



GEOTRACES Intercalibration of ^{230}Th , ^{232}Th , ^{231}Pa and prospects for ^{10}Be

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5556 Running Head = Intercalibration of ^{230}Th , ^{232}Th , ^{231}Pa and ^{10}Be
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For Review Only

Abstract

Nineteen labs representing nine nations participated in the GEOTRACES intercalibration initiative that determined concentrations of ^{232}Th , ^{230}Th , ^{231}Pa or ^{10}Be in seawater, suspended particles or sediments. Results generally demonstrated good agreement among labs that analyzed marine sediments. Two sets of seawater samples, aliquots of particulate material filtered in situ, and/or aliquots of biogenic sediments were distributed to participating labs. Internal consistency among participating labs improved substantially between the first and second set of seawater samples. Contamination was a serious problem for ^{232}Th . Standard Niskin™ bottles introduced no detectable contamination, whereas sample containers, reagents and labware were implicated as sources of contamination. No detectable differences in concentrations of dissolved ^{232}Th , ^{230}Th or ^{231}Pa were observed among samples of seawater filtered through Nuclepore™, Supor™ or QMA™ (quartz) filters with pore diameters ranging between 0.4 and 1.0 μm . Isotope yield monitors equilibrate with dissolved Th in seawater on a time scale of much less than one day. Samples of filtered seawater acidified to a pH between 1.7 and 1.8 experienced no detectable loss of dissolved Th or Pa during storage for up to three years. The Bermuda Atlantic Time Series station will serve as a GEOTRACES baseline station for future intercalibration of ^{232}Th and ^{230}Th concentrations in seawater. Efforts to improve blanks and standard calibration are ongoing, as is the development of methods to determine concentrations of particulate nuclides, tests of different filtration methods, and an increasing awareness of the need to define protocols for reporting uncertainties.

1.0 Introduction

Distributions of naturally occurring long-lived radionuclides provide quantitative information about the rates at which processes occur in the modern ocean (Francois 2007; Rutgers Van Der Loeff and Geibert 2008) while also providing insight into changes that have occurred in the past (Francois 2007; Henderson and Anderson 2003). As a consequence of their widespread use as tracers in ocean research, certain long-lived radionuclides (^{230}Th , ^{231}Pa) have been designated in the GEOTRACES Science Plan as “key” variables to be measured on all GEOTRACES ocean sections, while others (^{232}Th , ^{10}Be) are recommended to be measured where possible. Anticipating that concentrations of these nuclides will be determined by many labs worldwide throughout the duration of the GEOTRACES program, a substantial international intercalibration effort was organized to ensure accuracy and internal consistency throughout the global data set.

Analytical advances over the time span of a single generation have revolutionized the study of these radionuclides. The first reported measurements of dissolved ^{230}Th and ^{231}Pa in open-ocean seawater involved the shipboard processing of sample sizes between 130 and 190 liters. A volume this large was required to collect sufficient activity to determine nuclide concentrations by alpha decay counting techniques (Moore and Sackett 1964). Decay-counting methods remained the norm for the next two decades. Samples as large as 250 liters were collected to study these nuclides (Nozaki et al. 1981; Nozaki and Nakanishi 1985). As an alternative to shipboard processing of large volumes of seawater, MnO_2 -impregnated absorbers were also employed to extract Th and Pa in situ (Anderson et al. 1983a; Anderson et al. 1983b; Nozaki and Horibe 1983).

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3 The transition from decay counting to the counting of atoms by mass
4 spectrometry revolutionized the field, lowering sample size requirements by two orders
5 of magnitude or more depending on the nuclide (Chen et al. 1986). Many of the labs
6 participating in this intercalibration can now measure concentrations of ^{230}Th and ^{232}Th
7 reliably using sample volumes of one to two liters of seawater. Some labs employ
8 Thermal Ionization Mass Spectrometry (TIMS, e.g., Chen et al. 1986; Huh and Beasley
9 1987; Shen et al. 2003) while others use high-resolution Inductively-Coupled Plasma
10 Mass Spectrometry (ICP-MS, e.g., Choi et al. 2001; Fleisher and Anderson 2003; Shen et
11 al. 2002). This substantial reduction in sample size represents a critical advancement that
12 permits these nuclides to be measured routinely on GEOTRACES sections.
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27 Although these nuclides have been studied for decades, procedures for collecting
28 and processing large numbers of samples were relatively new and untested prior to
29 GEOTRACES. A systematic intercalibration among a large number of labs had not been
30 conducted previously, leaving some concern about the accuracy and internal consistency
31 of prior results. Because all labs use isotope dilution, variable chemical recovery was not
32 anticipated to be a problem. Furthermore, there is little industrial use of Th, and no
33 industrial use of Pa or of ^{10}Be , so contamination was also not expected to be a problem.
34 Factors that were considered to be of potential concern included:
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46 1) *Calibration of chemical yield monitors (spike isotopes):* The absence of readily
47 available and universally accepted standards for the determination of these nuclides led to
48 different strategies among participating labs for the calibration of chemical yield
49 monitors (spikes). These strategies had not been compared systematically prior to the
50 GEOTRACES initiative.
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3 2) *Sensitivity to filtration method*: Thorium isotopes, ^{231}Pa , and ^{10}Be are highly insoluble
4 in seawater. Thus, there was concern for loss of dissolved nuclides through sorption to
5 filter material or to the filtration apparatus when determining dissolved nuclide
6 concentrations.
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12 3) *Sample storage*: Loss of these dissolved nuclides by precipitation or by sorption to
13 container walls is another potential negative bias created by the insoluble nature of these
14 nuclides. Previous studies have shown that dissolved Th and Pa are lost during storage of
15 unacidified samples (Choi et al. 2001), and anecdotal accounts had suggested that
16 dissolved Th could be lost over time even from acidified samples. Spiking samples at sea
17 immediately following their collection would circumvent this problem. However, some
18 nations impose strict restrictions on the use of artificial radionuclides aboard research
19 vessels. Furthermore, the isotope of Pa used as a yield monitor (^{233}Pa) has a half-life of
20 only 27 days. It is simply impractical to prepare a yield monitor prior to sailing, and then
21 collect and process a large number of samples before ^{233}Pa concentrations fall below
22 detection limits. Storage of unspiked samples would be preferable provided this did not
23 bias the results.
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41 Perhaps the greatest surprise discovered during this intercalibration was the
42 severity of the contamination problems that plagued the determination of ^{232}Th
43 concentrations in seawater. The foremost outcome of this study is to identify the critical
44 nature of precautions to avoid contamination of seawater samples by extraneous sources
45 of ^{232}Th , and the importance of collecting blanks regularly during research cruises. By
46 contrast, no serious problems were identified pertaining to sample storage or to different
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3 types of filter material, although additional tests of filtration methods constitute one of
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5 the recommendations for future work.
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10 **2.0 Materials and Procedures**

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12 2.1 Solicitation of participants: Upon completion of the Science Plan in 2006,
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14 the GEOTRACES community turned its attention to intercalibration of methods as the
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16 next step in implementing the program. General information about the intercalibration
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18 process, together with a solicitation for participation by interested investigators, was
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20 distributed via the international GEOTRACES email list in July 2006, in January 2007
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22 and again in August 2007. Information about intercalibration was also posted on the
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24 GEOTRACES web site along with a list of all investigators interested in participating and
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26 the variable(s) that each investigator proposed to measure.
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32 2.2 Discussion of issues: In July 2007 the authors of this paper contacted
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34 those investigators who had expressed an interest in participating in the intercalibration of
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36 ^{230}Th , ^{232}Th , ^{231}Pa and/or ^{10}Be . Investigators were asked to identify the nuclide(s) to be
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38 measured as well as the type(s) of samples to be analyzed (e.g., seawater, sediments,
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40 particles). Investigators were also polled about specific tests of sampling and analytical
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42 methods that could be made by the authors. Principal concerns that needed to be
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44 addressed before the international community could embark on a global study of these
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46 nuclides include:
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50 1) *Sample collection*: With some exceptions (Moran et al. 1997; Moran et al.
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52 2002; Moran et al. 2001), seawater samples for these nuclides have generally been
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54 collected using Niskin™ bottles or similar devices that are not specifically designed to
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3 avoid trace metal contamination. Tests had never been performed to compare Niskin
4 bottles against trace metal-clean sampling systems. Consequently, intercalibration
5 protocols were designed to allow a comparison between results obtained using Niskin
6 bottles and results obtained using GO-Flo™ bottles (designed for trace metal sampling)
7 as a byproduct of other tests described below.
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15 2) *Sample filtration*: Filtration of seawater is necessary to determine
16 operationally defined concentrations of dissolved nuclides, but no convenient or
17 universally accepted procedure had been developed for samples as large as those required
18 here (5 to 20 liters). Prior to GEOTRACES some labs had filtered seawater by gravity
19 flow directly from Niskin bottles (e.g., Choi et al. 2001) while others transferred seawater
20 to secondary containers for pressure filtration. Where GO-Flo bottles have been used
21 previously to sample for these nuclides, seawater was filtered by pressurizing sample
22 bottles with nitrogen (Moran et al. 1997; Moran et al. 2002; Moran et al. 2001). Still
23 other investigators had limited their measurements to total (unfiltered) nuclide
24 concentrations (e.g., Chase et al. 2003). Furthermore, there was no consensus about the
25 best type of filter membrane to use. Following recommendations from the participants,
26 the authors designed a system (described below in Section 3.1) that allowed seawater to
27 be pressure filtered directly from Niskin bottles. Pressure filtration was then used to test
28 a variety of filter holders and filter membranes.
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48 3) *Processing at sea vs. sample storage*: Whether or not to spike and concentrate
49 samples at sea was the topic that generated the most discussion. The principal argument
50 in favor of processing samples at sea, at least through concentration by co-precipitation,
51 is the desire to reduce the volume and weight of samples that must be shipped back to the
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3 lab (3 to 5 thousand liters of water following a typical section cruise). Several concerns
4 and arguments against processing at sea were discussed as well. For example, different
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8 labs use different carriers ($\text{Fe}(\text{OH})_3$, MnO_2 , $\text{Mg}(\text{OH})_2$) to extract the nuclides from
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10 seawater. Furthermore, processing at sea requires the addition of ^{229}Th and ^{233}Pa yield
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12 monitors. As noted above, some nations impose severe restrictions on the use of artificial
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14 radionuclides aboard research vessels, and some participants would not be allowed to use
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17 ^{233}Pa on their cruises. All participants were concerned about the time frame for
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19 completion of sample analysis. Some cruises have a duration as much as two half lives
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21 of ^{233}Pa . More important, it is simply not possible to process hundreds of samples before
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23 the ^{233}Pa has decayed below routine limits of detection. The option of gamma counting
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25 samples at sea after concentration by co-precipitation was considered as a strategy to
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27 determine the chemical yield of Pa before decay of ^{233}Pa . Following decay of the initial
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29 ^{233}Pa , the oxyhydroxide carrier could then be spiked with a second aliquot of ^{233}Pa prior
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31 to completion of the analysis. Although processing of samples at sea was tested, in the
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33 end this proved to be a non-viable option because GEOTRACES cruises are so heavily
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35 subscribed that a sufficient number of berths are not available to accommodate the
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37 personnel who would be needed to collect and process the samples at sea. Consequently,
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39 samples for these nuclides collected during GEOTRACES cruises must be stored for
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41 processing in shore-based laboratories.
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48 A rigorous comparison of replicate samples stored for variable periods of time
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50 had never been carried out. Choi et al. (2001) reported that dissolved Th and Pa were lost
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52 from unacidified samples during storage, and that external reproducibility was degraded
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3 following storage of acidified but unfiltered samples. Here, testing the quality of results
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5 obtained following storage of filtered and acidified samples was set as a high priority.
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8 4) *Particulate nuclide concentrations*: A complete study of the marine
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10 biogeochemical cycles of ^{230}Th , ^{232}Th and ^{231}Pa requires the collection and analysis of
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12 samples for their particulate as well as their dissolved concentrations. Tests for
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14 particulate ^{10}Be were not conducted because it was anticipated that concentrations of
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16 particulate ^{10}Be would be below detection limits given the partition coefficients reported
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18 by Chase et al. (2002; 2003) and the expected dissolved ^{10}Be concentrations. Based on
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20 the detection limits reported by many labs, it was estimated that a minimum of 100 to 150
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22 liters of seawater must be filtered to measure concentrations of particulate ^{231}Pa . Samples
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24 of this size require in situ filtration, which was the purview of another intercalibration
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26 group led by investigators at the Woods Hole Oceanographic Institution (WHOI).
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28 Consequently, an agreement was negotiated with those investigators to provide filter
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30 material collected by in situ filtration for this intercalibration (see Section 2.3.3 below).
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32 In addition, a large amount of siliceous ooze from the Southern Ocean was homogenized
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34 and distributed to participants to serve as a proxy for biogenic-rich particulate matter.
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36 Distribution of sediment samples also allowed for the participation by several labs that
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38 routinely analyze sediment samples, but which do not analyze seawater for these
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40 nuclides.
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48 2.3 Intercalibration samples: Seawater and particulate material were collected
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50 during two intercalibration cruises aboard the *RV Knorr*: KN193-6, leg 2 (Bermuda to
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52 Norfolk, Virginia; July 2008) and KN195-8 (Honolulu, Hawaii to San Diego, California;
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54 May 2009).
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2.3.1 Collection of seawater: Two rosette systems were used to collect seawater on each cruise: a standard rosette with twenty-four 12-liter Niskin bottles and the U.S. GEOTRACES sampling system (Cutter and Bruland this volume), which operates twenty-four 12-liter GO-Flo bottles. In addition, near-surface seawater was collected using the “GeoFish” towed sampling system designed at the University of California Santa Cruz.

