The distribution of ²²⁷Ac along the GA01 section in the North Atlantic

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Abstract :

Actinium-227 is a powerful tool to study vertical mixing in the ocean, and was more recently proposed as a tracer of hydrothermal plumes. However, because 227Ac activities are especially low in the ocean, relatively few studies have been conducted to date, and none has been done on the large scale, as is reported in the present study. Here, we report a section of dissolved 227Ac in the North Atlantic Ocean between Portugal-Greenland-Canada based on nine full-depth profiles obtained in the framework of the international GEOTRACES program (GA01 section - GEOVIDE cruise, May–July 2014). The simultaneous determination of 231Pa, the parent nuclide of 227Ac, along the section allowed us to report a section of excess 227Ac activities (227Acex).

Actinium-227 activities were especially low along the section (<0.4 dpm m-3) compared to other ocean basins. In most cases, 227Ac activities reached secular equilibrium with 231Pa in the water column 500 m above the seafloor. Secular equilibrium is considered reached after ~five half-lives of the daughter. Secular equilibrium therefore suggests no external input over the last ~100 years. The highest 227Ac activities were often found close to the seafloor, indicating that 227Ac diffuses out of the sediments. However, two different patterns question the traditional one-dimensional vertical mixing model typically applied to 227Ac released by deep-sea sediments: first, significant 227Acex were found in the upper 500 m of the water column at several stations located near the Iberian margin. In the upper 500 m, the high 227Acex activities could indicate lateral advection of waters that interacted with the margins. Second, mid-water peaks of 227Acex were occasionally observed along the transect. At station 44 in the Irminger Basin, a peak of 227Acex activity at 2500 m that was co-located with high dissolved Fe concentrations could be interpreted as the signature of a hydrothermal plume. Near the seafloor, we often observed bottom nepheloid layers and we cannot exclude that these layers impact the 227Ac distributions.

Highlights

► Actinium-227 activities were especially low along the section (<0.4 dpm m⁻³) compared to other ocean basins. ► In most cases, ²²⁷Ac activities reached secular equilibrium with ²³¹Pa (suggesting no external input over the last ~100 years) in the water column 500 m above the seafloor. ► The highest ²²⁷Ac activities were often found close to the seafloor, indicating that ²²⁷Ac diffuses out of the sediments. ► In the upper 500 m, high ²²⁷Ac_{ex} activities could indicate lateral advection of waters that interacted with the margins. ► Near the seafloor, we often observed bottom nepheloid layers and we cannot exclude that these layers impact the ²²⁷Ac distributions, since manganese oxides were shown to be present in these layers and may thus scavenge ²²⁷Ac among other radionuclides.+

Keywords : GEOTRACES, Actinium, Actinium-227, North Atlantic

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1. Introduction

Actinium-227 (²²⁷Ac) is a deep-sea mixing tracer that combines the advantages of a source in the deep-sea sediments with a half-life of 21.8 y. ²²⁷Ac is produced by the decay of its parent nuclide protactinium-231 (²³¹Pa; $T_{1/2} = 32760$ y), ²³¹Pa being itself produced following the decay of uranium-235 (235 U; T_{1/2} = 7.04 10⁸ y). 235 U activities in the ocean arc mostly constant through space and time due to its long residence time (~0.5 Ma; Ku et al., 1977), which leads to a uniform production rate of ²³¹Pa in the water column. Once produced, ²³¹Pa rapidly acisetic onto particles. It is then transported to the seafloor and thus, accumulates slowly in the sediment, (Anderson et al., 1983). In the sediment, ²³¹Pa decays to ²²⁷Ac which is then released to the overly in water column due to its higher solubility (Anderson et al., 1983; Nozaki, 1984; Nozaki et al. 1990; Nozaki, 1993, p.199; Geibert et al., 2002; Kemnitz et al., 2022). ²²⁷Ac that diffuses out of the sediments is then redistributed in the ocean by transport (advection and upwelling) and mixin, (Nozaki, 1984; Geibert et al., 2002; Koch-Larrouy et al., 2015). The ²²⁷Ac that diffuses out of u.e sediment thus adds to the ²²⁷Ac produced in the water column (from ²³¹Pa) that is considered 10 of at secular equilibrium with ²³¹Pa, assuming steady-state conditions (Nozaki, 1984). Excess 227 Ac activities (227 Ac_{ex}, the difference between the total activities and ²²⁷Ac at secular equilibrium with ²³¹Pa) are thus often observed up to ca. 500 m above the seafloor due to vertical mixing (Geibert et 21, 2002; Nozaki, 1984). Nozaki (1984) proposed estimating vertical eddy diffusivity coefficients (X_{-}) from the ²²⁷Ac_{ex} vertical distribution. ²²⁷Ac_{ex} has also been used to estimate upwelling rates, since this process also brings ²²⁷Ac toward the surface (Geibert et al., 2002; Haskell et al., 2015; Kemnitz et al., 2022). Finally, Kipp et al. (2015) showed that hydrothermal vents constituted a source of ²²⁷Ac to the deep Atlantic Ocean that was equivalent to ~2-6% of the deep-sea sediment flux.

