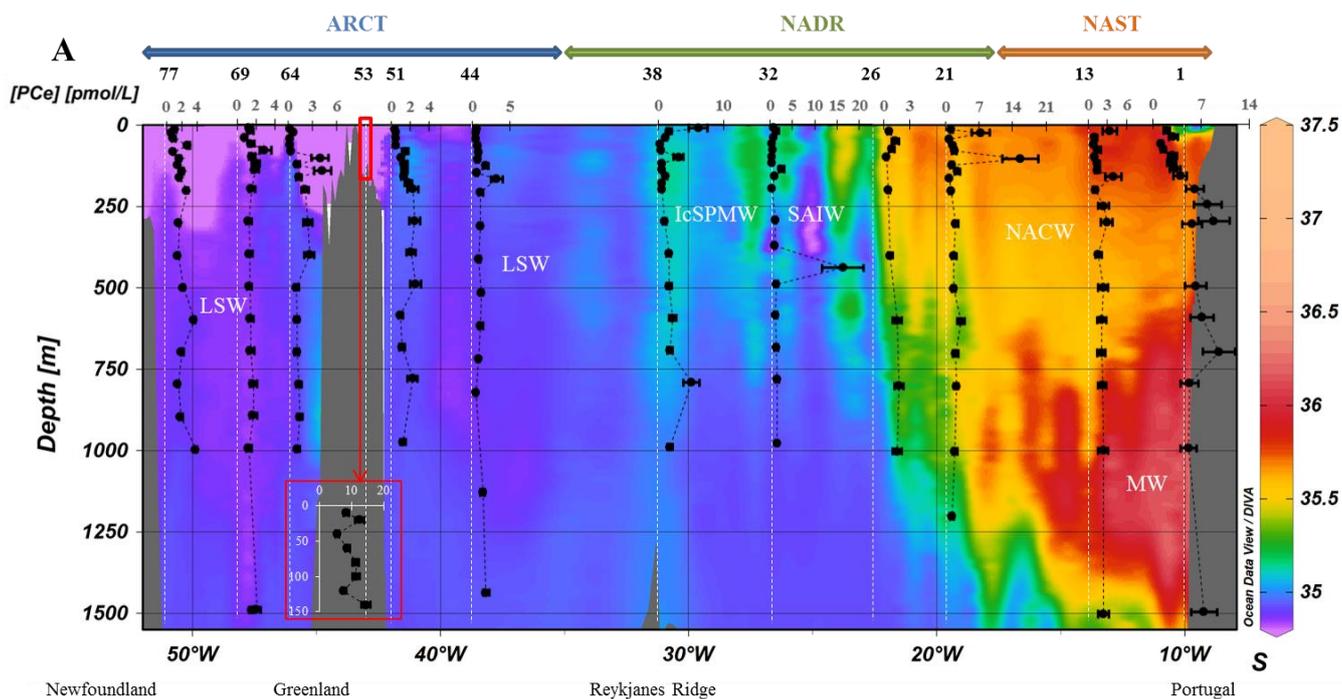


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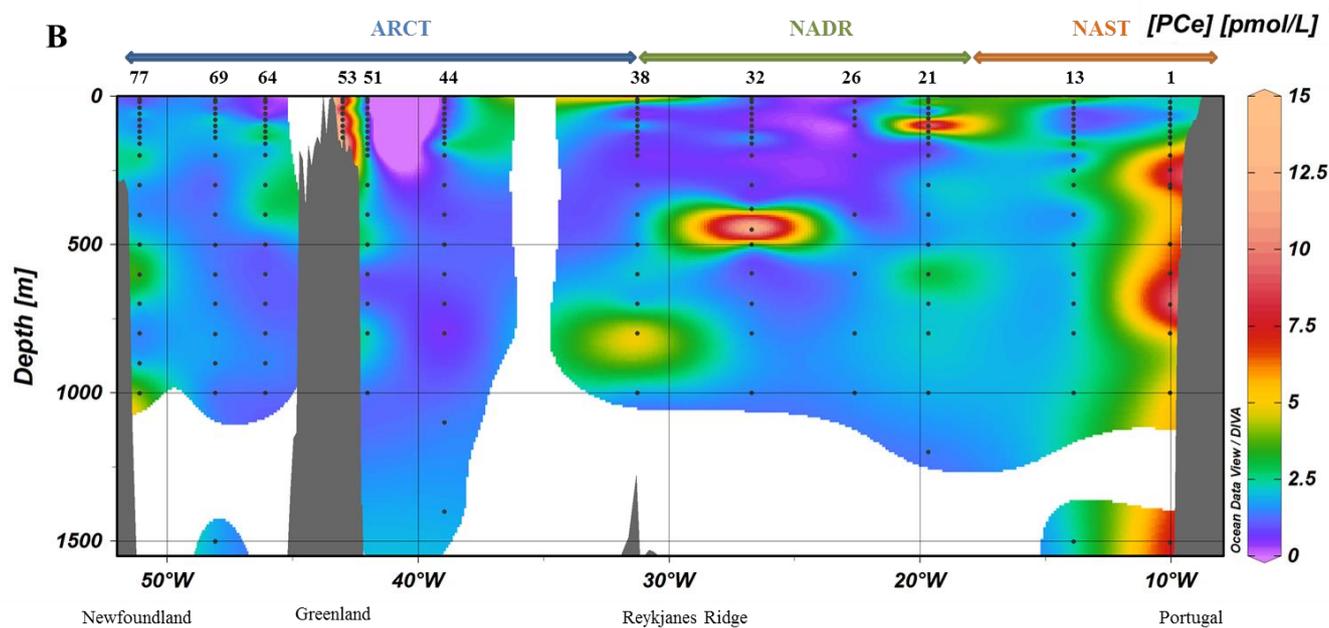