The standard Niskin rosette was owned by the ship operators (WHOI) and its operation during these cruises followed standard ship protocols, including the use of steel hydrowire and closing of bottles on the up cast of the package. Niskin bottles were equipped with nylon-coated closure springs and Viton™ O-rings. When the Niskin rosette was used to test different filtration methods all 24 bottles were closed at a single depth to provide replicate samples that were assumed to be from a homogeneous water mass. Typically, water from each Niskin bottle was filtered through a single filter, except in a few cases where it was necessary to change filters that had become clogged. The ship’s rosette was also used to produce full water column profiles of the concentrations of ^{230}Th , ^{232}Th and ^{231}Pa . There, three Niskin bottles were fired at each of eight depths to provide replicate samples (see Section 5.11 discussing “baseline” profiles).

Water to be distributed to the labs participating in this intercalibration was collected near the surface (5 to 15 m) using the GeoFish and from mid depth using the U.S. GEOTRACES sampling system. During the first cruise (KN193-6, Leg 2) mid-depth water was collected from 2000 m at the Bermuda Atlantic Time Series Station (BATS; roughly 75 km southeast of Bermuda at 31°50'N, 64°10'W). Surface water was also collected near the BATS site. Similar protocols were followed during the second

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3 cruise (KN195-8), where mid depth water was collected from 3000 m at a station located
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5 at 30°N and 140°W (also known as the SAFe station). At each location five casts of the
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7 U.S. GEOTRACES system were used to collect ~1000 liters of mid-depth seawater for
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9 this intercalibration.
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13 Water from GO-Flo bottles and from the fish was filtered through 10-inch
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15 Osmonics (Memtrex™) cartridges containing sequential 0.45 µm then 0.2 µm pore size
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17 polycarbonate, track etched (PCTE) pleated filter membrane, with a total filtration
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19 surface area of 1.6 square meters. All filtrations were performed inside a portable clean
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21 van. Filtered water was pumped into two 500 L acid-cleaned fluorinated low density
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23 polyethylene tanks through PFA Teflon™ tubing using an all PFA Teflon diaphragm
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25 pump. The two tanks were connected with all Teflon tubing, valves and fittings. Water
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27 was pumped continuously between the two 500-liter tanks to produce and maintain a
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29 homogenized sample. Once the tanks were filled, the water was acidified to pH ~1.7
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31 using either Fisher Optima™ HCl, Seastar™HCl, laboratory redistilled HCl or a
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33 combination of these sources. Individual aliquots of 1 to 20 liters were drawn from the
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35 filtered, acidified and homogenized seawater for distribution to participating labs.
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41 Each participating lab provided its own precleaned sample containers, which were
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43 shipped to the cruise and back to the home institution inside double plastic bags to reduce
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45 the chance of contamination. Each lab was normally provided with three 10-liter samples
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47 from each deep intercalibration depth and a single 10-liter sample from the near-surface
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49 depth. Triplicate samples from the deeper intercalibration depth were intended to provide
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51 a measure of internal reproducibility. Some labs subdivided their 10-liter samples to
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53 provide a larger number of replicates.
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4 2.3.2 Biogenic sediments: Silicious ooze sediment was collected by the LDEO
5 authors in 1996 during a cruise of the *RVIB Nathaniel B. Palmer* (NBP9604 Station 3,
6 Lat 61.95°S Lon 170.05°W, water depth 3404 meters). Approximately 120 grams dry
7 weight of near-surface sediment recovered from the trimmings of extruded multi-cores
8 was homogenized, freeze-dried and distributed to participating labs. The sediment had an
9 opal content of ~71% and a CaCO₃ content of ~7%.

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17 2.3.3 Collection of particles: In order to provide a sufficient mass of particulate
18 material for the measurement of ²³⁰Th and ²³¹Pa, and at the same time obtain comparable
19 samples from the same depth and time, investigators at WHOI designed and built a trace
20 metal clean pump-rosette (Maiti et al. this volume). The rosette frame was made of
21 epoxy-coated aluminum and deployed from a Kevlar™ wire. Up to eight battery-
22 powered McLane pumps were operated simultaneously at a single depth, effectively
23 providing samples of particulate material from the same water mass.
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34 A PVC device was designed to subsample filters. For subsampling, a filter was
35 placed on a base plate constructed of the same frit material as used in the McLane pump
36 filter-holders. The base of the device was attached to a vacuum pump to hold the filters in
37 place during subsampling, and the filters were sliced wet. A slotted PVC ring placed
38 around the outside of the filter allowed each filter to be subdivided into halves, quarters
39 or eighths with a ceramic knife. The base frit was replaced or acid washed between each
40 set of filters. All processing was carried out under clean conditions in a High Efficiency
41 Particulate Air (HEPA) flow bench. Under shore-based lab conditions blank filters were
42 sliced and weighed and the variability in weight between quarters was found to be less
43 than 2%, providing a measure of subsampling reproducibility. Subsamples from a single
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3 sample filter counted for ^{234}Th indicated intrafilter variability of ~2.5%, while differences
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5 between subsamples from filters housed in different pumps on the same rosette cast
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7 exhibited variability of between 15 and 20% (Maiti et al. this volume). Ultimately, this
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9 inter-pump variability places a practical limit on our ability to intercalibrate the
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11 determination of concentrations of particulate ^{230}Th , ^{232}Th and ^{231}Pa .
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15 Particles were collected for the Th-Pa intercalibration using 142 mm diameter
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17 Supor™ polyethersulfone filters (0.45 μm pore diameter). During a cast each pump
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19 generally filtered between 400 and 600 liters. Quarter-filters, sectioned as described
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21 above, were provided by investigators at WHOI and distributed by the authors to
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23 participating labs.
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27 A blank filter (dip blank) for each McLane pump was produced by inserting an
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29 individual 142-mm Supor filter into a 53 μm mesh bag (closed on all sides) and then
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31 placing the bag in a plastic box with holes drilled through its sides. Each pump had its
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33 own dip-blank box attached to its frame. Filters used for dip blanks were taken from the
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35 same batch as those used for the intercalibration samples. Dip-blank filters were
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37 sectioned following the same protocol that was used for the sample filters. Dip-blank
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39 filters were analyzed by participating labs to estimate the total procedural blank for
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41 particulate analysis. Although the dip blank filters do not provide a true procedural
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43 blank, it is thought that they provide a more accurate representation of the particulate
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45 blank than would be offered by a reagent blank alone.
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51 **2.4 Participating labs:** A summary of the laboratories that submitted results
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53 for this intercalibration is provided in Table 1. Also reported are the variables reported
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3 by each lab as well as a summary of the principal features of the analytical procedures
4 employed by each lab.
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10 **3.0 Procedures used at LDEO**

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12 The LDEO authors were responsible for a number of tests recommended by the
13 participants (Section 2.2). Therefore, the methods employed at LDEO are described here
14 in some detail.
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20 3.1 Seawater Samples and Blanks: Seawater samples were collected in low-
21 density polyethylene cubitainers (either 1 gallon or 10 liters volume; Hedwin Corp.) that
22 had been cleaned by soaking in 10% HCl (trace metal grade, Fisher Scientific) followed
23 by Mill-Q water rinse. Blanks were prepared at sea by adding at least two liters of Milli-
24 Q water to a cubitainer and then treating it as a sample.
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31
32 Niskin bottles were sampled on the open deck of the ship while still in place on
33 the rosette. Sampling under these conditions presented a potential source of
34 contamination, but no alternative was available. Sampling was completed as rapidly as
35 possible, and the sampling/filtration system was designed to minimize the potential for
36 contamination from the shipboard environment. For unfiltered samples, water was
37 drained via acid washed Teflon-lined Tygon™ tubing directly into the cubitainers.
38
39 Filtered samples were collected by in-line filtration from pressurized Niskin bottles.
40
41 Bottles were pressurized at 8-10 psi (54-70 kPa) with a system that distributed filtered air
42 from a compressor via a manifold to each bottle through a fitting placed in its air vent.
43
44 Each distribution line from the manifold was fitted with a valve so that each bottle could
45 be pressurized or vented independently, while still allowing for the possibility to
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3 simultaneously filter all the bottles. Pressurized bottles were held closed with Irwin
4 Quick-Grip™ bar clamps spanning the endcaps. No attempt was made to evaluate the
5
6 blank contributed by the pressurization scheme. However, pressurization of Niskin
7
8 bottles has been discontinued on subsequent U.S. GEOTRACES cruises in favor of
9
10 gravity filtration through Acropak capsules.
11
12
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14
15 The various filter types were precleaned in batches by soaking for several days in
16
17 10% HCl (trace metal grade, Fisher Scientific), followed by repeated soaking in Milli-Q
18
19 water until a neutral pH was measured. All of the shipboard filter handling, including
20
21 loading into Savillex™ filter holders (47mm or 90 mm) was carried out inside a HEPA
22
23 filtered laminar flow bench located within the main lab of the ship. Filter holders were
24
25 rinsed with Milli-Q water between each use. Teflon-lined Tygon tubing connected the
26
27 filter holders to the petcocks of the Niskin bottles. Each filter holder was held in place
28
29 directly above the mouth of the cubitainer, so that filtered water flowed directly into the
30
31 cubitainer. The first few milliliters of filtrate were used to rinse the cubitainer and then
32
33 discarded.
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38
39 SAFE tanks were located inside an enclosure constructed with plastic sheeting
40
41 held in place by a wooden frame. Water from the SAFE tanks was transferred to each
42
43 cubitainer through a length of plastic tubing connected to the Teflon diaphragm that
44
45 circulated water between the tanks. Sample containers (both those from LDEO and those
46
47 provided by participating labs) were rinsed with water from the tank prior to filling. Labs
48
49 participating in the intercalibration generally provided their own sample containers, each
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51 cleaned by the protocol normally used by the lab. Labs that requested samples after the
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3 completion of the first cruise were provided with seawater collected in cubitainers
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5 cleaned at LDEO and archived for future use.
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8 Samples from the Niskin bottles were acidified to between pH 1.7 and 1.8 using
9
10 Optima HCl (~40 ml of 6 Normal HCl for a 10 liter sample). All samples, whether
11
12 collected from Niskin bottles or from the SAFe tank, were double bagged for shipment to
13
14 their final destinations.
15
16

17 18 19 20 3.2 Coprecipitation and digestion of seawater samples 21

22 Unless otherwise noted, all of the acids and the ammonium hydroxide used at
23
24 LDEO in the procedures outlined below were Optima Grade from Fisher Scientific.
25
26

- 27 1) At LDEO, each seawater sample was weighed (corrected for the tare weight of the
28 cubitainer and for the weight of 6N HCl added at sea), spiked with known
29
30 (weighed) amounts of isotope yield monitors (about 20 pg ^{229}Th and 0.5 pg ^{233}Pa
31
32 for each sample), and 100 μl Fe carrier (FeCl_3 , ~150 mg Fe/ml, purified by
33
34 repeated extraction into isopropyl ether) was added.
35
36
- 37 2) The pH was raised to between 8.0 and 8.5 with NH_4OH to precipitate iron
38
39 hydroxide, which was concentrated by settling and centrifugation before digestion
40
41 in a sequence of HNO_3 , HClO_4 and HF. The HClO_4 and HF eliminate organic
42
43 matter and amorphous silicon that are carried by the iron hydroxide. All steps
44
45 following centrifugation took place in a HEPA filtered laminar bench.
46
47
- 48 3) The digested sample was taken up in dilute HCl and the iron hydroxide
49
50 precipitation step was repeated.
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3 4) The precipitate was dissolved in HCl, heated to dryness at 120°C, and dissolved
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5 in 1 ml concentrated HCl (heating at 50°C as needed to dissolve precipitate).
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10 3.3 Column chemistry

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12 Purification of Th and Pa was carried out by ion exchange using a series of three
13 columns. All of the column chemistry was conducted using columns with a bed volume
14 of 1ml BioRad AG1-X8 100-200 mesh anion resin (Cl⁻ form). Columns were
15 polypropylene with 45 μm porous polyethylene frits and a reservoir volume of ~6 ml.
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20 Step #1 (Separation of Th and Pa)

- 21
22 1) After preconditioning the column with concentrated HCl, the sample was loaded
23 onto the column in concentrated HCl (~1 ml, from above), which was collected
24 along with 6 ml (4 X 1.5 ml aliquots) concentrated HCl rinse as the Th fraction.
25
26
27 2) Pa was eluted with 3 X 2ml concentrated HCl/0.13N HF.
28
29 3) Iron was washed off the resin with 10-15 ml dilute (~1%) HCl and the column
30 was held for Step 2 below.
31
32
33 4) The Th and Pa fractions were dried down overnight at 105°C, after the addition of
34 0.5 ml of concentrated HNO₃ and 2 drops of concentrated HClO₄ to decompose
35 any organics that eluted from the resin. Samples were heated to a small drop of
36 HClO₄.
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48 Step #2 (Final purification of Th)

- 49
50 1) The Th fraction was converted to HNO₃, heated to fuming HClO₄, and taken up in
51 1 ml 8N HNO₃, before loading onto the original ion exchange column that was
52 reconditioned with 2 X 2ml 8N HNO₃.
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- 2) The beaker and column were rinsed with 4 X 1 ml 8N HNO₃.
- 3) Th was eluted with 200µl concentrated HCl, followed by 4 X 1.5ml concentrated HCl.
- 4) Two drops of HClO₄ and 0.5 ml concentrated HNO₃ were added and the solution was dried overnight at ~100°C.
- 5) The following day, 250 µl concentrated HNO₃ (and a drop of HClO₄ if the sample went dry overnight) were added and the solution was heated to a drop of fuming HClO₄ at 180°C. In preparation for analysis by ICP-MS, the sample was taken up in 1 ml of 1% HNO₃/0.1% HF, capped tightly and cooled. The small amount of HF was found to help reduce memory effects in the sample introduction system, including the Aridus desolvating nebulizer unit.