²²⁷Ac concentration in the ocean is, however, especially low. Its global oceanic inventory is only 37 mol or 8.4 kg (Geibert et al., 2008). In addition, interpretation of its distribution requires the knowledge of its ²³¹Pa parent distribution. As a result, the use of dissolved excess ²²⁷Ac as an oceanic tracer has been reported in few studies to date (Dulaiova et al., 2012; Geibert et al., 2008, 2002; Geibert and Vöge, 2008; Kipp et al., 2015; Le Roy et al., 2019; Koch-Larrouy et al., 2015; Nozaki, 1984, 1993; Nozaki et al., 1990; Levier et al., 2021; Kemnitz et al., 2022). In the present study, we report nine vertical profiles of ²²⁷Ac from a large-scale section in the North Atlantic Ocean and the Labrador Sea. 2018). The determination of both ²²⁷Ac and ²³¹Pa at the nine full-depth vertical profiles provides a unique section of $^{227}Ac_{ex}$ activities in the North Atlantic.

2. Material and Methods

2.1. GEOVIDE Cruise

The GEOVIDE section (GEOTRACES GA01; PIs: Géraldine Sarthou, LEMAR, France and Pascale Lherminier, LOPS, France) was conducted in the North Atlantic Ocean between Lisbon, Portugal and St John's, Newfoundland on board the R/V *Pourquoi Pas?* (15 May 2014-30 June 2014; Fig.1). The section crossed different topographic features and regions displaying contrasting biogeochemical patterns. It complements sections GA03 and GA? also conducted in the Atlantic Ocean in the framework of the GEOTRACES program.

2.2. Hydrodynamic Context

An Optimum Multiparameter Analysis (OMPA) was used to estimate the contributions of the different water masses found along the GEOVIDE section (García-Ibáñez et al., 2018). First, Central Waters, carried by the North Atlantic Current (NAC), oc cupy the upper eastern part of the GEOVIDE section from the Iberian Peninsula to the Roc'.a.' Lough (stations 1 to 26; Fig.2). Part of the Central Waters flows toward the Iceland Basin and the Irminger Sea and forms the Iceland Subpolar Mode Water (IcSPMW) in the Iceland Basin over the Reykjanes Ridges. Then, these waters follow the Irminger Current (Stations 32 and 38; Fig.2: Lacan and Jeandel, 2005; McCartney, 1992). Then, the Irminger Subpolar Mode Water (IrSTMW) results from the transformation of the Central Waters and the IcSPMW, northwest of the Inninger Sea (Krauss, 1995). The IrSPMW is located near Greenland (Fig.2) (García-Ibáñez et al., 2013; Lacan and Jeandel, 2004; Read, 2000). The Labrador Sea Water (LSW) is an intermediate vater class formed by deep winter convection in the Labrador Sea resulting from progressive cooling and ireshening of surface waters (station 69;Lazier, 1973). It is observed at multiple locations along the GEOVIDE section. It flows northward into the Irminger Sea (station 44) and eastward across the Mid-Atlantic Ridge (MAR). It then splits northward into the Iceland Basin (station 32), eastward into the West European Basin (stations 21 and 26), and equatorward. The Mediterranean Water (MW) enters the North Atlantic through the Gibraltar Strait and flows northwestward (García-Ibáñez et al., 2015). The Iceland-Scotland Overflow Water (ISOW) originates at the Iceland-Scotland sill and entrains the overlying warm saline Atlantic waters (SPMW) and LSW (van Aken, 2000). ISOW was found at stations located on the Eastern flank of the Reykjanes Ridge (stations 32 and 38) and near Greenland (station 64) at great depth (2000-3500 m) (Fig.2). Slightly deeper, the overflow waters coming from the Denmark Strait Overflow Water (DSOW) are then

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northern end of the Irminger Sea (station 44) and the deepest part of the Greenland continental slope (stations 69 and 77) (Fig.2). The Lower North East Atlantic Deep Water (NEADWl) is a water mass with a southern origin lying at the bottom of the West European Basin (stations 1 to 26 in Fig.2). It recirculates into the Rockall Trough and meets ISOW in the Iceland Basin (van Aken, 2000; McCartney, 1992; Schmitz and McCartney, 1993).

2.3. Sample collection

Due to the low ²²⁷Ac activities in the ocean (<5 dpm m⁻³, Geibert et al., 2002; 0.1 - 4.8 dpm m⁻³, (Levier et al., 2021), the analysis of ²²⁷Ac requires a pre-concentration step from large volumes of seawater. We used acrylic cartridges impregnated with manganese oxide (Mn-cartridges) mounted on large volume *in-situ* pumps, a method that is largely used to preconc, itrate radionuclides from seawater such as Ra and Th isotopes, as well as ²²⁷Ac (Henderson et al., 2013; Swarzenski and Baskaran, 2004). Pumps were deployed at 6 to 13 depths at each of the investigated stations. Pumping was carried out for 3 to 4 hours to filter large seawater volumes (420 to 1565 L). The resulting flow rates mostly ranged from 3 to 6 L min⁻¹. Seawater was first filtered *in situ* throug). Supor (Pall, 0.8 µm pore size) or QMA (Sartorius, 1 µm pore size, Ø 142 mm) membranes to collect suspended particles, and then through the cartridges impregnated with MnO₂ (Mn-cartridges' to collect dissolved ²²⁷Ac (as well as Ra isotopes). For the deep samples, two Mn-cartridges (ca cridge A and cartridge B) were placed in line to provide information on the yield of ²²⁷Ac fixation onto the Mn-cartridges. Following collection, each Mn-cartridge was rinsed with Ra-free milli-Q vater and slightly dried using compressed air. This protocol and the method used to determine the grie.⁴ of ²²⁷Ac fixation onto the Mn cartridges are presented in more detail in (Le Roy et al., 2019).