630 **Figure 1:** Map of the studied area (Subpolar North Atlantic, SPNA), including schematized circulation features, adapted from
García-Ibáñez et al. (2015). Bathymetry is plotted in color with interval boundaries at 100 m, at 1000 m, and every 1000 m below
635 1000 m. The red and green arrows represent the main surface currents, the pink and orange arrows represent currents at
intermediate depths, and the blue and purple arrows represent the deep currents. Diamonds indicate station positions, located in 3
distinct areas (grey squares): the North Atlantic Subtropical province (NAST), the North Atlantic Drift region (NADR), and the
Arctic region (ARCT). The approximate locations of the subarctic front (SAF; black bar crossing station #26) and the formation site
of the Labrador Sea Water (LSW form.) are indicated. The section used in ODV figures is symbolized by the thick grey line. From
Lemaitre et al. (2018).

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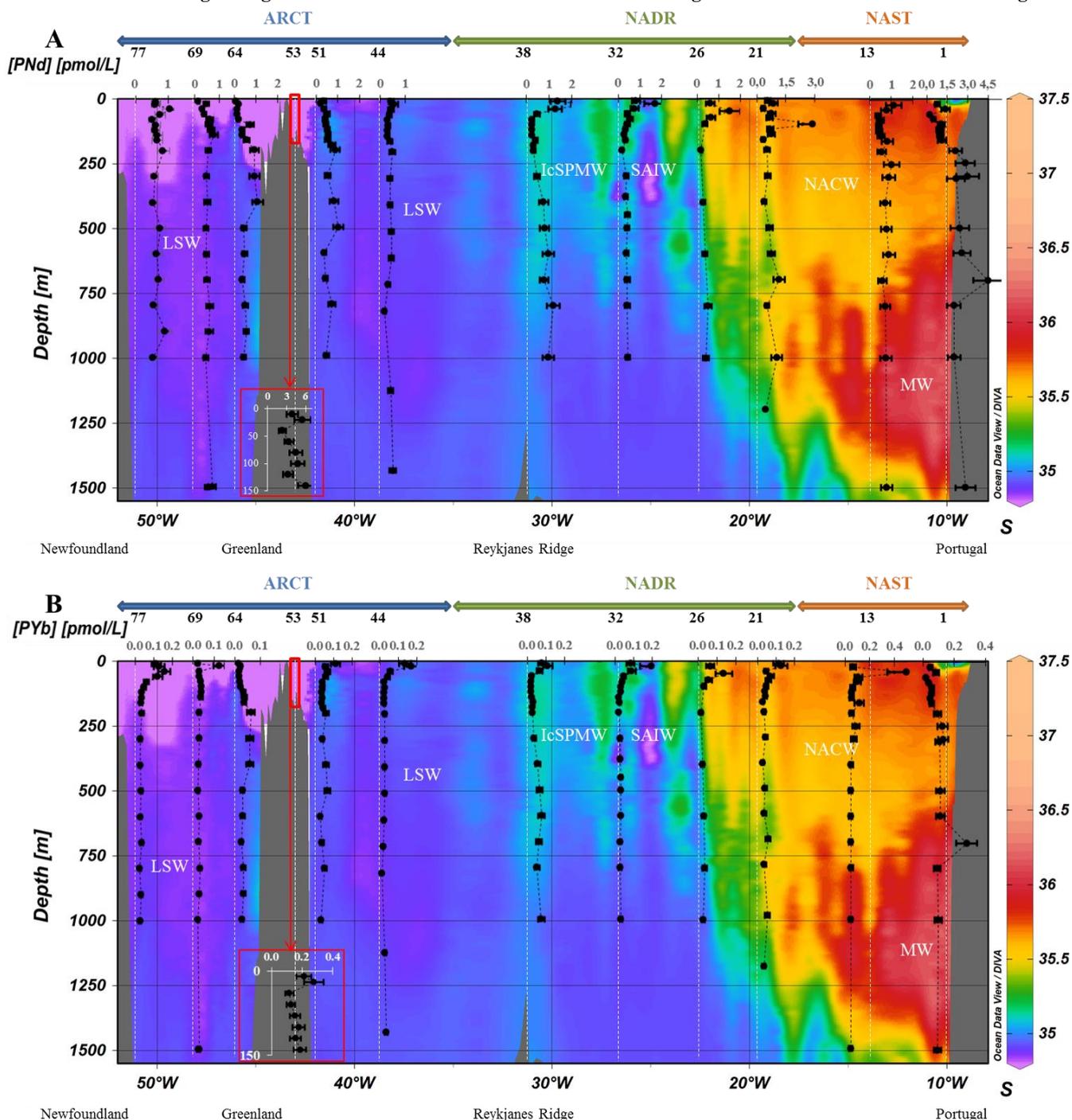
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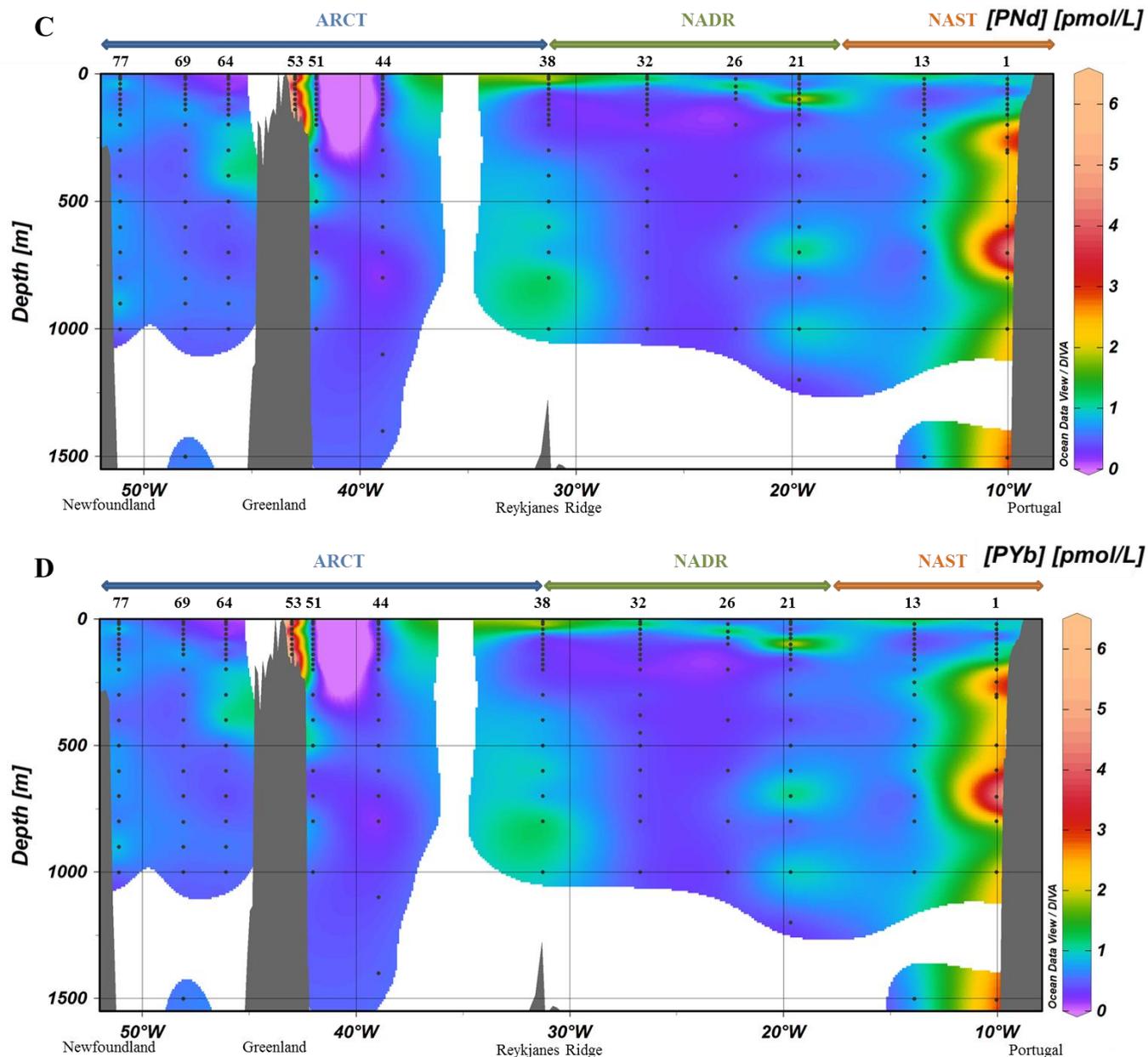