Step #3 (Final purification of Pa)

- 1) A drop of HClO₄ and 200 µl concentrated HNO₃ were added to the Pa fraction, which was heated at 180°C to fuming HClO₄ in order to drive off all remaining HF. The sample was taken up in a few drops of concentrated HCl and heated again to fuming HClO₄ at 180°C. Two drops of concentrated HCl were added and the beaker was cooled before adding 1 ml concentrated HCl.
- 2) The column used for Th was reconditioned with 3 ml trace metal grade concentrated HCl/0.13N HF, 2 full reservoirs Milli-Q water, and finally 2 X 2ml concentrated Optima Grade HCl.
- 3) The sample was loaded on the column in 1 ml concentrated HCl, followed by six 1-ml concentrated HCl rinses, after which Pa was eluted with 3 X 2ml concentrated HCl/0.13N HF.

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- 4) Two drops of HClO_4 and 0.5 ml concentrated HNO_3 were added and the solution was dried overnight at $\sim 100^\circ\text{C}$.
 - 5) Another drop of HClO_4 and 200 μl concentrated HNO_3 were added as needed, the sample was dried at 100°C to a drop of HClO_4 and take up in 0.75 ml of 1% $\text{HNO}_3/0.1\%$ HF.

3.4 Filter digestion

Supor polyethersulfone filters were used for the GEOTRACES intercalibration of particulate Th and Pa concentrations. Total digestion of these filters presented problems for several groups, so the method used at LDEO to successfully digest this filter material is described below.

Lab equipment

- 1) This procedure uses perchloric acid, so a hood certified for HClO_4 use is absolutely required.
- 2) A hot plate capable of reaching $200\text{-}220^\circ\text{C}$.
- 3) 60 ml Savillex Teflon jars or equivalent.
- 4) Concave “White” Teflon watch glasses to fit the 60 ml Savillex jars.

Laboratory procedure

- 1) Weighed spikes, 100 μl of Fe carrier (Section 3.2) and 5 ml concentrated HNO_3 were added to the filter in a 60 ml Savillex jar.
- 2) The sample was heated overnight at 150°C in the Savillex jar covered with a white Teflon watch glass. HNO_3 was added as needed to maintain volume.

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- 3) The sample was allowed to cool, after which 3 ml concentrated HClO_4 were added and the temperature was raised to 200°C . The sample was heated uncovered until white HClO_4 fumes appear.
- 4) The hotplate temperature was reduced to 180°C and the sample was covered again with the white Teflon watch glass. After about 30 minutes, the HClO_4 began to oxidize the Supor filter (as evidenced by foaming on the filter surface). Soon thereafter the oxidation reaction accelerated, sometimes becoming violent enough that some of the filter material splattered onto the watch glass and walls of the jar.
- 5) Upon completion of the oxidation, the walls and watch glass were rinsed with water into the jar, and the contents were heated to fuming HClO_4 at 200°C . If necessary, 1 or 2 ml additional HClO_4 were added. About an hour was needed, depending on the amount of water and HClO_4 added. Dissolution was complete when the solution was visibly free of any remaining filter material. Heating was continued until dense HClO_4 fumes appeared.
- 6) The sample was allowed to cool briefly, after which 10 drops ($200\text{-}300\ \mu\text{l}$) concentrated HF were added to attack the silicate minerals and opal in the particles. The sample was heated until HClO_4 fumes appeared and then an additional 5 drops of HF were added.
- 7) The sample was heated to the fuming HClO_4 stage again, after which the jar walls were rinsed with concentrated HNO_3 .
- 8) Heating continued until less than 0.5 ml solution remained.
- 9) 0.5 ml concentrated HCl and then 5 ml water were added to the jar.

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3 10) The sample was heated for a few minutes on the hot plate, then removed from the
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5 hot plate, covered, and cooled. The solution had a yellow/orange color from the
6
7 iron, but it was free of visible solids.
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9

10 At this point, the solution was transferred to a 50 ml centrifuge tube with a few
11
12 5ml water rinses of the sample jar. NH_4OH was used to raise the pH to ~8 and iron
13
14 hydroxide precipitated. After centrifuging, the $\text{Fe}(\text{OH})_3$ was washed in Milli-Q water,
15
16 centrifuged again and dissolved in HCl. From that point the column chemistry was the
17
18 same as that described above for seawater samples.
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22 NOTE that step 4 involves a violent oxidation reaction. Test this procedure using a
23
24 small piece of filter material before applying it to a complete sample.
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27 28 29 3.5 ICP-MS run information

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31 Analyses were made on a VG Elemental AXIOM Single Collector Magnetic
32
33 Sector ICP-MS with a Resolving Power of ~400 to ensure the highest sensitivity. All
34
35 measurements were done using a peak jumping routine in ion counting mode. A solution
36
37 of SRM129, a natural U standard, was run to determine the mass bias correction
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39 (assuming that the mass fractionation for Th and Pa are the same as for U). The
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41 concentration of U in the SRM129 solution was 100 pg/ml.
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46 An aliquot of the final Th solution, typically 100 μl , was diluted to 800 μl with the
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48 LDEO “run solution” (1% HNO_3 /0.1% HF) for the ^{232}Th measurement. The remaining
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50 900 μl of the final Th solution was used without further dilution for the ^{230}Th
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52 measurement. This scheme was more time consuming than measuring all of the Th
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54 isotopes at once, but it allowed ^{232}Th to be measured on the multiplier so there was no
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3 need to make a Faraday/multiplier gain correction for the ^{232}Th analyses. SRM129 was
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5 run three times each day with both the Th and Pa analyses. Each sample measurement
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7 was bracketed by measurement of an aliquot of the run solution, used to correct for the
8
9 instrument background count rates on the masses measured.
10
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12 To correct for potential tailing of ^{232}Th into the minor Th and Pa isotopes, beam
13
14 intensities were measured at the half masses above and below each mass for ^{230}Th , ^{231}Pa
15
16 and ^{233}Pa . Tailing under each minor isotope was estimated as the log mean intensity of
17
18 the half masses on either side of each minor isotope.
19
20

21 For all results from LDEO, error bars represent the propagation of one sigma
22
23 errors based on the standard deviation of five sequences of isotope ratios collected by
24
25 ICP-MS, our estimated error in the ^{229}Th or ^{233}Pa spike concentration, and the blank
26
27 correction of the individual isotopes for each sample batch. Blanks processed
28
29 concurrently with each batch of samples were used rather than a grand mean blank
30
31 covering the entire intercalibration effort because blank levels were reduced significantly
32
33 during the course of this work, especially for ^{232}Th (from ~100 pg initially to ~10 pg for
34
35 the last samples processed). Blanks were reduced primarily by carrying out sample
36
37 digestions and column chemistry inside a laminar flow bench. Further reduction of the
38
39 blank was achieved by adopting a modified version of the procedure for cleaning anion
40
41 exchange resin described by Andersson et al. (this volume). Acid cleaning the
42
43 microcentrifuge tubes used as sample vials for the ICP-MS also helped reduce the blank.
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50 Additional information about sampling and purification methods, as well as
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52 information about offline processing of ICP-MS data, is available on request from the
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54 LDEO authors.
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4.0 Developing a Baseline Station at BATS

Two additional sets of results from the Bermuda Atlantic Time Series site were combined with those from the first GEOTRACES intercalibration cruise to establish a baseline station for future intercalibration of Th and Pa (see Section 5.11 below). One set of seawater samples collected prior to GEOTRACES was analyzed at the University of Minnesota. The other, collected subsequent to the GEOTRACES cruise, was processed at WHOI. Procedures employed in processing and analyzing the second set of samples are described in an accompanying paper (Auro et al. this volume). Methods used at the University of Minnesota are described below.

4.1 Minnesota chemical procedures

Two-liter seawater samples were collected in July 1993 and analyzed for ^{230}Th later that year. Chemical purification of Th for these samples was performed either at the U. Minnesota (by J. Hoff) or at WHOI (by K. Buesseler, R.A. Belastock, and S.B. Moran). Purified Th separates were all analyzed at the U. Minnesota using TIMS. The University of Minnesota group also participated in the GEOTRACES intercalibration but using techniques detailed in a companion paper (Wang et al., in preparation for this volume) that had evolved substantially since 1993.

Seawater samples were processed in clean rooms at WHOI or Minnesota using ultrapure reagents following similar separation procedures. Seawater samples were transferred on board the ship into acid-cleaned polyethylene bottles. Upon return to the lab the samples were weighed, acidified by addition of 1 ml of concentrated HCl, and spiked with ~50 pg of ^{229}Th tracer and ~6 mg of Fe carrier. The sample/spike mixture

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3 was gently heated ($\sim 75^\circ \text{C}$) for 2 or more days to ensure sample/spike equilibration.
4
5 Thorium was co-precipitated with $\text{Fe}(\text{OH})_3$ by addition of NH_4OH until the seawater
6
7 solution reached a pH of 8 to 9 (Goldberg et al. 1963). The precipitate was allowed to
8
9 settle for at least one day and then either vacuum filtered onto $0.45 \mu\text{m}$ Durapore filters
10
11 (WHOI) or isolated using centrifugation (Minnesota). The filtrate was then dissolved in
12
13 $\sim 1 \text{ ml}$ of 8N HNO_3 and transferred to a Teflon vial for separation chemistry. Chemical
14
15 separation procedures for Th were modified from Chen et al. (1986) and have been
16
17 summarized elsewhere (Moran et al. 1997). Two anion-exchange columns (BioRad AG
18
19 1-X8, 100-200 mesh) were used to separate Th. The initial column contained an $800 \mu\text{l}$
20
21 resin bed and the second column had a $175 \mu\text{l}$ resin bed. For both columns the same
22
23 procedures were used, including preconditioning with 3 column volumes (cv) of 6N HCl ,
24
25 3 cv of H_2O and 3 cv of 7N HNO_3 . Fe was eluted with 3 cv of 7N HNO_3 and then the Th
26
27 fraction was collected by adding 3 cv of 6N HCl . The Th fraction was then dried and re-
28
29 dissolved in $\sim 200 \mu\text{l}$ of 7N HNO_3 for the small volume clean-up columns. The final Th
30
31 fraction was dried and taken up in 1 N HNO_3 for loading for TIMS.
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39 4.2 Blanks: Chemistry blanks were processed with each batch of samples by
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41 adding Th spike, Fe carrier, $\sim 250 \text{ ml}$ of de-ionized water, and 1 ml of concentrated HCl
42
43 to a sample container and gently heating ($\sim 75^\circ \text{C}$) along with seawater samples for a
44
45 minimum of 2 days. Precipitation and separation chemistry followed that described
46
47 above. Blanks for the 2-liter samples used to generate the baseline profile were $0.23 \pm$
48
49 0.33 fg ($1.0 \pm 1.5 \times 10^{-5} \text{ dpm}$) ^{230}Th and $8.2 \pm 5.9 \text{ pg}$ ($2.0 \pm 1.6 \times 10^{-6} \text{ dpm}$) ^{232}Th based
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51 on 13 measurements.
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3 4.3 Mass Spectrometry: TIMS Th analyses were performed on a Finnigan
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5 MAT 262 RPQ mass spectrometer, using a secondary electron multiplier in pulse
6
7 counting mode. All filaments used for analyses were first loaded with a small amount of
8
9 colloidal graphite and analyzed by TIMS at temperatures of 1750° to 2050°C to monitor
10
11 signals in the Th mass range. Filaments with count rates at masses 229 and 230 that were
12
13 significantly higher than multiplier dark noise were discarded from use. All filaments had
14
15 measurable 232 count rates. Those with rates higher than 500 cps were discarded. This
16
17 count rate is equivalent to the beam produced by about 2 pg ($\sim 5 \times 10^{-7}$ dpm) of ^{232}Th .
18
19 Correction for the 232 filament blank was included in the total chemistry blank. Samples
20
21 were loaded onto the acceptable filaments using $\sim 0.1\text{N HNO}_3$, evaporated to dryness,
22
23 covered with colloidal graphite, and then heated using a current of ~ 2 amps for ~ 20
24
25 seconds to dry the graphite. Typical ion yields were 0.5%.