2.4. Quantification of the ²²⁷Ac vivities

Following Shaw and Moore (2002), we used Radium Delayed Coincidence Counters (RaDeCC, Scientific Computer Instruments, USA; Moore and Arnold, 1996) to quantify ²²⁷Ac activities on the Mn-cartridges. The ²²³Ra in equilibrium with ²²⁷Ac was determined by measuring the ²¹⁹Rn activities that are in secular equilibrium with ²²³Ra and ²²⁷Ac. This method is extensively described in (Le Roy et al., 2019) and briefly summarized below. The partially dried Mn-cartridges were placed in plastic cartridge holders in a closed helium circulation loop. Helium was allowed to circulate through the Mn-cartridges and carried the ²¹⁹Rn to the scintillation cell coated with ZnS, where alpha particles produced by the ²¹⁹Rn decaying into ²¹⁵Po were detected. A delayed coincidence system (originally developed by (Giffin et al., 1963) and then adapted by Moore and Arnold, (1996) allowed us to discriminate the signal associated to ²¹⁹Rn from those of other Rn isotopes (²²⁰Rn and ²²²Rn) which were not associated with ²²⁷Ac. Corrections for chance coincidence counts were performed following Moore and Arnold

Journal Pre-proof a Mn-cartridge containing a known amount of ²²⁷Ac (Le Roy et al., 2019). Dissolved ²²⁷Ac activities were then determined using the Mn-cartridge extraction efficiency determined from the two Mn cartridges placed in series (Le Roy et al., 2019). A mean extraction efficiency of $47 \pm 12\%$ was applied to all Mn-cartridges (1 SD, n=21; (Le Roy et al., 2019). Repeatability experiments allowed us to

estimate the uncertainty associated with the ²²⁷Ac analysis (19%, 1SD; Le Roy et al., 2019).

Excess 227 Ac activities (227 Ac_{ex}) were then calculated following:

$$^{227}Ac_{ex} = {}^{227}Ac - {}^{231}Pa$$

Equation (1)

This is assuming that a fraction of the ²²⁷Ac present in the water column is at secular equilibrium with ²³¹Pa. In some cases, the dissolved ²³¹Pa activities were collected using a separate sampling system and therefore not determined at the exact same water depth as the ²²⁷Ac activities (Deng et al., 2018). We thus used the depth interpolated ²³¹Pa activity determined from complex collected above and below the ²²⁷Ac sample.

3. Results

3.1. ²²⁷Ac distribution along the GEOVIDE section

The distribution of ²²⁷Ac activities along the GFOVIDE section is shown in figure 3. As a comparison, the ²³¹Pa activities are also reported (isolines). Most samples displayed activities ranging between 0.02 and 0.4 dpm m⁻³, with higher ²²⁷Ac activities observed in deep waters near the seafloor: deep waters at the bottom of the West European Basin and in the Irminger basin at 2500 m display ²²⁷Ac activities of 0.6 and 1.1 cpm m⁻³, respectively. In upper waters (~500 m), slightly higher activities can be found near the r argins (e.g., stations 1 and 13 located on the Iberian margin or station 64 located near Greenland).

Excluding the enrichments of served in deep waters, ²²⁷Ac activities determined along the section were in the lower range of the "Ac activities reported by Nozaki et al. (1998) in the northeast Pacific: 0.05 dpm m⁻³ in the upper water column to 2.68 dpm m⁻³ near the seafloor. However, the ²²⁷Ac activities reported here were in the same order of magnitude as those reported by (Geibert et al., 2002) in the central Arctic (²²⁷Ac activities ranging from 0.08 dpm m⁻³ at 1220 m to 0.28 dpm m⁻³ at 4220 m). In the Atlantic Ocean, (Kipp et al., 2015) reported ²²⁷Ac activities as low as 0.16 dpm m⁻³ at 1060 m and up to 0.55 dpm m⁻³ at 4700 m; higher ²²⁷Ac activities (up to 1.44 dpm m⁻³) were associated with a neutrally buoyant hydrothermal vent plume.

3.2. ²²⁷Ac vs ²³¹Pa activities and ²²⁷Ac_{ex} distribution

In general, ²²⁷Ac activities gradually increased with increasing depth at stations located in the West European Basin (stations 1, 13, 21 and 26; Fig. 4, Table 1). In contrast, the increase with depth was vertical distribution of ²³¹Pa activities that increased with increasing depth in the West European basin from 0.04 dpm m⁻³ in the surface waters to 0.28 dpm m⁻³ in the deep waters (Deng et al., 2018). In contrast, a lower and narrower range of ²³¹Pa activities was observed in the Irminger and Labrador basins (0.05 - 0.15 dpm m⁻³; (Deng et al., 2018). At intermediate depths (below ca. 500 m and above 500 m above the seafloor), ²²⁷Ac and ²³¹Pa activities were similar for most of the investigated stations. This suggests that at these depths, ²²⁷Ac originated from the decay of ²³¹Pa in the water column. However, at several intermediate depths, ²²⁷Ac activities were slightly lower than ²³¹Pa (e.g., stations 13, 21, 38).

The ²²⁷Ac_{ex} section is shown in Fig.5. The ²²⁷Ac_{ex} distribution mirrors that of ²²⁷Ac (Fig.3). Significant ²²⁷Ac_{ex} activities were found at several stations near the seafloor (13, 26, 32, 44, 64 and 69), but this was not the case for all stations, with excess activities reach c_{y} up to 1.0 dpm m⁻³ (Fig.4 and 5). Occasional isolated peaks in ²²⁷Ac_{ex} activities were found at deptl s av ay from the seafloor (station 44; 2500 m; stations 1 and 21; 500 m). Surprisingly, surface waters also sometimes exhibited significant ²²⁷Ac_{ex} activities (stations 1, 13, 32, 44 and 64).