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Figure 2: A. Profiles of particulate [Ce] concentrations superimposed on salinity (S) measured by CTD at every GEOVIDE station (Lherminier and Sarthou, 2017); in white, water masses characterized by a multiparametric (OMP) analysis (García-Ibáñez et al., 2018). For the station #53, profiles are shifted to the bottom at a lower scale because of the shallow depth of the station. This map and the following were created with the software Ocean Data View (Schlitzer, 2016). B. Particulate [Ce] concentrations interpolated with the DIVA gridding function of Ocean Data View along the section defined in Fig. 1.





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Figure 3: A. Profiles of particulate [Nd] and B. [Yb] concentrations superimposed on salinity (S) measured by CTD at every GEOVIDE station (Lherminier and Sarthou, 2017); in white, water masses characterized by OMP analysis (García-Ibáñez et al., 2018). At station #53, profiles are shifted to the bottom at a lower scale because of the shallow depth of the station. C. Particulate [Nd] and D. [Yb] concentrations interpolated with the DIVA gridding function of Ocean Data View along the section defined in Fig. 1.

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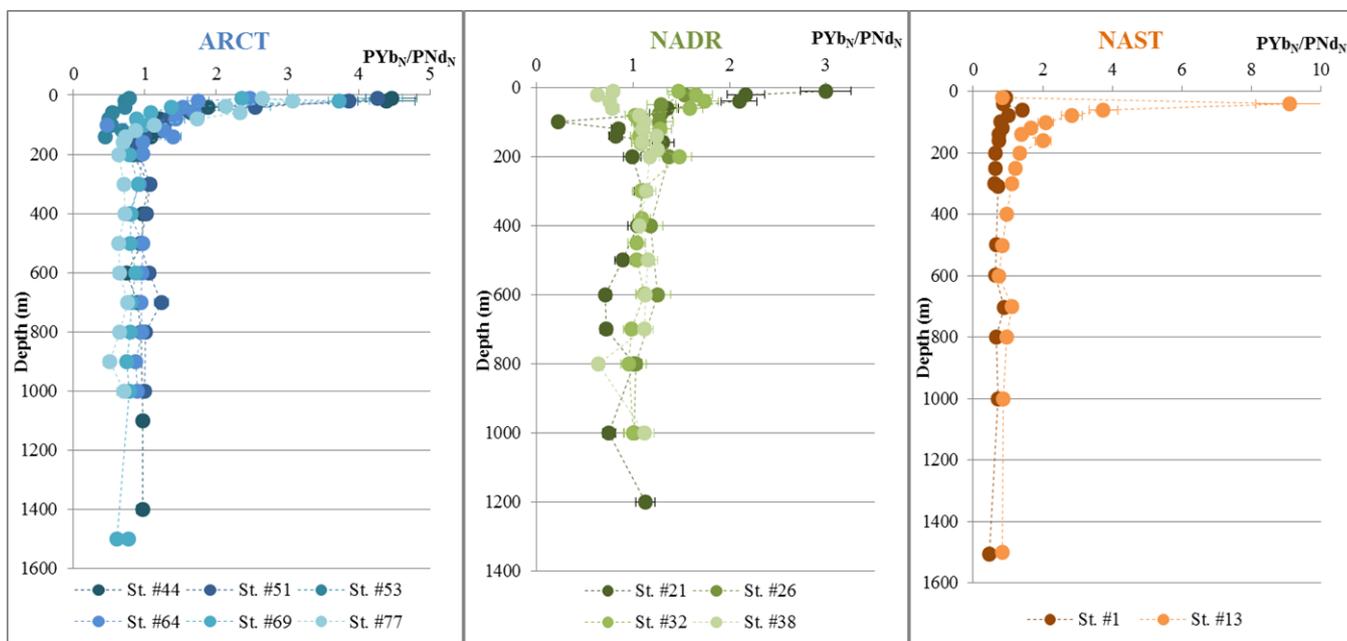
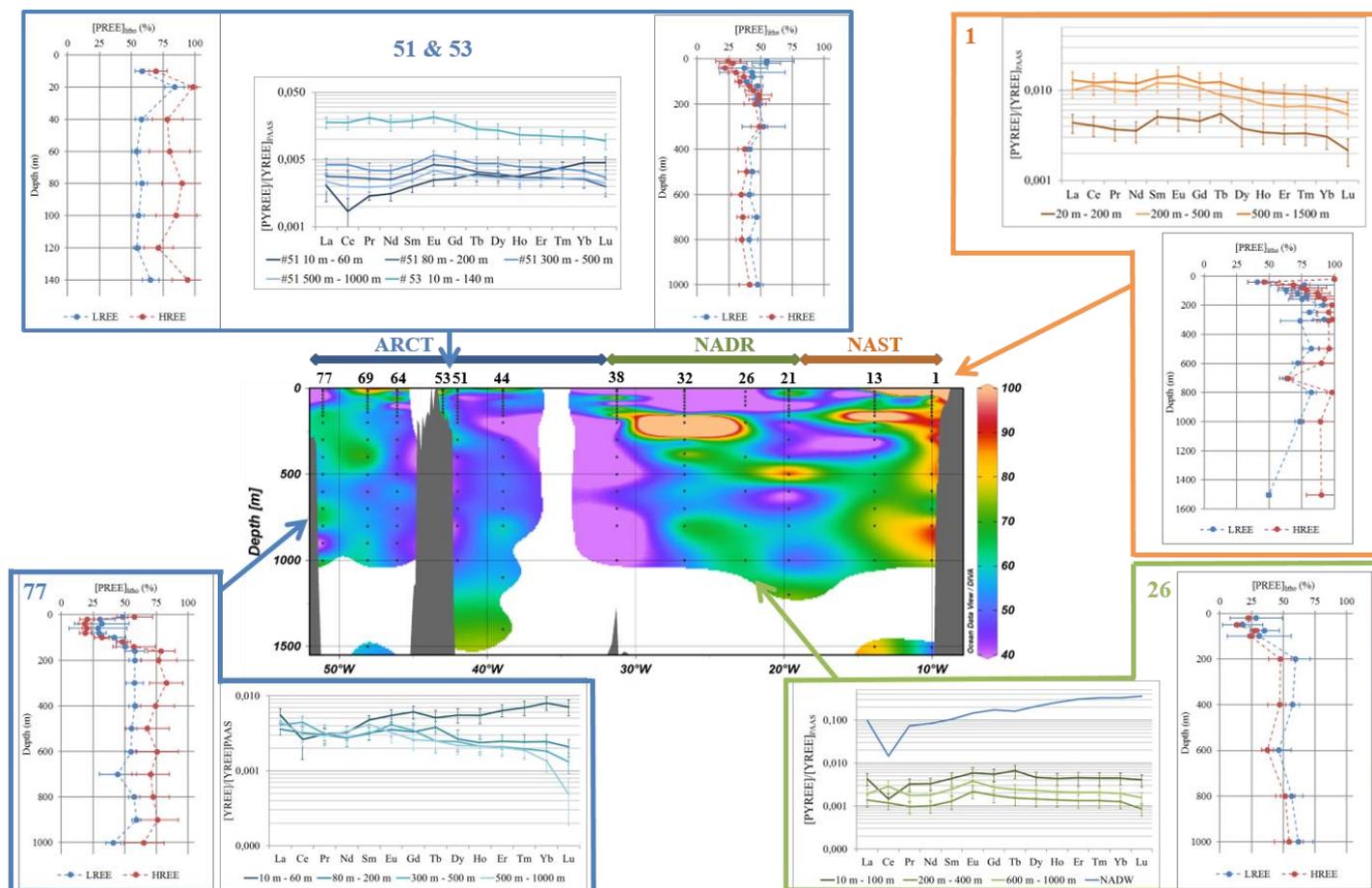
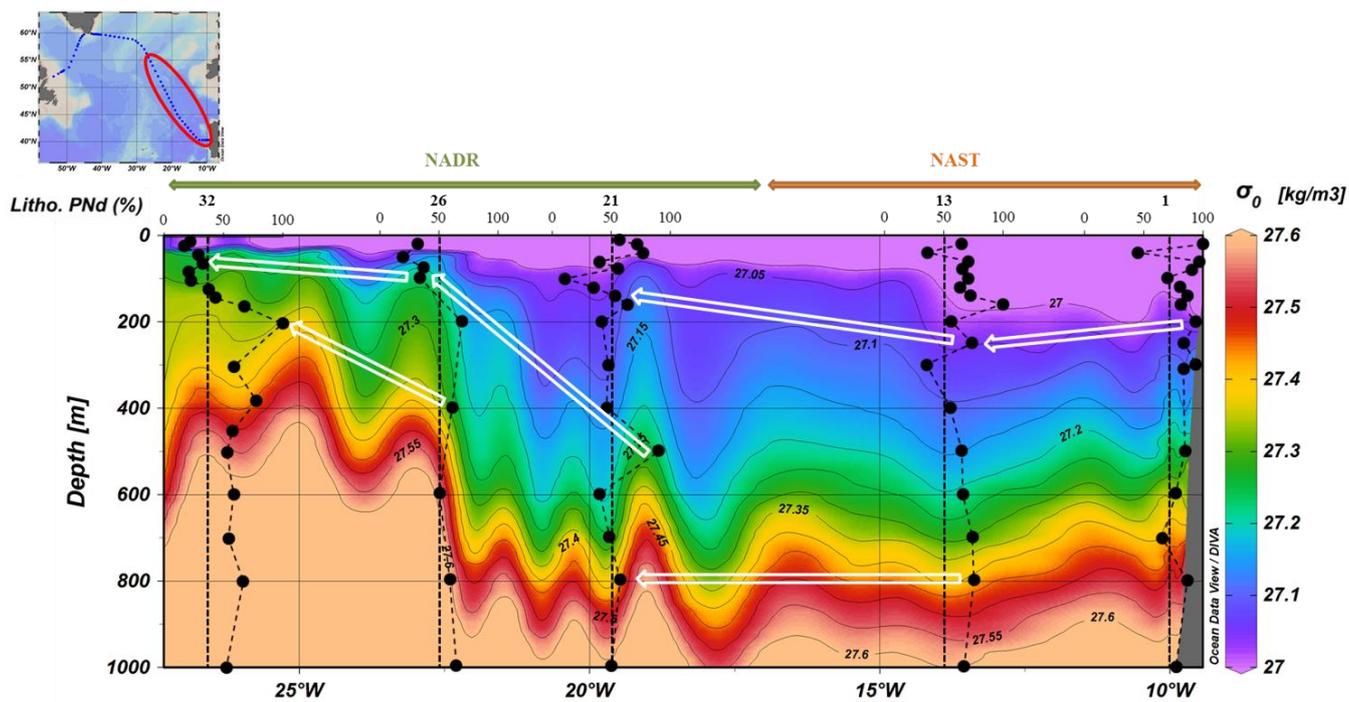