32 33 34 **5.0 Assessment**

35
36 5.1 Spiking at Sea: Sample integrity during transportation and storage was
37
38 tested in two steps. The first step determined whether any bias was introduced by
39
40 deferring the addition of chemical yield monitors until the samples (filtered and acidified)
41
42 had been delivered to the shore-based laboratory. The second step, described below,
43
44 determined whether or not there was a detectable change in the concentration of
45
46 dissolved nuclides during storage of filtered and acidified (but unspiked) samples for
47
48 periods of up to three years. All of the water used for these tests was drawn from the
49
50 1000-liters of seawater, collected from 2000 m at the BATS site, and homogenized in the
51
52 SAFe tanks (Section 2.3.1) for the main intercalibration effort.
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3 The first test consisted of three sets of samples, 12 in total. Set 1 included four
4 samples that were spiked at sea. Nuclides were concentrated at sea by co-precipitation
5 with FeCl₃ carrier following the LDEO procedure described above. Iron hydroxide was
6 concentrated aboard the ship by centrifugation and returned to LDEO in the centrifuge
7 tubes. Set 2 included 5 samples that were spiked at sea, using the same isotope yield
8 monitors that were used for Set 1, and then shipped to LDEO before the addition of FeCl₃
9 carrier or any further processing. Set 3 included three samples that were not spiked until
10 they were processed at LDEO. Set 3 was processed at the same time as Set 2. Sample
11 sizes for Sets 2 and 3 were determined by weighing samples at LDEO. Sample sizes for
12 Set 1 were of necessity determined by measuring their volume in a graduated cylinder at
13 sea and, therefore, those sample sizes may have a larger but indeterminable uncertainty.
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29 Concentrations of ²³²Th (Figure 1A) and ²³⁰Th (Figure 1C) are systematically
30 lower in the samples for which the nuclides were concentrated by co-precipitation at sea
31 (samples 1 - 4 in Figure 1). The reason for this offset is unknown. Concentrations of
32 ²³²Th and ²³⁰Th in samples spiked at sea but co-precipitated at LDEO (samples 5 - 9) are
33 indistinguishable from concentrations in samples that were spiked after the cruise, in the
34 lab at LDEO (samples 10 - 12). Therefore, we rule out an artifact created by delaying the
35 spiking until the samples were processed at LDEO. An overestimation of the size of
36 samples processed at sea (1 - 4) could account for lower Th concentrations in those
37 samples, but the uniformity of ²³¹Pa concentrations throughout all of the samples (Figure
38 1B) is inconsistent with this explanation, and it is unlikely that sample volumes would be
39 in error by as much as 10%. Leaching of Th from sample containers during transport and
40 storage is unlikely as well, because this would require the source of leached Th to have a
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3 $^{232}\text{Th}/^{230}\text{Th}$ ratio similar to that of seawater, roughly two orders of magnitude smaller
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5 than the ratio in average crustal material, a more likely source of contamination.
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8 Furthermore, clean but unused sample containers were filled at sea with distilled water
9
10 and processed as blanks. They showed no evidence for leaching of Th from the container
11
12 material. Iron carrier and ammonia used to process samples at sea came from the same
13
14 batches as the reagents used at LDEO, so contamination of these reagents is also unlikely
15
16 to be responsible for the observed differences.
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19
20 Unfortunately, variability among results for the main intercalibration exercise
21
22 using samples from the 2000-m SAFe tank was so large (see discussion below pertaining
23
24 to Figure 7) that they are of little help in determining which set of results in Figure 1 is
25
26 more accurate. Nevertheless, if one makes an informed subjective decision about the
27
28 consensus values for ^{232}Th and ^{230}Th based on the main intercalibration samples (see text
29
30 below related to Figure 7), then results from sets 2 and 3 (samples 5-12 in Figure 1) are
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32 more consistent with the consensus values than are the results from Set 1.
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35
36 Whatever the cause of the lower Th concentrations in samples co-precipitated at
37
38 sea (Set 1), results from this test demonstrate that Th is not lost from seawater samples
39
40 acidified to pH ~1.7 during transport from the ship to the shore-based lab.
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43 5.2 Sample Storage: Analysis of samples collected during KN193-6 showed
44
45 no systematic trend in concentration of ^{230}Th , ^{232}Th or ^{231}Pa over a three-year period
46
47 following collection in July 2008 (Figure 2). Thus, we conclude that samples may be
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49 stored for a period of at least three years without loss of Th or of Pa by precipitation or by
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51 sorption to container walls, provided that samples are filtered and acidified to pH ~ 1.7,
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53 and provided that appropriate corrections for ingrowth are made.
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Corrections for ^{230}Th ingrowth must be made if samples are stored for more than a few months (see Figure 8 in Robinson et al. 2004). Concentrations of dissolved ^{230}Th in seawater are on the order of 0.01% of the level that would occur if ^{230}Th was in radioactive secular equilibrium with ^{234}U (this is an approximate value as the concentration of ^{230}Th increases with depth in the water column whereas the concentration of its parent, ^{234}U , is relatively uniform throughout the ocean). Low concentrations of dissolved ^{230}Th in the ocean are a consequence of its high affinity for sorption to particle surfaces. Dissolved ^{230}Th concentrations in filtered and acidified samples increase at a rate of about $0.027 \text{ dpm m}^{-3} \text{ yr}^{-1}$ (or $\sim 0.57 \text{ fg kg}^{-1} \text{ yr}^{-1}$) due to ingrowth and the elimination of these removal processes (the actual rate varies from sample to sample depending on the salinity and concentration of uranium). The impact of ingrowth is evident in Figure 2C where concentrations of ^{230}Th uncorrected for ingrowth (grey symbols) diverge from the corrected values (black symbols) by an amount that increases with storage time.

Ingrowth corrections were made assuming a constant ratio of dissolved uranium to salinity (Chen et al. 1986) and the measured salinity at each sample location (often using the CTD results rather than bottle salinity). However, there is evidence that the U/Salinity ratio may not be constant in some cases (Owens et al. 2011; Robinson et al. 2004). Errors introduced by assuming a constant U/Salinity ratio will be small for short sample storage times, but it may be necessary to measure the concentration of uranium when making ingrowth corrections for ^{230}Th if samples are stored for long periods of time.

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3 Here we corrected concentrations of dissolved ^{231}Pa for ingrowth as well although
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5 the correction, expressed as percent of the initial concentration, was much smaller than
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7 for ^{230}Th . Ingrowth imposes a smaller factor for ^{231}Pa because its concentrations in the
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9 deep sea are on the order of 0.2% of those that would occur in the case of radioactive
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11 secular equilibrium with ^{235}U . Although still low, this is significantly greater than for
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13 ^{230}Th (~0.01%), reflecting the lower affinity of Pa for sorption to most particulate phases
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15 (excepting biogenic opal and some metal oxide phases, e.g., Anderson et al. 1983a;
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17 Anderson et al. 1983b). After three years of storage the ingrowth corrections were still at
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19 about the level of the overall analytical uncertainty in the measurements (see grey
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21 symbols in Figure 2B).
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27 5.3 Niskin vs. GO-Flo: The storage experiment offered an opportunity to test
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29 for contamination by sampling with standard Niskin bottles. The primary samples for the
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31 storage tests were taken from the SAFe tanks, which were filled using five casts of the
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33 U.S. GEOTRACES rosette where all 24 GO-Flo bottles were tripped at a depth of 2000
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35 m (Section 2.3.1). Results from two samples of seawater collected at 2000 m using the
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37 Niskin rosette and analyzed as part of this study are shown as open circles in Figure 2.
38
39 Neither Niskin sample shows evidence for contamination. If anything, concentrations of
40
41 ^{230}Th and ^{232}Th are slightly lower in the Niskin samples, although the concentrations
42
43 overlap within the reported analytical uncertainty. The lower Th concentrations in Niskin
44
45 samples might indicate the loss of Th by sorption to bottle walls, but it could also reflect
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47 natural variability created by the fact that the Niskin and GO-Flo casts were taken on
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49 different days and therefore do not reflect true replicate samples. A larger number of
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51 analyses would be necessary to evaluate this potential bias more rigorously.
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3 5.4 Spike equilibration: The corollary to the question “how long can samples
4 be stored?” is “how long must yield monitors equilibrate before samples can be
5 processed?” Isotopes added in acidic solutions as yield monitors would be expected to
6 have a chemical speciation very different from that of isotopes of the same element
7 dissolved in seawater. Acidifying the seawater sample prior to addition of the yield
8 monitors may partly compensate for this, but it is still unclear if the speciation is
9 fundamentally different, or how much time is required for isotopes added as yield
10 monitors to equilibrate with the isotopes present naturally in the sample.
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22 Traditionally, the authors and other investigators have allowed at least a day for
23 spikes to equilibrate with sample isotopes before extracting the nuclides from bulk
24 seawater, and this was the case for samples coprecipitated at sea during the first
25 intercalibration cruise (Section 5.1). To test whether this is sufficient, or even necessary,
26 a suite of 1-liter aliquots of deep-sea water were spiked with a dual ^{229}Th and ^{236}U spike
27 at the University of Oxford. Uranium and Th were extracted from the seawater by co-
28 precipitation with iron hydroxide at various times after the spikes were added, purified,
29 and analyzed by MC-ICP-MS. Results of these analyses (Figure 3) indicate that the
30 traditional one-day spike equilibration should be sufficient. If anything, the isotopes of
31 “insoluble” thorium (Figure 3A, B) seem to reach their final values very rapidly, whereas
32 uranium (Figure 3 C) may take up to a day to equilibrate, as indicated by the asymptotic
33 approach toward final values of ^{238}U concentration. Rapid equilibration with a Th
34 isotope yield monitor was also observed in a study of ^{234}Th (Wei and Hung 1993).
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52 Together, these results indicate that the common practice of allowing one day between
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3 spiking and co-precipitation is more than adequate for Th, but a full day may be
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5 necessary for spike equilibration when measuring U concentrations in seawater.
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8 5.5 Test of different filters: Historically, various types of filter material have
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10 been used by different labs when determining concentrations of dissolved trace elements
11 and radionuclides, but no systematic test had been made to compare results for Th and Pa
12 using different filter materials. Participants in this intercalibration felt that such a test
13 was necessary to establish whether all labs should use the same type of filters, or if each
14 lab could use its preferred filter type without introducing a significant bias.
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22 Filter tests were conducted by tripping all 24 bottles on the Niskin rosette at a
23 single depth. The entire contents of each Niskin bottle were filtered through a single
24 filter, except in a few cases where Nuclepore™ membranes had to be changed in mid
25 filtration due to clogging. Tests were conducted at two locations during KN193-6: (1) at a
26 depth of 2000 m at the BATS station, corresponding to the location of water collected for
27 intercalibration, and roughly corresponding to the mid depth minimum in particle
28 concentration, and (2) at a depth of 635 m at a site over the continental slope (37° 2.0' N,
29 74° 23.87' W) where the concentration of particles was expected to be greater than at the
30 BATS site. Particle concentrations at these two sites were expected to span the
31 approximate range of concentrations in subsurface waters on open-ocean sections.
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33 Although dry weights of particle mass were not determined, transmissometer profiles
34 collected during full depth casts at each station confirmed that the 2000-m depth at the
35 BATS station was close to the mid depth particle minimum, and that concentrations of
36 particles at the slope station were much greater than at the BATS station (data not
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3 shown). A smaller selection of filter types was compared at the slope station than at the
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5
6 BATS site.