4. Discussion

4.1. Influence of the water masses on the ²²⁷Ac di 'cil ution

²²⁷Ac is at secular equilibrium with ²³¹Pa for nost of the stations between 500 m below the surface and ca. 500 m above the seafloor, excluding the sporadic higher ²²⁷Ac values that were sometimes observed in the water column (Fig.3). Indee 4, ²²⁷Ac_{ex} is in most cases relatively low when compared to other oceanic regions (Geibert et al., 2008, 2002; Nozaki, 1984; Levier et al., 2021). Water masses that display ²²⁷Ac at secular equilibrium subgrest that they did not enter in contact with sediments over the last 100 years (five ²²⁷Ac half-life). East of the section, the vertical ²³¹Pa activities increase with increasing depth and may reflect reversible scavenging. In particular, the NEADWI, located in the eastern part of the section, he is a southern component (Fig. 2). NEADWI, with an apparent age of at least 400 years, and is an old water mass (not recently ventilated) compared to the other water masses along the section that are less than 100 years old (Deng et al., 2018). NEADWI is thus characterized by relatively high ²³¹Pa activities, since ²³¹Pa activities increase with age (Deng et al., 2018). Consequently, the ²²⁷Ac activities that are at secular equilibrium with ²³¹Pa also increase with increasing depth in the West European Basin (Fig.3).

In the western part of the section, the formation of deep-waters transports the low ²³¹Pa concentrations from surface waters into the deep ocean, where ²³¹Pa activities increase as the water mass is transported (Deng et al., 2018). In the Labrador and Irminger Basins, low ²³¹Pa (and thus ²²⁷Ac) activities throughout the water column are thus explained by the presence of recently ventilated (young)

found in the Arctic Ocean, even close to the seafloor (Geibert et al., 2002).

4.2. ²²⁷Ac_{ex} activities Near the Seafloor

The ²²⁷Ac_{ex} activities increase near the seafloor at stations 13, 26, 32, 64 and 69 (Fig.5), but this pattern is not observed at all stations as would be expected from a tracer originating from deep-sea sediments. The highest ²²⁷Ac_{ex} activity near the seafloor is found at station 13 (0.42 dpm m⁻³), which is the deepest station investigated along the GEOVIDE section (5500 m). Since the accumulation rate of ²³¹Pa in sediments increases with increasing water column depth, the flux of ²²⁷Ac diffusing out of the sediment is also a function of the water column thickness (Geibert et al., 2002). The same explanation holds for the three deep stations (13, 21, 26) located in the deep basins (Iberian Abyssal plain; West European basin) that display significant ²²⁷Ac_{ex} activities near the afloor. In addition, these stations are likely less impacted by advection and strong currents—since the basins are located away from boundary currents—than the other stations located west of the section. The ²²⁷Ac enrichments near the seafloor are, however, restricted to the deep-sea and are rat.¹v transported by vertical mixing further than 500 m above the seafloor (Fig. 4). This latter patter or currants with the profiles observed in the Pacific Ocean (Nozaki, 1984), where significant ²²⁷Ac_{ex} activities can be found at a greater distance from the seafloor.

Several vertical profiles of $^{227}Ac_{ex}$ do not nov an increase near the seafloor, where $^{227}Ac_{ex}$ activities are expected (Nozaki, 1984). Benthic nepheloid la yers were observed along the GEOVIDE section from the Iceland basin to the Labrador Basin (Stations 26 to 69), as deduced from decreases in the transmissiometry near the seafloor, indicative of an increase in the particle concentration (Fig.4). These bottom nepheloid layers may impact the distribution of radionuclides in the dissolved phase. In particular, the particulate Mn concentrations (PMn) also increase near the seafloor within the bottom nepheloid layers at all these rations (Fig.4). PMn is well known to scavenge radionuclides, including radionuclides reactive to particles such as Th or Pa. Scavenging near the seafloor has been reported by previous studies in the Labrador Basin (Bacon and Anderson, 1982; Deng et al., 2014, 2018). This is potentially also the case for radionuclides such as Ra isotopes and ²²⁷Ac, although this effect has never been investigated in past studies. It is the property of MnO_2 to adsorb radionuclides that has led to the use of fibers and cartridges impregnated with Mn to pre-concentrate radionuclides from seawater samples (Moore and Reid, 1973) and to co-p+recipitate Ra (Ghaleb et al., 2004) and Ac (Levier et al. 2021). Such impact, however, has never been clearly reported in oceanic studies. Some high manganese oxide concentrations are associated with particles resuspended from the sediment. However, we could not find, any statistical relationship between PMn concentrations and ²²⁷Ac, ²²⁶Ra and ²³¹Pa patterns near the seafloor (Fig.4). Moreover, it is unclear why ²²⁷Ac would be preferentially scavenged compared to ²³¹Pa. One hypothesis would be that recent re-suspension of surface sediments, with high

this process may really impact the distribution of radionuclides such as Ra and Ac.

4.3. ²²⁷Ac_{ex} Enrichments in the Water Column

In the Irminger and Labrador Basins, some higher $^{227}Ac_{ex}$ activities were observed in deep waters, at stations 64 and 69, or even more clearly at station 44. Peaks of $^{227}Ac_{ex}$ may be indicative of a recent (younger than ~ 100 years) contact of the water mass with ocean boundaries such as margins and deep sediments (Nozaki, 1984; Nozaki et al., 1990) or indicative of a hydrothermal plume (Kipp et al., 2015).