Figure 4: PYb/PNd ratio normalized to PAAS in each biogeochemical province

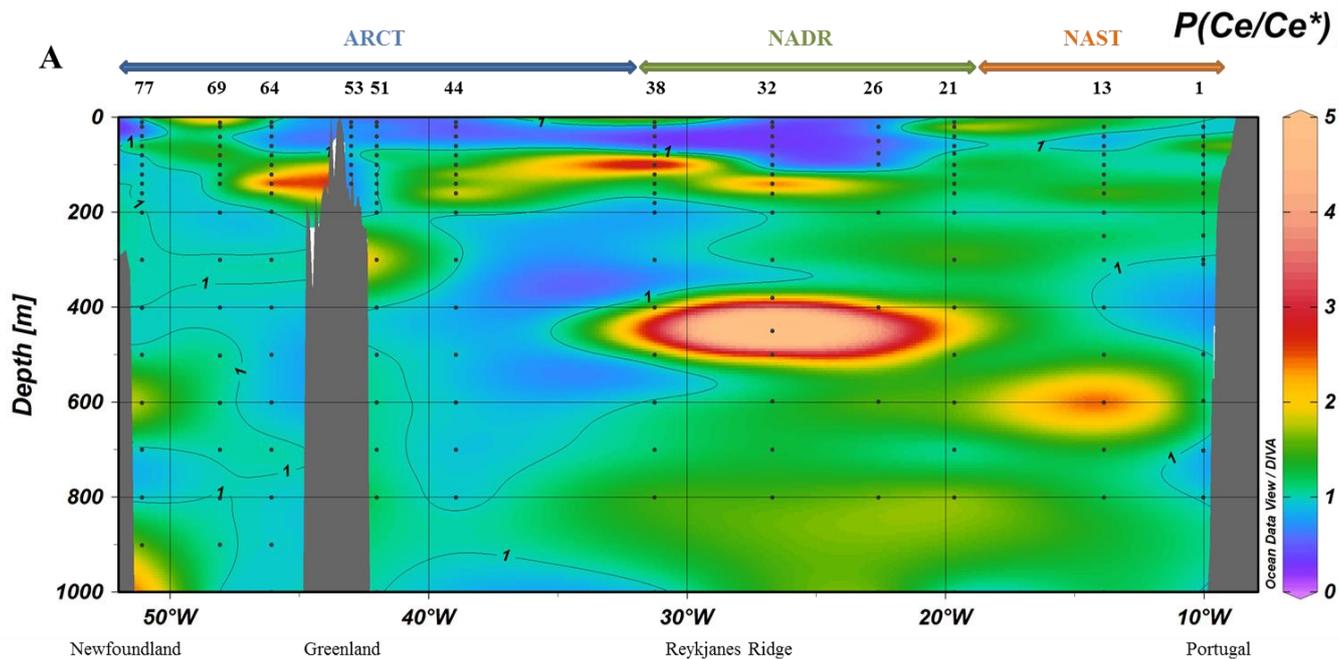


665 **Figure 5:** Center: proportion of lithogenic PNd along the GEOVIDE section (in %); Borders: vertical profiles of the lithogenic fraction of LREEs (except Ce, blue lines) and HREEs (red lines) and PAAS-normalized REE patterns of the total fraction, averaged by depth layers, at stations #1, #26, #51, #53 and #77.



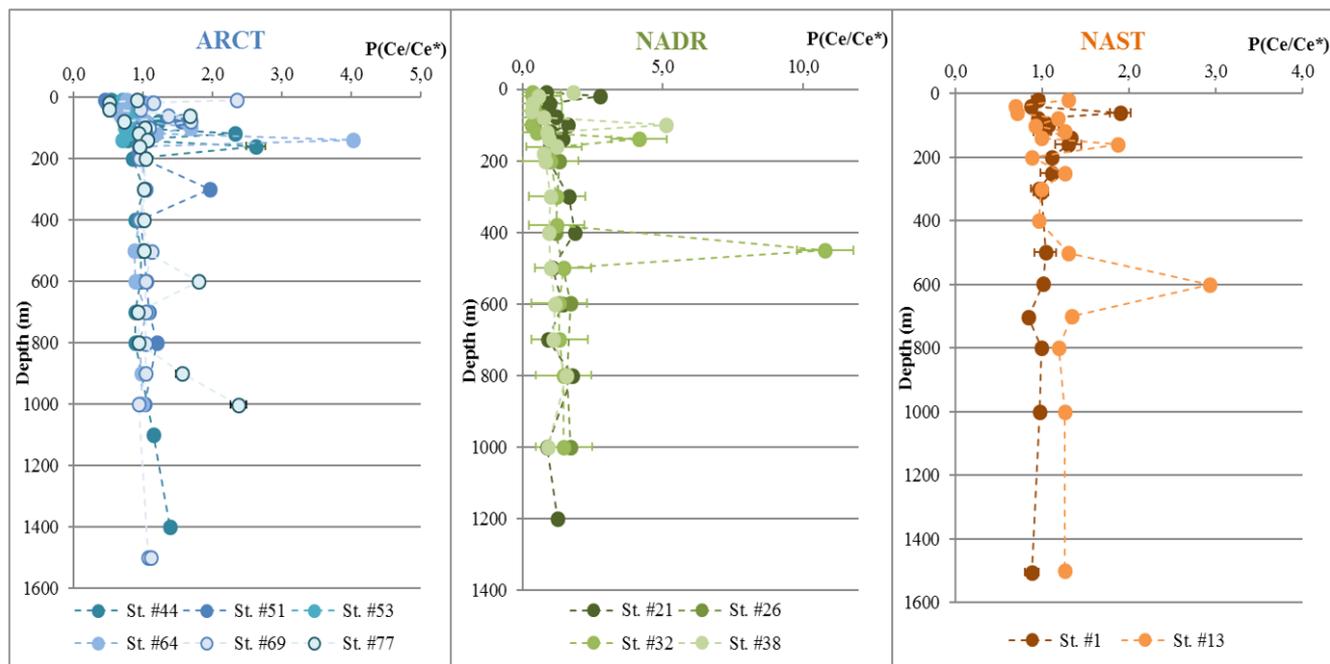
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Figure 6: Proportion of lithogenic PNd profiles superposed to density from station #1 to #32