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8 Filtration through all types of filters reduced the concentrations of ^{230}Th and
9
10 ^{232}Th substantially compared to unfiltered samples at both sites (Figures 4 and 5).
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12 Concentrations of ^{231}Pa were not significantly lower in filtered samples, consistent with
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14 the lower affinity of Pa for particles compared to Th. A larger fraction of the total
15
16 (unfiltered) Th concentration was associated with filterable particles (i.e., unfiltered
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18 minus average filtered Th concentration) at the slope station than at the BATS site (18%
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20 of ^{230}Th at BATS; 37% of ^{230}Th at the slope station), as expected given the greater
21
22 abundance of particles at the slope site. A larger fraction of the total concentration of
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24 ^{232}Th was associated with filterable particles than for ^{230}Th (31% of ^{232}Th at the BATS
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26 site versus 18% for ^{230}Th), which is also expected given that there is no in situ production
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28 of dissolved ^{232}Th by radioactive decay of a parent isotope.
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34 Among the standard materials that were tested as individual filters of 47 or 90-
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36 mm diameter (Nuclepore, QMATM, Supor) there was no detectable difference in the
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38 concentrations of any of the three nuclides (Figures 4 and 5). Furthermore, there was no
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40 systematic difference between pore diameters (0.4 μm vs. 1.0 μm for Nuclepore; 0.45 μm
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42 vs. 0.8 μm for Supor). This latter comparison suggests that most of the particulate Th
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44 that is retained by 0.4 μm pore diameter filters, as estimated by comparing Th
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46 concentrations in filtered and in unfiltered samples, resides on particles greater than 1.0
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48 μm in diameter. By contrast, concentrations of Th (both ^{232}Th and ^{230}Th) were
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50 systematically (albeit slightly) lower in samples filtered through Osmonics cartridges
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52 with 0.22 μm pore diameter membranes. Whether the lower Th concentrations in
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3 samples filtered through the Osmonics cartridges reflect the presence of a significant
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5 fraction of Th on particles between 0.2 and 0.4 μm in diameter, or reflect sorption of
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7 dissolved Th to the filter cartridges, cannot be assessed with the existing information.
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10 The Osmonics cartridge used here was of the same type used to filter 1000 liters of
11
12 seawater when filling the SAFe tanks (1.6 square meters effective surface area; see
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14 above), and thus it was much larger than the filters and filter holders used for the other
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16 filtration tests (either 47-mm or 90-mm diameter). Therefore, dissolved Th may have
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18 been lost by sorption to the Osmonics cartridge or to its filter membrane. Also, the dead
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20 volume inside the Osmonics cartridge was more than one liter, and it is possible that the
21
22 cartridge was flushed insufficiently before sample collection commenced. Although
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24 cartridges as large as this might not be used to filter routinely from Niskin bottles,
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26 intermediate-sized filtration cartridges may be used on certain cruises to filter all samples
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28 consecutively in a clean van before distributing the unfiltered samples. This is an
29
30 excellent way to filter large volumes of samples under the cleanest possible conditions,
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32 similar to the method used for trace metals. Consequently, we recommend that future
33
34 studies should be carried out to determine the fraction of Th in the deep sea that resides
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36 on particles smaller than 0.4 μm in diameter, and to determine if sampling biases are
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38 introduced when using large filtration cartridges.
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46 Concentrations of dissolved nuclides obtained by filtration through quartz (QMA)
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48 filters were not significantly different from those obtained with the individual (47 or 90
49
50 mm) membrane filters for any of the three nuclides (Figure 4). This is somewhat
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52 surprising because Pa is known to have a high affinity for sorption to glass and to
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54 amorphous silica (e.g., many labs use silica gel to adsorb ^{233}Pa when milking it from
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3 ²³⁷Np). Further testing would be necessary before we would confidently conclude that
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6 QMA filters may be used to determine concentrations of dissolved Th and Pa on
7
8 GEOTRACES cruises.
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10 5.6 Intercalibration - Sediment analyses: The most comprehensive assessment
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12 of intercalibration among labs that measure Th and Pa was provided by the analysis of
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14 siliceous ooze sediments. Concentrations of ²³²Th agreed nearly as well between labs as
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16 between replicated analyses from a single lab with the exception of two high values
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18 reported initially by Lab 15 using alpha spectrometry (Figure 6A). When Lab 15
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20 reanalyzed the sample by ICP-MS (hollow grey symbol in Figure 6A) the ²³²Th
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22 concentration fell into line with the values from other labs. The source of the high ²³²Th
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24 values remains undetermined, but it may involve an insufficient tail correction in the
25
26 alpha spectrum. Excluding these high values, the mean and standard deviation of lab
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28 mean values (i.e., averaging for each lab where replicate results were reported) is $0.059 \pm$
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30 0.005 dpm/g (0.242 ± 0.021 $\mu\text{g/g}$; 9% 1 sigma).
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36 Reported concentrations of ²³⁰Th vary by as much as 20% (full range of values)
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38 among labs (Figure 6C), more than would be expected given the agreement for ²³²Th.
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40 That is, the variability does not seem to be due to differences among labs in the
41
42 calibration of the ²²⁹Th yield monitors, with the possible exception of Lab 16 for which
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44 the reported ²³²Th and ²³⁰Th values were each about 20% less than the respective means
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46 of the other labs. Including results from all labs, the mean and standard deviation of lab
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48 mean values is 5.38 ± 0.42 dpm/g (7.7% 1 sigma). Excluding the low mean values from
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50 Labs 15 and 16 the mean and standard deviation of the lab mean values for the remaining
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52 11 labs becomes 5.53 ± 0.22 dpm/g (4% 1 sigma). Agreement for ²³¹Pa (Figure 6B) is
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3 similar to that for ^{230}Th with the exception of one lab (5) that reported a ^{231}Pa
4 concentration that was 55% of the mean value from other labs. Excluding this one low
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6 value, the mean and standard deviation of lab mean values is 1.25 ± 0.07 dpm/g (6% 1
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8 sigma).
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12 Discussion of these results among group participants raised questions about the
13 homogeneity of the siliceous ooze, given that variability among results from a single lab
14 for ^{230}Th (e.g., Lab 1) is similar to the variability between all labs (Figure 6C). Intra-lab
15 variability is not the result of spike calibration or other systematic errors. If a large
16 portion of the intra-lab variability evident in Figure 6C is due to sample heterogeneity,
17 then this is a concern for anyone measuring these nuclides in sediments. The siliceous
18 ooze was homogenized more thoroughly than is common during the analysis of marine
19 sediments in anticipation that it would be used for intercalibration. It would be a concern
20 if variability among aliquots of as much as 20% survived this level of homogenization.
21
22 In response to this concern about sample heterogeneity, the LDEO group prepared a
23 mixed isotope solution containing ^{232}Th , ^{230}Th and ^{231}Pa that could be analyzed as a
24 working standard. Results from that solution are described in Section 5.9 below.
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41 5.7 Intercalibration - Seawater analyses: Initial intercalibration results from
42 seawater collected during the first cruise (KN193-6, BATS site) revealed unexpected
43 levels of contamination by sources of ^{232}Th . Further investigation indicated that
44 contamination was sometimes derived from sample containers and sometimes from
45 laboratory procedures (see also accompanying papers by Wang et al., in preparation for
46 this volume, Andersson and Schöberg this volume; Auro et al. this volume; Roy-Barman
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3 this volume). As noted above, there was no indication of contamination from the Niskin
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6 sampling system used at the BATS site.
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8 Triplicate analyses by Lab 7 of each of three samples from 2000 m at the BATS
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10 site indicated that Th contamination was present in sample containers (Figure 7A).
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12 Internal consistency within each set of triplicate samples, together with the large
13
14 differences in ^{232}Th concentration between individual samples, is consistent with
15
16 contamination of the sample containers rather than the random introduction of
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18 contamination during processing of the samples. After identifying this problem Lab 7
19
20 analyzed samples that had been collected in a different set of containers (smaller
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22 containers also provided by Lab 7) and obtained ^{232}Th concentrations (hollow grey
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24 symbols in Figure 7A) close to those from other labs. By plotting $^{232}\text{Th}/^{230}\text{Th}$ ratios vs.
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26 ^{232}Th concentrations Lab 7 was able to establish that the source of contamination had a
27
28 crustal $^{232}\text{Th}/^{230}\text{Th}$ ratio (not shown). Lab 8 also encountered ^{232}Th contamination. Their
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30 blanks were so large that they precluded reporting ^{232}Th concentrations during the first
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32 set of intercalibration tests (BATS seawater).
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38 Contamination introduced a smaller but still significant bias in ^{230}Th
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40 concentrations, as is evident by circled values from Lab 7 in Figure 7C, which are greater
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42 than ^{230}Th concentrations reported by other labs. These results were flagged as
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44 contaminated by Lab 7 at the time that they were first reported.
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48 An important lesson from this experience is that sample containers may be a
49
50 significant source of contamination. Blanks must be determined using sample containers
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52 that are cleaned and handled by the same procedure as the samples. A weakness of this
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54 intercalibration exercise is that empty sample containers from each lab were not
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3 processed at sea as blanks. We recommend that blanks be processed regularly at sea. On
4
5 U.S. GEOTRACES section cruises, for example, a seawater blank is processed at each
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7 station by adding 2 to 3 liters of Milli-Q water to a sample container and treating it as a
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9 sample.
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13 Discovery of the unexpected contamination problems described above, as well as
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15 others that were later identified, stimulated an extensive discussion and collaboration
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17 among participants to locate sources of contamination and to eliminate them.
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20 Investigators must be cautious about checking blanks, especially when changes are made
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22 in analytical methods. For example, the LDEO authors of this manuscript discovered that
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24 a new source of high purity iron carrier was heavily contaminated with Th. Repeated
25
26 extraction of iron from 7N HCl into isopropyl ether eventually reduced the
27
28 contamination. Similarly, the WHOI author of this manuscript discovered that some
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30 batches of ion-exchange resin contain large amounts of Th, and that some labware
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32 adsorbs Th irreversibly, leading to low chemical yields. Those experiences are described
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34 in greater detail in a companion manuscript (Auro et al. this volume). The lesson here is
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36 that the clean-up procedure that may be acceptable for one batch of Fe carrier or resin
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38 may be inadequate for another.
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44 As challenging as the contamination issue has proven to be, it can be (and has
45
46 been) addressed through systematic testing of individual steps in the collection and
47
48 analysis of seawater samples. A more perplexing problem that was discovered during the
49
50 seawater intercalibration involves poor reproducibility among results from a single lab.
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52 Sometimes this is manifest as low concentrations that cannot be explained by
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54 contamination (e.g., see ^{230}Th results from Labs 2 and 5 in Figure 7C), and sometimes as
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3 high concentrations for ^{231}Pa in samples where no corresponding contamination of Th
4 was evident (e.g., see ^{231}Pa results for Labs 2 and 8 in Figure 7B). The source(s) of this
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high concentrations for ^{231}Pa in samples where no corresponding contamination of Th was evident (e.g., see ^{231}Pa results for Labs 2 and 8 in Figure 7B). The source(s) of this variability is (are) still under investigation.

Analysis of surface waters (from a depth of between 5 and 15 m) is more challenging because of the much lower concentrations of dissolved ^{230}Th and ^{231}Pa (compare Figure 7 panels B and C with Figure 8 panels B and C). Despite the lower levels, surface ^{230}Th concentrations were within reported uncertainty of 0.08 dpm m^{-3} (Figure 8C), except for the circled values from Lab 7, indicating results that were flagged by the investigators as likely suffering from contamination, as well as results from Lab 8, which also reported high ^{232}Th blanks (see above). Similarly, surface concentrations of ^{231}Pa are generally within reported error of 0.05 dpm m^{-3} , except for Lab 2 and Lab 12. Three labs reported surface concentrations of ^{232}Th within reported uncertainty of 0.02 dpm m^{-3} , including the results by Lab 7 from samples collected in smaller sample containers (Figure 8A). Concentrations of ^{232}Th from Labs 2 and 4 are significantly higher, as are those flagged by Lab 7 as likely suffering from contamination (circled in Figure 8A). Although results from only three labs are insufficient to conclude that 0.02 dpm m^{-3} is the correct value for the concentration of ^{232}Th in the near-surface BATS sample, the higher values from other labs would be consistent with the widespread contamination by extraneous sources of Th described above.

Evaluation and testing of Th blanks was still underway at the time of the second intercalibration cruise (KN195-8). Although the extensive efforts by participating labs in response to the results from the first cruise had improved their blank levels substantially, evidence for ^{232}Th contamination can still be found in the results from the second cruise.