The highest ²²⁷Ac_{ex} activities across the section (1.12 dpm m^3) were observed at 2500 m at station 44 in association with high dissolved iron (DFe) concentrations—up to 2.5 nmol L⁻¹—between 2000 m and 2800 m (Fig.4; Tonnard et al., 2018). The DFe and ²²⁷Ac patterns observed at depth at station 44 may be attributed to the signature of a hydrothermal plu. Although there is no additional supporting evidence for hydrothermal activity (e.g. no paralle δ ³He measurements were made), hydrothermal vents have been reported along the nearby Reykjan 28 Ridge (Le Roy et al., 2019). At station 64, a slight increase of DFe concentration—up to 2.1 n nol L⁻¹—was observed between 2000 and 3000 m (Fig.4). The high ²²⁷Ac_{ex} activities observed at stations 64 and 69 in deep waters may be explained by lateral advection of waters that interacted with the margins. The Irminger and Labrador Basins are not as deep as West European Basin and the "Deep Western Boundary Current (DWBC). These features may foster interaction between the water masses and the sediments.

Slightly higher ²²⁷Ac activities were forma at stations near the Iberian margin, at 500 m (stations 1 and 21) and in the upper 500 m at station 1. This pattern is similar to that reported in the Central Arctic where an input from Siberian shelves was suspected (Geibert et al., 2002). A similar increase at 500 m in the ²²⁷Ac activities has also been recently reported in a vertical profile in the Weddell Gyre (Levier et al., 2021). Such high $^{227}Ac_{e_s}$ activities at relatively shallow depths may be explained by lateral advection of waters that interacted with shallow sediments. In coastal regions, boundary scavenging leads to preferential ²³¹Pa removal in areas of high particle flux and therefore accumulation of ²³¹Pa in margin sediments (Anderson et al., 1983; Bacon et al., 1976). However that flux maybe diluted by a higher detrital input than in deep waters. Another hypothesis is extensive irrigation of sediments by fauna that could produce high ²²⁷Ac flux. Epping et al. (2002) showed that the geochemistry of the Nazaré Canyon sediments may be significantly affected by benthic macrofauna activity. Margin sediments are thus likely a source of ²²⁷Ac to the water column. In particular, station 1 is located on the Iberian margin; the high ²²⁷Ac_{ex} activity found at relatively shallow depth could result from a lateral input of ²²⁷Ac released by the shallow sediments deposited onto the margin, a signal that is then be advected toward offshore waters, thus reaching station 13 and station 21 via different paths (Barbot et al., 2022). Near the Iberian margin (station 1), transmissiometry data indicate a shallow nepheloid layer

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Barbot et al. (2022) established that internal tides (ITs) generated in the Bay of Biscay and along the Iberian slope could facilitate sediment resuspension and could be the major mechanism for the resuspension happening below 300 m. As hypothesized for bottom waters, we cannot exclude that the shallow nepheloid layers impact the distribution of radionuclides such as Ra or ²²⁷Ac (Fig.4); in some cases, these nepheloid layers may act as a sink when particles scavenge chemical elements (e.g., radionuclides), while in other cases, they may act as a source when chemical elements are released from the particles.

4.4. Estimate of vertical eddy diffusivity coefficients (Kz)

When the vertical profiles of ²²⁷Ac result from the diffusion of ²²⁷Ac from the sediment and are not impacted by any other input or removal (e.g., bottom nepheloid la_y, r), the vertical profiles of ²²⁷Ac_{ex} activities can be used to derive vertical eddy diffusivity coefficients r_z (Nozaki, 1984). Vertical eddy diffusivity coefficients can be determined using a simplified vertical one-dimension model, assuming steady state conditions and that the impact of advection can be neglected (Nozaki, 1984):

$$\frac{\partial}{\partial z} \left(K_z \frac{\partial A}{\partial z} \right) - \lambda A = 0$$
 Equation 2

Where K_z is the vertical eddy diffusivity coefficient: Λ is the ²²⁷Ac_{ex} activity; z is the depth above the bottom; λ is the ²²⁷Ac decay constant.

The solution of Equation 2 is given by:

$$A = A_{j,z} \left(- \frac{1}{K_{z}} \right)$$
 Equation 3

Where $A=A_0$ at z=0 and A=0 at z ∞ .

We applied this equation at station 13 located in the Iberian Abyssal Plain (Fig.6).

Station 13 was selected because it is the deepest station (5455 m) of the section and it displays significant ²²⁷Ac_{ex} activities (up to 0.42 dpm m⁻³) near the seafloor. We do not observe a bottom nepheloid layer at this station (Fig.4), as it was the case at several other stations where we cannot exclude that resuspended Mn oxides may impact ²²⁷Ac activities near the seafloor. The K_z value derived from the vertical profile of ²²⁷Ac_{ex} is 0.68 \pm 0.49 cm ² s⁻¹ at that station (Fig.6). This value is within the lower range of K_z values usually reported for the deep ocean (0.1–50 cm ² s⁻¹;Huh and Ku, 1998; Kaufman et al., 1973; Koch-Larrouy et al., 2015; Nozaki, 1984; Nozaki et al., 1990). This low K_z can be explained by the topography of the Iberian Abyssal Plain since vertical mixing is linked to the underlying bathymetry and especially seafloor roughness (Polzin et al., 1997; Mauritzen et al., 2002). In

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compared to the interior of the basins (Hasumi and Suginohara, 1999; Mauritzen et al., 2002).