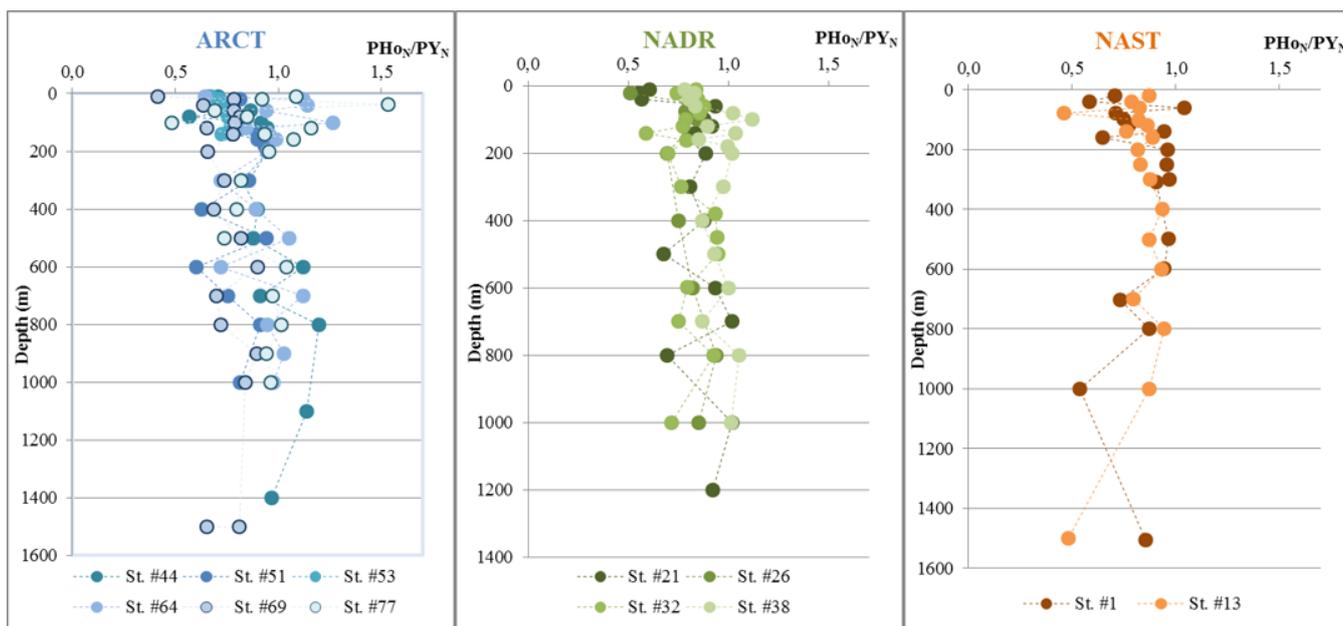




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675 **Figure 7: A. Particulate Ce anomaly (Ce/Ce^*) along the GEOVIDE section, interpolated with the DIVA gridding function of Ocean Data View and B. Ce/Ce^* profiles by biogeochimical provinces.**



680 **Figure 8: PAAS-normalized PHo/PY profiles by biogeochimical provinces.**

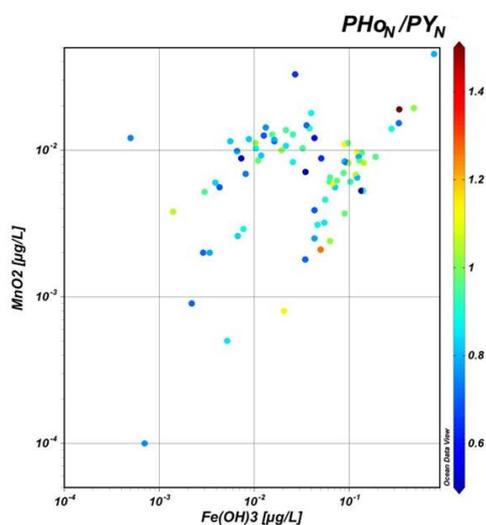


Figure 9: PAAS-normalized PHO/PY ratio as a function of Fe(OH)₃ and MnO₂ concentrations (in μg.L⁻¹).

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Author contribution

N.L. did the sampling during the cruise, helped by C.J. N.L. did the leaching on the filter and first Ba measurements. C.J., M.B., M.G. and M.L. did REE measurements. ML wrote the manuscript, and C.J., H.P., M.G., N.L. and P.L. did the proofreading.

Competing interests

690 The authors declare that they have no conflict of interest.

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