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3 Lab 9 flagged high ^{232}Th values in duplicate analyses of one sample of water from 3000
4 m at the SAFE station (circled values in Figure 9A) as having suffered from
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6 contamination. Reproducible results were obtained from duplicate analyses of this
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8 sample, indicating that the contamination likely occurred in the sample container rather
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10 than in the lab procedure. This is reminiscent of container contamination experienced by
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12 Lab 7 during the analysis of samples from KN193-6. The source of the container
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14 contamination is unknown. Containers were cleaned by each lab prior to each cruise and
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16 transported to the ship enclosed in double plastic bags. The SAFE tanks (see above),
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18 located inside a hangar aboard the ship, were enclosed within a temporary structure (tent)
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20 constructed of plastic sheeting to minimize contamination by aerosols. Sample
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22 containers were removed from their bags inside the tent, and then rinsed and filled inside
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24 the tent before rebagging for return to their final destinations. These procedures follow
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26 those adopted for collecting large numbers of replicate seawater samples for trace metal
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28 analysis, which have been found to be free of trace metal contamination. Consequently,
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30 the source of Th contamination in containers remains undetermined. Lab 8 reported that
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32 they were still undertaking tests to reduce their Th blanks at the time these samples were
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34 analyzed, and the variability among the initial results for Lab 8 (Figure 9A, black points)
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36 indicate that their efforts were initially only partially successful. Following further work
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38 to improve their blank, Lab 8 was sent two additional samples that had been archived at
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40 LDEO. Results from those samples (grey points in Figure 9) are in good agreement with
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42 the results from other labs, consistent with the much lower blanks reported by Lab 8 at
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44 the time those samples were analyzed. Note that the results for ^{231}Pa reported by Lab 8
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46 also improved after their ^{232}Th blanks were lowered (grey points in Figure 9B), reflecting
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3 in part the reduced uncertainty in the tail correction for ^{232}Th that remained in the Pa
4 fraction. Lab 10 indicated that their ^{232}Th analyses required large blank corrections,
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6 adding to the variability among their results (Figure 9A). Lab 10 has subsequently
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8 reported that their ongoing methods development, stimulated by their participation in the
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10 GEOTRACES intercalibration, has reduced their ^{232}Th blank level by about a factor of
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12 four since their SAFe samples were analyzed.
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18 Determining concentrations of ^{232}Th in surface waters at the SAFe station was
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20 more challenging than at BATS because ^{232}Th concentrations in surface waters of the
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22 central Pacific are much lower than in the North Atlantic, reflecting the much lower
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24 fluxes of mineral aerosols (dust, the principal source of ^{232}Th in the open ocean) to the
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26 Pacific. The average concentration of ^{232}Th in the surface sample from the SAFe station
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28 (0.0041 ± 0.0009 dpm m^{-3} ; mean and standard deviation of lab mean values; Figure 10A)
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30 is only about 20% of the average surface concentration at the BATS site (0.022 ± 0.004
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32 dpm m^{-3} ; mean and standard deviation of lab mean values excluding values flagged by
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34 Lab 7, Figure 8A). Despite the low concentration of dissolved ^{232}Th , high levels of
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36 contamination are not evident in any of the reported results. Labs 1 and 4 reported ^{232}Th
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38 concentrations below and above the group mean, respectively, by more than their
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40 reported uncertainties. Blank corrections were reported to be substantial for each of these
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42 labs, so the departures from the mean ^{232}Th concentration may reflect an inaccurate
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44 evaluation of the blank (i.e., an overestimation and an underestimation of the blank
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46 correction by Labs 1 and 4, respectively, with a corresponding underestimation of the
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48 propagated uncertainty).
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4 Results from the analysis of deep (3000 m) water from the SAFe station indicate
5 an improvement for ^{230}Th and for ^{231}Pa relative to results from the first cruise (compare
6 Figure 7 panels B and C with Figure 9 panels B and C). The mean and standard
7
8 deviation of lab mean values for ^{230}Th at 3000 m (Figure 9C) is 1.16 ± 0.07 dpm m^{-3} (6%
9 one sigma). Labs 1, 9 and 10 reported average ^{230}Th concentrations that deviated from
10 the group mean by more than their reported analytical uncertainties, with average values
11 for Labs 1 and 10 falling below the group mean and for Lab 9 falling above. Despite
12 these differences, without eliminating any data the average ^{230}Th concentration reported
13 by each lab fell within two sigma of the mean of all labs that reported results for the
14 SAFe 3000-m sample. For ^{231}Pa the situation is nearly as good, where the mean and
15 standard deviation of lab mean values is 0.44 ± 0.03 dpm m^{-3} (7% one sigma) if one
16 anomalously high value from Lab 2 and one low value from Lab 12 are excluded (Figure
17 9B).
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34 As noted above, the analysis of surface waters for ^{230}Th and ^{231}Pa is challenging
35 because of the low concentrations that are present, and this is reflected in the variability
36 of ^{230}Th concentrations reported by Lab 2 and by the low ^{231}Pa concentration reported by
37 Lab 4 (Figure 10). Nevertheless, excluding these results the generally good agreement
38 among labs indicates that the low concentrations of these nuclides can be measured in
39 surface waters and thereby contribute to the study of trace element scavenging and
40 removal at shallow depths. Collection and analysis of replicate samples from surface
41 waters are recommended to improve confidence in the results.
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53 5.8 Intercalibration - Particle analyses: Determining concentrations of
54 particulate ^{230}Th , ^{232}Th and ^{231}Pa requires volumes of seawater greater than can be
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3 obtained using standard Niskin bottles because only a small fraction of the total Th and
4 Pa in the open-ocean water column is associated with particulate material (compare
5 results for filtered and unfiltered samples in Figure 4). Consequently, as described above,
6 particles were collected for intercalibration by filtering 400 to 600 liters of water using in
7 situ pumps. Two locations were selected: (1) a depth of 2000 m at the BATS site,
8 which allows particulate results to be compared with those from the intercalibration of
9 dissolved nuclides (Figure 7), and (2) at a depth of 80 m at a station on the continental
10 slope (37° 01.68' N, 74° 23.88' W). The slope site provided an opportunity for
11 intercalibration in shallow water, where concentrations of dissolved ^{230}Th and ^{231}Pa are
12 much lower than in the deep open ocean, but where concentrations of particulate material
13 are also likely to be much greater than in the open ocean. A one-quarter section of a filter
14 from each site was distributed to each participating lab, along with a dip blank (see
15 Section 2.3.3 above for methods) from each site.
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34 Five labs reported concentrations of particulate Th for the 2000-m samples at the
35 BATS site while six labs reported concentrations of particulate ^{231}Pa (Figure 11). The
36 average particulate ^{230}Th concentration at 2000 m is $\sim 0.05 \text{ dpm m}^{-3}$ for four of the five
37 reporting labs with Lab 9 reporting a substantially lower value (Figure 11A).
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43 Investigators at the Woods Hole Oceanographic Institution determined that variability
44 among subsamples from Supor filters deployed on different pumps, but sampling at the
45 same time and location on the pump rosette, can be as large as 20% (see Section 2.3.3
46 above and Maiti et al. this volume). Therefore, the low value reported by Lab 9 may
47 reflect a real difference between filter subsamples rather than an analytical bias. Setting
48 aside the result from Lab 9, the average concentration of particulate ^{230}Th at 2000 m
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3 (Figure 11A) is about 10 to 12% of the average concentration of dissolved ^{230}Th (Figure
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6 7C). This is somewhat less than, but nevertheless consistent with, the difference between
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8 filtered and unfiltered samples from 2000 m analyzed as part of the comparison of
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10 different filter membranes (estimated as 18% of ^{230}Th associated with filterable particles;
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12 see Section 5.5 above and Figure 5C).
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16 Given the potential for substantial heterogeneity within and between filters, it is
17
18 informative to examine particulate $^{232}\text{Th}/^{230}\text{Th}$ ratios, which should be relatively
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20 insensitive to the heterogeneous distribution of particulate material over the surface of a
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22 filter provided that there is no fractionation by particle composition created by
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24 heterogeneous particle deposition on the membrane surface. We are aware of no
25
26 evidence that such fractionation occurs. Here, in contrast to particulate Th
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28 concentrations, where the value reported by Lab 9 was low (Figure 11A), we find that the
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30 $^{232}\text{Th}/^{230}\text{Th}$ ratio reported by Lab 9 is consistent with the results from three other labs,
31
32 allowing for the possibility that the amount of particulate material deposited on the
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34 section of filter analyzed by Lab 9 was below average. By contrast, the $^{232}\text{Th}/^{230}\text{Th}$ ratio
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36 reported by Lab 2 is greater than for the other four labs (Figure 11B). Whether this
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38 reflects an unrecognized ^{232}Th blank or another factor affecting the results from Lab 2 is
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40 unknown.
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46 Concentrations of particulate ^{231}Pa were below the limit of detection for three of
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48 the six labs that reported results (Figure 11C). Results from the remaining three labs are
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50 internally consistent, and indicate a concentration of particulate ^{231}Pa of about 0.0018
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52 dpm m^{-3} (Figure 11C), or about 1% of the concentration of dissolved ^{231}Pa at the same
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54 location (Figure 7C). These results are consistent with previous studies, which have
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3 determined that the fractionation factor representing the preferential scavenging of Th
4 from seawater relative to Pa, defined (Anderson et al. 1983b) as the ratio {(particulate
5 $^{230}\text{Th}/\text{particulate } ^{231}\text{Pa}$)/(dissolved $^{230}\text{Th}/\text{dissolved } ^{231}\text{Pa}$)}, has a value of about 10 at
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11 2000 m at the BATS site (Moran et al. 2002).

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13 Concentrations of particulate ^{230}Th in surface waters at the slope station (Figure
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15 12A) are only about one quarter of the average reported for 2000 m at the BATS site
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17 (Figure 11A), and the lower concentrations may contribute to the poorer reproducibility
18
19 there among labs. In particular, the particulate ^{230}Th concentration reported by Lab 1 was
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21 only about one third of the average concentration reported by the other four labs,
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23 departing from the mean by more than can be attributed to heterogeneous particle
24
25 collection (Section 2.3.3). The low particulate ^{230}Th concentration, together with the high
26
27 particulate $^{232}\text{Th}/^{230}\text{Th}$ ratio reported by Lab 1 (Figure 12B), suggests a problem with the
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29 ^{230}Th . Five labs reported concentrations of particulate ^{231}Pa for the shallow slope site.
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31 For three of the five labs the particulate ^{231}Pa concentration was below their detection
32
33 limit (note the position of zero concentration on the y-axis in Figure 12C). For the two
34
35 labs that reported particulate ^{231}Pa concentrations significantly greater than their detection
36
37 limits (Labs 1 and 2), the reported values differ by about a factor of three, too large to be
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39 attributable to sample heterogeneity. Determining concentrations of particulate ^{230}Th and
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41 ^{231}Pa in surface waters is clearly challenging, and further work is needed to develop
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52 methods for the reliable determination of these concentrations at the low levels that occur
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60 in surface waters.

53 **5.9 Intercalibration - Working Standards:** Following discussion of the issues
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55 identified above, the participants determined that it would be beneficial to prepare and
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3 distribute a working solution that could be analyzed repeatedly to monitor the
4 performance of each lab while also providing a further intercalibration of lab methods. A
5 mixed solution was prepared at LDEO containing ^{232}Th , ^{230}Th and ^{231}Pa in quantities
6 such that 0.5 g of the solution contained approximately the same amount of each nuclide
7 as would be present in 10 liters of seawater recovered from 2000 m at the BATS site.
8 The solution was then split into two aliquots, and ^{229}Th was added to one of the two
9 portions. The portion without ^{229}Th , labeled SW STD 2010-1, was to be treated as an
10 unknown and processed as for any other intercalibration. Each lab was asked to add its
11 own yield monitors (^{233}Pa and ^{229}Th) and process the standard through normal chemical
12 procedures. Comparison of results was expected to yield information about overall
13 consistency among labs, including blanks and the calibration of yield monitors.
14 Participants were asked to process the portion containing ^{229}Th (labeled SW STD 2010-2)
15 without adding yield monitors to determine Th atom ratios in the solution as a measure
16 primarily of the mass biases inherent in mass spectrometers, although blanks would affect
17 these ratios as well.

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39 Different procedures were used among the labs that have submitted results for the
40 working solutions. At one extreme, SW STD 2010-1 was added to several liters of
41 distilled water that was then processed as a full seawater sample. At the other extreme,
42 the analyst simply spiked a weighed aliquot of the SW STD 2010-1 solution, which was
43 then evaporated, reconstituted and injected directly into the mass spectrometer. Results
44 are reported in Figure 13 without distinction among the types of procedures applied.
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53 Also, participating labs have been renamed here. That is, Lab A in the results for the
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3 working solutions is not Lab 1 in the results for seawater, sediments and particles, and so
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6 forth.