Conclusion

This study reports an oceanic section of ²²⁷Ac activities built from 9 full-depth vertical profiles determined between Portugal and Canada in the North Atlantic. By combining the ²²⁷Ac activities with the ²³¹Pa activities, we provide an entire section of ²²⁷Ac_{ex} activities. ²²⁷Ac activities were relatively low along that section compared to other oceanic regions (e.g., Pacific Ocean). In most cases, ²²⁷Ac was in secular equilibrium with ²³¹Pa. This suggests that at these depths, ²²⁷Ac was only produced from the decay of ²³¹Pa in the water column and that there was no external input - younger than ~100 years - of ²²⁷Ac. East of the section, the vertical ²²⁷Ac and ²³¹Pa activities increased with increasing depth, a pattern that can be explained by both the origin and age of the deep waters in the West European Basin that drove the ²³¹Pa activities (reversible exchange model). West on the section, the presence of recently ventilated (young) water masses leads to low and relatively antiform ²²⁷Ac and ²³¹Pa activities throughout the water column.

Below 2000 m, several vertical profiles were character. ed by higher ${}^{227}Ac_{ex}$ activities near the seafloor, which reflect the input of ${}^{227}Ac$ that diffu e 1 out of the sediment. Sporadic enrichments of ${}^{227}Ac_{ex}$ were observed either in surface/subsurf... of deep waters, which could be attributed to waters that interacted with the margins. At station 4 in the Irminger Basin, the high ${}^{227}Ac_{ex}$ activities at 2500 m were accompanied by high DFe concentrations; these two enrichments may indicate the presence of a hydrothermal plume. Finally, bottom nepheloid layers were observed along the section. High PMn concentrations were found in these layers and may contribute to scavenging radionuclides such as ${}^{227}Ac$ (and ${}^{226}Ra$), a process that may impact the vertical profiles of ${}^{227}Ac_{ex}$ (and Ra). At station 13 (Iberian Abyssal Plain, where no bottom nepheloid layer was observed, we estimated a vertical eddy diffusivity confineent (K_z) of 0.67 ± 0.14 cm 2 s⁻¹ from the ${}^{227}Ac$ vertical distribution; this value is consistent with K_z values previously documented for the deep ocean.

Figure 1: Location of stations investigated for ²²⁷Ac (and ²³¹Pa) along the GEOVIDE section (GEOTRACES GA01). Surface currents are displayed in warm colors while deeper currents are displayed in cold colors.

Figure 2: Distribution of salinity (CTD data) along the GEOVIDE section. The different water masses are indicated (MW: Mediterranean Water, LSW: Labrador Sea Water, NEADWI: Lower North East Atlantic Deep Water, ISOW: Iceland-Scotland Overflow Water, DSOW: Denmark Strait Overflow Water). The station numbers are found on top of the figure. The sampling depths for ²²⁷Ac are shown for each vertical profile (black dots). The isohalines are represented (white lines). This section was created with Ocean Data View (2022).

Figure 3: Distribution of dissolved ²²⁷Ac activities along the GEOVIDE section with the dissolved ²³¹Pa activities (dpm m⁻³) in contours. Station numbers are found on top of the panel. The sampling depths for ²²⁷Ac are shown for each vertical profile (black dots). This section was created with Ocean Data View (2022).

Figure 4: Vertical profiles of dissolved ²²⁷Ac with ²³¹Pa, ²⁷Ac_{ex}, ²²⁶Ra, DFe, PMn and transmissiometry determined along the GEOVIDE section. Note that the scale may be different from one station to the other. The seafloor is represented by the bottom line. Nepheloid layers are represented by the shaded area. DFe and PMn concentrations are reported in Tonnard et al. (2018) and Gourain et al. (2018), respectively and are available in the GEOTRACES Intermediate Data Product (2021). This section was created with Ocean Data View (2022).

Figure 5: Distribution of dissolved $^{227}Ac_{ex}$ activities along the GEOVIDE section. Station numbers are found on top of the panel. The sampling depths for ^{227}Ac are shown for each vertical profile (black dots).

Figure 6: Vertical profiles of ${}^{227}Ac_{ex}$ activities (dpm m⁻³) determined at station 13 in the 1_{ast} 1500 m. The exponential fit used to estimate the vertical eddy diffusivity (Kz in cm² s⁻¹) is shown (red line) with the interval of possib 2 values (dashed grey lines).

Table 1: Activities of ²²⁷Ac, and ²²⁷Ac_{ex} along the GEOVIDE section.