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8 Among the eight labs that have submitted results to date for SW STD 2010-1,
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10 concentrations of ^{230}Th agreed within the uncertainties reported by each lab (Figure 13A).
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12 The same is true for ^{232}Th (Figure 13B) although it would be desirable to have more
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14 results from each lab to assess the overall internal consistency. For ^{230}Th , the average
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16 and standard deviation of the lab mean values is 247 ± 3 pg/g, from which it can be
17
18 inferred that the calibration of yield monitors for Th is internally consistent among labs to
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20 within about 1%. In principle, if other factors (e.g., blanks) can be controlled, then it
21
22 should be possible to constrain global variability in the concentration of ^{230}Th at nearly
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24 this level of precision.
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29 In light of the good agreement among labs that reported concentrations of ^{230}Th
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31 and ^{232}Th for SW STD 2010-1, it is surprising that labs differ by as much as they do in
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33 results for SW STD 2010-2 (pre-spiked with ^{229}Th). The largest reported uncertainties
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35 were experienced for $^{230}\text{Th}/^{229}\text{Th}$ (Figure 14A) and $^{232}\text{Th}/^{230}\text{Th}$ (Figure 14C) ratios
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37 because ^{230}Th was present in the lowest abundance among the three Th isotopes.
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39 Differences between labs for these ratios are just at the limit of exceeding reported
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41 uncertainties, so it is difficult to establish their significance. Nevertheless, among the
42
43 nine labs reporting results for SW STD 2010-2 the mean and standard deviation of lab
44
45 mean $^{230}\text{Th}/^{229}\text{Th}$ atom ratios is 0.01338 ± 0.00022 (1.6% 1 sigma). Finding a slightly
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47 poorer reproducibility here than for the determination of the ^{230}Th concentration in SW
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49 STD 2010-1 (247 ± 3 pg/g, see above) points to blanks and instrumental errors rather
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3 than spike calibration as the current factor limiting interlab reproducibility in determining
4 the concentration of ^{230}Th .
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8 By contrast, ratios reported for the two more abundant isotopes ($^{232}\text{Th}/^{229}\text{Th}$)
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10 differ by more than the reported uncertainties (Figure 14B). It appears as if there are two
11 clusters of $^{232}\text{Th}/^{229}\text{Th}$ ratios, with results from Labs A, F, G and H falling about the
12 lower value and those from Labs B, C and D falling about the higher value. Lab I
13 reported $^{232}\text{Th}/^{229}\text{Th}$ ratios significantly below the range of values reported by other labs,
14 even though the $^{230}\text{Th}/^{229}\text{Th}$ and $^{232}\text{Th}/^{230}\text{Th}$ ratios from Lab I are consistent with those
15 from other labs. The source of the disagreement remains undetermined. Lab B reported
16 a large number of results, among which the $^{232}\text{Th}/^{229}\text{Th}$ ratios seem to fall within one or
17 the other of the clusters defined by seven of the other labs (excluding Lab I). Keeping in
18 mind that each lab was running the same solution with the same $^{232}\text{Th}/^{229}\text{Th}$ ratio, the
19 differences reported here are interpreted to indicate instrumental bias in the reported atom
20 ratios. The differences appear to be too large to have been caused by a mass bias issue,
21 but may reflect a problem with the calibration of the efficiency of the ion counter relative
22 to the Faraday cups on certain instruments. There is no correlation with the type of mass
23 spectrometer employed (TIMS vs. ICP-MS). Whatever its origin, the consistency among
24 labs for the Th concentrations reported for SW STD 2010-1 (Figure 13) suggests that any
25 instrument bias in atom ratios cancels out when the same bias is first applied to
26 calibration of the ^{229}Th yield monitor used by each lab, and later the yield monitor is used
27 to determine the concentrations of ^{232}Th and ^{230}Th in an unknown.
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53 We note that the ^{229}Th content of SW STD 2010-2 was greater than the amount
54 that some labs typically use when spiking seawater samples. Consequently, some
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3 instruments may have had trouble with the large range in intensities compared to normal
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5 efforts for a well-matched $^{230}\text{Th}/^{229}\text{Th}$ ratio in spiked samples.
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8 5.10 Intercalibration - ^{10}Be in seawater: Beryllium-10 was included with this
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10 group for intercalibration because, historically, ^{10}Be concentrations in seawater and in
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12 sediments have been determined by many of the same labs that determine concentrations
13
14 of Th and Pa. Furthermore, it is hoped that it will be possible at some point in the future
15
16 to use the spatial variability in the ratio of dissolved ^{10}Be (cosmogenic) to ^9Be
17
18 (lithogenic) to constrain rates of mixing in the ocean. In light of this anticipated
19
20 application that will be of interest to the GEOTRACES community, we present
21
22 intercalibration results for dissolved ^{10}Be even though only three labs have reported.
23
24 Seawater from 2000 m at the BATS site was distributed for intercalibration of ^{10}Be .
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26 Results to date are internally consistent within the uncertainties reported by each lab
27
28 (Figure 15). The fraction of ^{10}Be in seawater that is associated with filterable particles is
29
30 expected to be much less than for ^{231}Pa (i.e., $\ll 1\%$; Section 2.2), so no attempt was
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32 made to measure ^{10}Be concentrations in particulate material recovered by in situ
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34 filtration.
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41 5.11 Baseline station - BATS: One of the objectives of the GEOTRACES
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43 intercalibration initiative is to establish “baseline stations” where concentration profiles
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45 of trace elements and their isotopes will be determined by several laboratories repeatedly
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47 throughout the duration of the program. Repeated analyses at these stations provide an
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49 ongoing measure of intercalibration. Baseline stations also provide a location where any
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51 lab can collect samples and expect that the concentrations of trace elements or isotopes
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53 are known reasonably well, at least in the deep ocean where concentrations are not
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3 expected to change significantly on decadal time scales. Here we illustrate this principle
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5 using concentrations of ^{230}Th and of ^{232}Th measured at the BATS site. In the next
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8 section, we describe evidence from the Pacific to illustrate some caveats concerning the
9
10 baseline station concept.
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13 Three of the labs participating in the intercalibration of Th and Pa have generated
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15 concentration profiles of ^{230}Th and of ^{232}Th at the BATS site over a period of 16 years.
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17 Duplicate profiles (i.e., beginning with separate Niskin bottles and processing samples
18
19 independently) collected during KN193-6 (July 2008) and analyzed at LDEO were
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21 filtered at the time of collection through 0.45 μm pore size Supor filters, like those used
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23 in the filtration tests (e.g., Figure 4). The profiles analyzed by the other two labs were
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25 processed without filtration, so this difference in processing must be considered when
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27 comparing results. Two-liter samples of unfiltered seawater collected on July 7, 1993
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29 during the *R/V Weatherbird* cruise 93-06 were analyzed shortly after collection at the
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31 University of Minnesota using TIMS. Samples from > 200 m were collected in 10 liter
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33 Niskin bottles using a CTD/Rosette deployed on a steel hydrowire. Samples from < 200
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35 m were collected either using GO-Flo bottles suspended on a stainless steel wire or on a
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37 separate cast using trace metal clean bottles suspended on a Kevlar line provided by Dr.
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39 Jim Moffet (WHOI). The second set of unfiltered samples was collected in September
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41 2009 by Ken Buesseler on cruise AE0908 of the *R/V Atlantic Explorer*. Samples were
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43 collected via Masterflex Tygon tubing directly from the Niskins into cubitainers. Each
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45 10L cubitainer was rinsed with ~ 50 ml of seawater then filled. Optima grade HCl (1ml
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47 concentrated HCl per liter seawater) was added to each sample in a clean hood.
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3 In each profile the concentration of ^{230}Th increases with depth from the surface to
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5 ~3000 m. Below ~3000 m the concentration of dissolved ^{230}Th decreases toward the
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7 bottom (Figure 16A), as is often observed in the North and South Atlantic Ocean (Luo et
8
9 al. 2010; Moran et al. 2002). Concentrations of total ^{230}Th in the two sets of unfiltered
10
11 samples (U-MN and WHOI) are in good agreement above 3000 m despite the 16 years
12
13 between collection of the two sets of samples. This agreement illustrates the principle
14
15 that baseline stations can be used to provide access to seawater with known
16
17 concentrations of trace elements and their isotopes in stable regions of the ocean interior
18
19 (but see caveats below). Concentrations of dissolved ^{230}Th (Figure 16A; LDEO) exhibit
20
21 good agreement between replicate profiles, and they are slightly but systematically lower
22
23 than concentrations in the unfiltered samples (Figure 16A; U-MN and WHOI). This is to
24
25 be expected given that about 10% of the total ^{230}Th in mid-depth waters resides on
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27 particles that can be removed by filtration (Section 5.5).
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34 Concentrations of total ^{232}Th agree well between the two sets of unfiltered
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36 samples (WHOI and U-MN), and the total ^{232}Th concentration is uniformly greater than
37
38 the dissolved (<0.45 μm ; LDEO) ^{232}Th concentration throughout the water column
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40 (Figure 16B). The first set of replicate samples from KN193-6 (LDEO) experienced high
41
42 ^{232}Th blanks, so the results are not reported here. Below 3000 m the profiles of filtered
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44 and unfiltered samples diverge, more so for ^{232}Th than for ^{230}Th (Figure 16). We
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46 interpret this to indicate the presence of resuspended sediment particles throughout the
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48 lower ~2 km of the water column, which increases the concentration of total Th relative
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50 to mid water column depths. The presence of resuspended sediments is supported by
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52 transmissometer profiles collected during KN193-6, which show beam transmission
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3 decreasing below ~3740 m at the BATS site (not shown). The abundance of resuspended
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5 sediments may vary with time, a factor that should be taken into account when
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7 interpreting Th concentrations in near-bottom waters from the BATS site.
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10 Concentrations of ^{230}Th and of ^{232}Th measured as part of the baseline profile, both
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12 with and without filtration (Figure 16), are consistent with the concentrations reported for
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14 filtered and unfiltered samples as part of the test of different filter membranes (Figure 4).
15
16 They are also consistent with the concentrations of dissolved (Figure 7C) and particulate
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18 (Figure 11A) ^{230}Th determined through the intercalibration exercise, with the caveat that
19
20 there was substantial variability among participating labs in the results that were reported.
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22 Based on the internal consistency among results from different labs and from samples
23
24 collected at different times, we conclude that mid-depth water at the BATS site can serve
25
26 as a baseline station for ongoing intercalibration of ^{230}Th and ^{232}Th .
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32 5.12 Baseline station - Pacific: Results from two locations in the central north
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34 Pacific Ocean suggest that the assumption of constant or known nuclide concentrations in
35
36 deep waters may not always be valid. Results from samples collected in 1994 at the
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38 Hawaii Ocean Time Series (HOT) Station Aloha ($22^{\circ}45'\text{N}$, 158°W) show the
39
40 concentration of ^{230}Th increasing linearly with depth (Figure 17A, Roy-Barman et al.,
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42 1996), as expected for removal by reversible scavenging to sinking particles (Bacon and
43
44 Anderson 1982). Concentrations of dissolved ^{230}Th derived from samples collected at the
45
46 HOT site in February 2002 agree well with the results from samples collected in 1994
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48 (Figure 17A, Francois 2007). However, the concentration profile of ^{230}Th determined
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50 from samples collected during a reoccupation of Station Aloha six months later, analyzed
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52 by the same lab and using the same methods as for the February 2002 samples, depart
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3 significantly from results obtained during two previous occupations (Figure 17A, R.
4 Francois, unpublished data). Surprisingly, whereas concentrations of ^{230}Th in the upper
5 1000 m of the water column agree within error for the two occupations in 2002, results
6 diverge with increasing water depth, differing by nearly 20% for the deepest samples. It
7
8 is unlikely that nearly 20% of the ^{230}Th in deep water could have been removed by
9 scavenging over an interval of about six months, as this would imply removal of ^{230}Th
10 much faster than is inferred from its overall residence time of several decades. Whether
11 the divergence of deepwater results between the February and August 2002 occupations
12 reflects the movement of a deep water mass with lower ^{230}Th concentration into the
13 vicinity of the Aloha station or some other factor remains to be determined.
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27 A similarly disconcerting finding is evident in results from the second
28 GEOTRACES intercalibration cruise at the SAFe site. There, the concentration of
29 dissolved ^{230}Th increases linearly with depth, much as in the 1994 results from the Aloha
30 station (Figure 17B). However, concentrations of dissolved ^{230}Th determined using
31 samples from the 3000-m SAFe tank (Figure 9) fall between 20 and 25% below the value
32 from an equivalent depth in the profile obtained using Niskin bottles (Figure 17B).
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34 Triplicate samples from the SAFe tank (3000 m) and triplicate Niskin samples (3042 m)
35 were each processed at LDEO using the same spike and the same analytical procedure.
36
37 Internal consistency among each set of samples indicates that random contamination was
38 not responsible for the systematic difference. Although the SAFe tank samples were
39 processed through filters with a smaller effective pore diameter (0.22 μm) than the Niskin
40 samples (0.45 μm), tests at the BATS site indicate that differences this large would not be
41 expected for the different filtration procedures (Figure 4). Niskin-profile and SAFe-tank
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3 samples were collected within a few days of one another at the same location. Therefore,
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5 neither scavenging nor displacement of deep water masses can be invoked to explain the
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7 divergent results. The cause of this difference remains unresolved. Future tests of filter
8
9 pore diameter should be conducted to establish the abundance of colloidal ($<0.45 \mu\text{m}$) Th
10
11 and the sensitivity of dissolved ^{230}Th concentrations to filter pore size.
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15 Good agreement between results from samples collected in 1994 and in February
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17 2002 at the HOT site provides reason for optimism that different labs, using different
18
19 methods, can obtain consistent results over a period of more than a decade. However,
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21 results from the two occupations of the HOT site in 2002 indicate that concentration of
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23 ^{230}Th in the deep ocean may vary on subannual time scales by nearly 20%. Results from
24
25 the SAFe site also indicate that factors still to be identified may have a substantial (20-
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27 25%) effect on reported ^{230}Th concentrations. Additional tests will be required to
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29 determine whether these factors involve unknown analytical artifacts. Whether or not
30
31 such artifacts exist, the results from the HOT site offer reason to suspect that nature may
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33 not be as stable as thought, that is, that concentrations of nuclides in the deep ocean
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35 cannot be assumed to remain constant over multi-decadal time scales in some locations.
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37 If so, then another strategy for sustained intercalibration must be sought. Further
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39 discussion among the community making these measurements will determine whether
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41 archived aliquots of homogenized seawater samples, such as those collected from the
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43 SAFe tank (see above), or artificial seawater standards (Figure 13), are best suited for this
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45 purpose.
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53 5.13 Prior intercalibration efforts: Prior to the GEOTRACES effort, the author
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55 of this manuscript from the Alfred Wegener Institute (Bremerhaven, Germany) initiated
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3 an intercalibration of labs that measure Th and Pa in seawater. Two sets of samples were
4 distributed. The first set was collected aboard the *R/V Polarstern* from 2000 m depth in
5 the Angola Basin on 9 November 2005 at station PS69/021 (11° 51.80' S, 2° 30.72' W).
6 Unfiltered water, collected with a standard rosette, was acidified with 1 ml/liter quartz-
7 distilled HCl. A second set of samples was collected from a depth of 2000 m on 2
8 September 2007 aboard the *R/V Polarstern* during cruise ARK XXII/2 in the Makarov
9 Basin (Arctic Ocean). Two subsets of Arctic samples were collected, one using the
10 standard ship's rosette and one using the trace metal-clean sampling system of the
11 Netherlands Institute of Sea Research (De Baar et al. 2008). Unfiltered Arctic samples
12 were acidified with 1 ml/liter Seastar™ concentrated HNO₃.
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27 Results from both sets of samples are reported together in Figure 18 because of
28 the relatively small number of data. Reporting the results together is not meant to imply
29 that concentrations of Th and Pa in the Angola Basin are expected to be the same as in
30 the Arctic Ocean. Only two labs (Labs W and Z) reported concentrations of ²³²Th
31 (Figure 18A). Values reported by Labs W and Z are in agreement for the Angola Basin
32 samples, indicating that there was no bias associated with spike calibration. Therefore, it
33 is unclear why the concentration of ²³²Th reported by Lab Z for the Arctic Ocean is
34 significantly less than that reported by Lab W. In light of the widespread but variable
35 contamination problems encountered for ²³²Th during the GEOTRACES intercalibration
36 (Section 5.7), contamination may have been a factor. The average concentration of ²³⁰Th
37 reported for the Angola Basin is 19.7 fg/kg (Figure 18C). Only Lab W reported values
38 within their reported uncertainty of the overall average. Concentrations reported by other
39 labs differed from the mean by 10 to 15%. Concentrations of ²³¹Pa reported for Angola
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3 Basin samples range over a factor of four (Figure 18B), indicating a problem with the
4 determination of this nuclide. Concentrations reported by Labs W and Z agree to within
5 reported uncertainty while values from Labs X and Y are much lower than those reported
6 by Labs W and Z.
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12 As for the Angola Basin, Labs W and Z reported concentrations of ^{230}Th and ^{231}Pa
13 for the Arctic samples that either agree within the reported uncertainties, or differ by
14 slightly more than the reported uncertainties. Concentrations of ^{230}Th and ^{231}Pa reported
15 by Lab Y were lower than values reported by Labs W and Z. The difference was slightly
16 greater than the uncertainty reported by Lab Y for ^{230}Th (Figure 18C), but much greater
17 for ^{231}Pa (Figure 18B). Lab X did not report a ^{231}Pa result for the Arctic samples, and
18 reported a concentration of ^{230}Th below those of Labs W, Y and Z.
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29 These results indicate that problems existed among some labs measuring Th and
30 Pa in seawater before the GEOTRACES intercalibration was implemented. One
31 encouraging result from the pre-GEOTRACES intercalibration is that with one exception
32 (Lab X, ^{230}Th), there is no significant difference between results from the standard rosette
33 and those from the trace metal-clean system where samples from the two systems were
34 analyzed by the same lab. This is consistent with the conclusion from the GEOTRACES
35 intercalibration that seawater can be collected without contamination for Th and Pa using
36 standard rosettes (Section 5.3).
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51 **6.0 Discussion**