Station	Longitude	Latitude	Depth	²²⁷ Ac	²²⁷ Ac _{ex}	
	(°E)	(°N)	(m)	(dpm m ⁻³)	(dpm m ⁻³)	
1	-10.04	40.33	45	$0.13 \hspace{0.2cm} \pm \hspace{0.2cm} 0.02$	$0.08 \hspace{0.2cm} \pm \hspace{0.2cm} 0.03$	
1	-10.04	40.33	170	$0.06 \hspace{0.2cm} \pm \hspace{0.2cm} 0.01$	$0.00 \hspace{0.1in} \pm \hspace{0.1in} 0.01$	
1	-10.04	40.33	350	$0.06 \hspace{0.2cm} \pm \hspace{0.2cm} 0.01$	$\textbf{-0.01} \hspace{0.1in} \pm \hspace{0.1in} 0.02$	
1	-10.04	40.33	500	$0.30 \hspace{0.2cm} \pm \hspace{0.2cm} 0.05$	$0.21 \hspace{.1in} \pm \hspace{.1in} 0.06$	
1	-10.04	40.33	550	$0.13 \hspace{0.1in} \pm \hspace{0.1in} 0.02$	0.04 ± 0.03	
1	-10.04	40.33	800	$0.11 \hspace{.1in} \pm \hspace{.1in} 0.02$	0.00 ± 5	
1	-10.04	40.33	1050	$0.11 \hspace{.1in} \pm \hspace{.1in} 0.02$	$-0.0^{-1} \pm 0.03$	
1	-10.04	40.33	2000	$0.14 \hspace{0.1in} \pm \hspace{0.1in} 0.03$	-0.03 ± 0.03	
1	-10.04	40.33	2500	$0.20 \hspace{0.1in} \pm \hspace{0.1in} 0.04$	-0.01 ± 0.05	
1	-10.04	40.33	3000	$0.20 \hspace{0.2cm} \pm \hspace{0.2cm} 0.04$	-v ^4 ± 0.05	
1	-10.04	40.33	3450	0.14 ± 0.03	0.13 ± 0.04	
13	-13.89	41.38	4	0.27 ± 0.0 .	0.21 ± 0.05	
13	-13.89	41.38	40	0.10 + 0.0 -	$0.05 \hspace{0.2cm} \pm \hspace{0.2cm} 0.02$	
13	-13.89	41.38	170	0.1.5 0.02	0.07 ± 0.03	
13	-13.89	41.38	350	0	0.03 ± 0.02	
13	-13.89	41.38	450	$0.1' \pm 0.02$	0.05 ± 0.03	
13	-13.89	41.38	1500	0.07 ± 0.01	$\textbf{-0.09} \hspace{0.2cm} \pm \hspace{0.2cm} 0.03$	
13	-13.89	41.38	3000	$0.19 \hspace{0.2cm} \pm \hspace{0.2cm} 0.03$	$\textbf{-0.05} \hspace{0.2cm} \pm \hspace{0.2cm} 0.05$	
13	-13.89	41.38	4000	$0.14 \hspace{0.1in} \pm \hspace{0.1in} 0.03$	$\textbf{-0.10} \hspace{0.2cm} \pm \hspace{0.2cm} 0.04$	
13	-13.89	41.38	4850	$0.29 \hspace{0.2cm} \pm \hspace{0.2cm} 0.05$	$0.08 \hspace{0.1in} \pm \hspace{0.1in} 0.06$	
13	-13.89	41.38	5280	$0.65 \hspace{0.2cm} \pm \hspace{0.2cm} 0.12$	$0.42 \hspace{.1in} \pm \hspace{.1in} 0.12$	
21	-19.67	46.54	4	$0.00 \hspace{0.1 in} \pm \hspace{0.1 in} 0.00$	$\textbf{-0.06} \hspace{0.2cm} \pm \hspace{0.2cm} \textbf{0.01}$	
21	-19.67	46.54	300	$0.03 \hspace{0.1in} \pm \hspace{0.1in} 0.01$	$\textbf{-0.03} \hspace{0.1in} \pm \hspace{0.1in} 0.01$	
21	-19.67	46.54	450	$0.16 \hspace{0.2cm} \pm \hspace{0.2cm} 0.03$	$0.09 \hspace{0.1in} \pm \hspace{0.1in} 0.03$	
21	-19.67	46.54	800	$0.07 \hspace{0.1in} \pm \hspace{0.1in} 0.01$	$\textbf{-0.02} \hspace{0.2cm} \pm \hspace{0.2cm} 0.02$	
21	-19.67	46.54	1500	$0.04 \hspace{0.1in} \pm \hspace{0.1in} 0.01$	$\textbf{-0.07} \hspace{0.1in} \pm \hspace{0.1in} 0.02$	
21	-19.67	46.54	2700	$0.08 \hspace{0.2cm} \pm \hspace{0.2cm} 0.02$	$0.00 \hspace{0.1in} \pm \hspace{0.1in} 0.02$	
21	-19.67	46.54	3500	$0.17 \hspace{0.1in} \pm \hspace{0.1in} 0.03$	$\textbf{-0.04} \hspace{0.1in} \pm \hspace{0.1in} 0.04$	
21	-19.67	46.54	3944	$0.17 \hspace{0.1in} \pm \hspace{0.1in} 0.03$	$\textbf{-0.05} \hspace{0.2cm} \pm \hspace{0.2cm} 0.04$	
21	-19.67	46.54	4444	$0.31 \hspace{.1in} \pm \hspace{.1in} 0.06$	0.05 ± 0.07	
26	-22.60	50.28	1013	$0.09 \hspace{0.2cm} \pm \hspace{0.2cm} 0.02$	$0.02 \hspace{0.1in} \pm \hspace{0.1in} 0.02$	

26	22.60	L 60.00 - 00	~~			0.02	
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26	-22.60	50.28 35	63	$0.14 \hspace{0.1in} \pm \hspace{0.1in} 0.03$	-0.05 ±	0.04	
26	-22.60	50.28 38	68	$0.44 \hspace{0.1in} \pm \hspace{0.1in} 0.08$	0.23 ±	0.09	
26	-22.60	50.28 41	26	$0.29 \hspace{0.2cm} \pm \hspace{0.2cm} 0.05$	0.16 \pm	0.06	
32	-26.71	55.51	ŀ	$0.18 \hspace{0.2cm} \pm \hspace{0.2cm} 0.03$	0.08 \pm	0.03	
32	-26.71	55.51 43	50	$0.09 \hspace{0.2cm} \pm \hspace{0.2cm} 0.02$	-0.01 ±	0.02	
32	-26.71	55.51 80	00	$0.06 \hspace{0.2cm} \pm \hspace{0.2cm} 0.01$	-0.04 \pm	0.02	
32	-26.71	55.51 14	81	$0.10 \hspace{0.1in} \pm \hspace{0.1in} 0.02$	-0.03 ±	0.03	
32	-26.71	55.51 22	67	$0.07 \hspace{0.1in} \pm \hspace{0.1in} 0.01$	-0.02 \pm	0.02	
32	-26.71	55.51 28	54	$0.08 \hspace{0.2cm} \pm \hspace{0.2cm} 0.02$	-0.02 \pm	0.02	
32	-26.71	55.51 30	50	$0.21 \hspace{.1in} \pm \hspace{.1in} 0.04$	0.09 \pm	0.04	
32	-26.71	55.51 31	70	$0.18 \hspace{0.2cm} \pm \hspace{0.2cm} 0.03$	0.07 \pm	0.04	
38	-31.27	58.84 10	00	$0.09 \hspace{0.2cm} \pm \hspace{0.2cm} 0.02$	-0.04 ±	0.02	
38	-31.27	58.84 29	97	$0.02 \hspace{0.1in} \pm \hspace{0.1in} 0.00$	-0.11 ±	0.02	
38	-31.27	58.84 64	13	$0.12 \hspace{.1in} \pm \hspace{.1in} 0.02$	-0.05 \pm	0.03	
38	-31.27	58.84 84	0	$0.08 \hspace{0.2cm} \pm \hspace{0.2cm} 0.01$	-0.02 \pm	0.02	
38	-31.27	58.84 10	87	$0.08 \hspace{0.2cm} \pm \hspace{0.2cm} 0.01$	-0.03 \pm	0.02	
38	-31.27	58.84 12	35	$0.15 \hspace{0.1in} \pm \hspace{0.1in} 0.03$	$0.00 \pm$	0.03	
38	-31.27	58.84 13	16	$0.07 \hspace{0.1in} \pm \hspace{0.1in} 0.01$	-0.09 \pm	0.03	
44	-38.95	59.62	ŀ	$0.14 \hspace{0.1in} \pm \hspace{0.1in} 0.03$	0.05 \pm	0.02	
44	-38.95	59.62 50	00	$0.14 \hspace{0.1in} \pm \hspace{0.1in} 0.03$	0.03 \pm	0.03	
44	-38.95	59.62 11	36	$0.06 \hspace{0.1in} \pm \hspace{0.1in} 0.01$	-0.03 \pm	0.02	
44	-38.95	59.62 19	72	$0.09 \hspace{0.2cm} \pm \hspace{0.2cm} 0.02$	-0.05 \pm	0.03	
44	-38.95	59.62 25	60	$1.12 \hspace{.1in} \pm \hspace{.1in} 0.21$	$1.00 \pm$	0.21	0
44	-38.95	59.62 27	46	$0.06 \hspace{0.1in} \pm \hspace{0.1in} 0.01$	-0.01 \pm	0.02	
44	-38.95	59.62 28	80	$0.17 \hspace{.1in} \pm \hspace{.1in} 0.03$	0.08 \pm	0.03	
64	-46.08	59.07 4	0	$0.16 \hspace{0.2cm} \pm \hspace{0.2cm} 0.03$	0.06 ±	$\overline{0.0^{7}}$	
64	-46.08	59.07 40	00	$0.13 \hspace{.1in} \pm \hspace{.1in} 0.02$	0.04 ±	0.v `	
64	-46.08	59.07 90	00	$0.07 \hspace{0.1in} \pm \hspace{0.1in} 0.01$	-0.05 -	0.02	
64	-46.08	59.07 13	72	$0.13 \hspace{.1in} \pm \hspace{.1in} 0.02$	0. ^r s ~	0.03	
64	-46.08	59.07 17	16	$0.09 \hspace{0.2cm} \pm \hspace{0.2cm} 0.02$	-0.03 ±	0.02	
64	-46.08	59.07 20	60	0.21 ± 0.0^{A}	$0.1. \pm$	0.04	
64	-46.08	59.07 23	13	0.08 ± C 02	-v. 92 ±	0.02	
64	-46.08	59.07 24	43	0.29 ± \05	0.20 ±	0.05	
69	-48.09	55.84 40	50	0.08 = 0.01	0.01 ±	0.02	
69	-48.09	55.84 17	78	$0 \ (3 \ \pm \ 0.02)$	$0.01 \pm$	0.03	
69	-48.09	55.84 24	63	<u>`31 £ 0.06</u>	0.25 ±	0.06	
69	-48.09	55.84 29	53	0.1 ± 0.03	$0.08 \pm$	0.03	
69	-48.09	55.84 34	40	$0.23 \hspace{.1in} \pm \hspace{.1in} 0.04$	$0.17 \pm$	0.04	
69	-48.09	55.84 36	16	0.12 ± 0.02	$0.08 \pm$	0.02	
77	-51.09	52.99	ŀ	0.27 ± 0.05	0.16 ±	0.05	
77	-51.09	52.99	ŀ	$0.08 \hspace{0.2cm} \pm \hspace{0.2cm} 0.02$	-0.02 ±	0.00	
77	-51.09	52.99 64	3	$0.03 \hspace{0.1in} \pm \hspace{0.1in} 0.01$	-0.06 \pm	0.01	

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Figure 1



Figure 2



Figure 3







Figure 5



Figure 6