52 The GEOTRACES intercalibration brought together many members of the
53 international community of investigators who determine the concentrations of long-lived
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3 radionuclides in marine samples, many of whom had had no prior interaction. This
4
5 community-building effort will facilitate future collaboration in methods development as
6
7 well as in research on marine processes. Results from the analysis of marine sediments
8
9 (Figure 6) represent the largest effort to date to establish the internal consistency among
10
11 labs that determine ^{232}Th , ^{230}Th and ^{231}Pa concentrations in marine samples. Despite the
12
13 evidence for some heterogeneity of the sediment, with a few exceptions the results
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15 indicated good agreement among the participating labs.
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20 Results from the first GEOTRACES intercalibration cruise (Figures 7, 8, 11, 12)
21
22 indicated that some labs were unable to determine concentrations of dissolved and
23
24 particulate ^{232}Th , ^{230}Th and ^{231}Pa with sufficient accuracy and precision to undertake a
25
26 global survey of the distribution of these nuclides in the ocean. Poor reproducibility was
27
28 also evident in the results from a previous intercalibration among four of the labs that
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30 later participated in the GEOTRACES intercalibration (Figure 18).
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34 A substantial improvement in the performance of the participating labs was
35
36 evident in the results reported over the course of the GEOTRACES intercalibration
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38 initiative, illustrating the benefits of such intercalibrations to the marine geochemistry
39
40 community. By the time of the second intercalibration cruise, many labs had made
41
42 progress in reducing their ^{232}Th blanks, although others were still working on the
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44 problem. Results from the second cruise demonstrated that participating labs can
45
46 determine concentrations of dissolved ^{230}Th and ^{231}Pa in the deep sea that are internally
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48 consistent to within 7% of the mean value (Figure 9). Results from eight labs that
49
50 analyzed a solution containing unknown concentrations of ^{232}Th and ^{230}Th indicate that
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52 calibration of Th isotope yield monitors is consistent among the labs to within about 1%
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3 (Figure 13). Therefore, much of the variability observed in the results from seawater
4
5 analysis must be attributed to factors other than errors in the calibration of isotope yield
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7 monitors.
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10 Numerous sources of ^{232}Th contamination were encountered by participants,
11
12 including sample containers, reagents and labware. These sources of contamination were
13
14 identified and eliminated or reduced (see above as well as accompanying papers in this
15
16 volume by Wang et al., in preparation, Andersson and Schöberg this volume; Auro et al.
17
18 this volume; Roy-Barman this volume).
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22 Special systems designed to collect samples for the analysis of contamination-
23
24 prone trace metals are not necessary when sampling seawater to determine concentrations
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26 of dissolved ^{232}Th , ^{230}Th and ^{231}Pa (Section 5.3 and Figure 2). Standard Niskin bottles
27
28 can be used although precautions should be taken to minimize chances for contamination
29
30 during sampling.
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34 Concentrations of dissolved ^{232}Th , ^{230}Th and ^{231}Pa can be determined by filtering
35
36 with either Nuclepore or Supor membranes (Figures 4, 5). No significant differences in
37
38 dissolved concentrations of ^{232}Th , ^{230}Th or ^{231}Pa were detected in using filter membranes
39
40 with pore diameters ranging between 0.4 and 1.0 μm . A single test suggested that QMA
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42 (quartz) filters may be used as well, although this finding needs verification, especially
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44 for ^{231}Pa .
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48 Isotope yield monitors need not be added to samples at sea. There was no
49
50 detectable loss of dissolved ^{232}Th , ^{230}Th or ^{231}Pa during storage for up to three years from
51
52 seawater that had been filtered and acidified at sea (Figure 2). Isotope yield monitors
53
54 equilibrate rapidly with Th dissolved in seawater (too rapidly to establish a rate constant
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3 using the data presented here) whereas spike equilibration requires about a day for U
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6 (Figure 3).

7
8 Sampling at the BATS site on three occasions over a period of 16 years, by three
9
10 different labs, produced internally consistent concentration profiles of ^{232}Th and ^{230}Th
11
12 (Figure 16). Consistency among these results demonstrates the value of the BATS site as
13
14 a GEOTRACES baseline station. Other labs can use the relatively unvarying
15
16 concentrations of ^{232}Th and ^{230}Th at mid-depths for future intercalibration, with
17
18 reasonable confidence that the concentrations of ^{232}Th and ^{230}Th will be close to those
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20 reported in this study. Concentrations in surface and bottom waters may change,
21
22 however, and should not be used for intercalibration. Results from the Pacific (HOT site)
23
24 indicate that concentrations of dissolved ^{230}Th may vary by up to 20% at subannual time
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26 scales even at mid depths (Figure 17), a possibility that warrants further investigation.
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34 **7.0 Recommendations**

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36 Taking into account the lessons learned through the intercalibration initiative, we
37
38 now consider steps to bring the overall consistency among labs that determine Th and Pa
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40 concentrations in seawater from its current best-case level ($\pm \sim 7\%$, Figure 9) toward the
41
42 reproducibility demonstrated during the analysis of SW STD 2010-1 ($\pm \sim 1\%$; Figure 13).
43
44 First we note contributions to what we have identified as the three principal sources of
45
46 error (sampling, chemical purification procedures, and instrumental analyses), and then
47
48 we make specific recommendations for near-term testing and development, as well as
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50 general recommendations for procedures to be adopted during the remainder of the
51
52 GEOTRACES program.
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3 7.1 Sampling errors: Three sampling issues were identified during the
4
5 intercalibration:
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8 1) The sensitivity of operationally defined concentrations of dissolved Th and Pa
9
10 to filtration method remains incompletely resolved. Although there was no detectable
11
12 difference among concentrations of dissolved Th and Pa determined by filtration through
13
14 membranes of different composition and with pore diameters between 0.4 and 1.0 μm
15
16 (Figures 4 and 5), concentrations of Th were significantly lower in samples filtered
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18 through a membrane with a pore diameter of 0.22 μm (Figure 4). The extent to which Th
19
20 in seawater resides on particles $<0.4 \mu\text{m}$ in diameter should be tested more completely.
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24 2) The possibility of filtration artifacts was also identified. Cartridges used to
25
26 filter through a 0.22 μm pore diameter membrane were much larger (1.6 square meters
27
28 membrane surface area; > 1 liter dead volume) than the individual membrane filters (47
29
30 or 90-mm diameter) and Acropak capsules (500 cm^2 membrane surface area; 30 ml dead
31
32 volume) used to filter other samples. Whether the lower Th concentrations in samples
33
34 filtered through 0.22 μm pore diameter membranes is a filtration artifact, related to
35
36 sorption of dissolved Th to cartridge and membrane surfaces, or a measure of the true
37
38 amount of Th associated with particles with an effective diameter between 0.22 and 0.4
39
40 μm , remains unresolved.
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46 Pall AcropakTM 500 capsule filters, or an equivalent product, were recommended
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48 for use on GEOTRACES cruises (see the document “Sampling and Sample-handling
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50 Protocols for GEOTRACES Cruises” available via the GEOTRACES web site at
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52 <http://www.obs->
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54 [vlfr.fr/GEOTRACES/libraries/documents/Intercalibration/Cookbook.pdf](http://www.obs-vlfr.fr/GEOTRACES/libraries/documents/Intercalibration/Cookbook.pdf)) based on their
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3 flow rates (resistance to clogging) and the ability to clean them against trace metal
4
5 contamination. This recommendation was made before the filter tests shown in Figures 4
6
7 and 5 were completed, and the Acropak capsules have never been tested directly against
8
9 individual (47-mm or 90-mm) membrane filters or against filters with smaller effective
10
11 pore diameters for the determination of dissolved Th and Pa concentrations. These tests
12
13 need to be carried out together with further testing of large cartridges, such as the
14
15 Osmonics cartridge used to fill the SAFe tanks.
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20 3) Contamination of sample containers with Th was evident in a small number of
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22 cases during each intercalibration cruise (e.g., Figure 7 Lab 7; Figure 9 Lab 9). In some
23
24 cases the contaminated containers were cleaned by procedures that left no detectable
25
26 contamination in other containers of the same batch. Whether the observed
27
28 contamination occurred in the shore-based labs or during sample collection at sea is
29
30 unknown, but the experience emphasizes the need for precautions against contamination
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32 at all stages of sampling and sample handling.
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36 7.3 Chemical Purification: Both blanks and chemical yields are issues of
37
38 concern.
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41 1) High chemical yields are crucial because, as noted above, the determination of
42
43 dissolved and particulate Th and Pa concentrations in seawater in many cases is near the
44
45 detection limit for these nuclides. In some cases (not described in this manuscript),
46
47 participating labs reported low chemical yields. In some cases the cause of low yields
48
49 remains unknown, but in one case (Auro et al. this volume) it was traced to the
50
51 irreversible sorption of Th by material used in the construction of ion exchange columns.
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53 Investigators are encouraged to report the cause of low yields when they are identified.
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3 2) Reagent blanks can be large. As noted above, high purity iron carrier may
4 contain large amounts of Th. Ion exchange resin (e.g., Auro et al. this volume) can also
5 be a source of substantial Th contamination. Contamination can vary by a large amount
6 from one batch of reagents to another. Each batch of reagents should be tested carefully
7 and processed if possible (e.g., for iron carrier or for ion exchange resin) to reduce blanks
8 to tolerable levels.
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17 3) Isobaric interferences from molecular species were not discussed in this paper,
18 but they are a concern when measuring Th and Pa by mass spectrometry. Careful
19 purification of Th and Pa will help reduce isobaric interferences.
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24 7.3 Instrumentation: The three principal types of mass spectrometers used to
25 measure concentrations of Th and Pa were identified in the Introduction. Some issues are
26 unique to one type of instrument while other issues are common to all.
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32 1) Filament blanks, applicable to TIMS, are contributed from the rhenium
33 filament as well as from colloidal graphite applied to facilitate ionization of Th (e.g., see
34 above as well as Andersson and Schöberg this volume; Edwards et al. 1987; Roy-Barman
35 this volume). The procedure followed at the University of Minnesota for quality control
36 of filaments is described in Section 4.3 as an example that may be followed by other labs.
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43 2) Mass bias corrections can be large during analyses by ICP-MS. For example,
44 SRM 129 with a natural $^{238}\text{U}/^{235}\text{U}$ abundance of 137.88 is used at LDEO to determine
45 mass bias. Measured 238/235 ratios accompanying seawater analyses at LDEO varied
46 between 138 and 141. Mass bias must be measured frequently and incorporated into data
47 reduction routines. It is generally assumed that mass bias for U is not significantly
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3 different from that for Th and Pa, although most labs have not tested this assumption on
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5 their own instruments.
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8 3) Isobaric interferences, mentioned above, can be controlled to some extent
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10 during ICP-MS analysis by membrane desolvation and by adjusting instrument settings
11
12 (e.g., Choi et al. 2001). Investigators should take into account isobaric interferences as
13
14 well as sensitivity, abundance sensitivity, and other factors when optimizing each
15
16 instrument.
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19 4) Variable gain among multiple detectors adds to the uncertainty in measured
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21 atom ratios. This includes the variability in gain between an ion counter and a Faraday
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23 cup as well as variability between multiple ion counters.
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27 5) The linearity of detector response as a function of absolute signal intensity was
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29 generally assumed by participating labs not to be an issue, but this is a potential source of
30
31 uncertainty that should be considered in future studies.
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35 Many of the items listed above may have contributed to the observed variability
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37 of $^{232}\text{Th}/^{229}\text{Th}$ ratios during the analysis of SW STD 2010-2 (Figure 14). Each of these
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39 topics warrants further examination as the community measuring concentrations of Th
40
41 and Pa in seawater works toward improving internal consistency.
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44 7.4 General recommendations: The following recommendations expand upon
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46 the specific issues identified above. They were developed in the context of improving
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48 results from future GEOTRACES cruises although they apply to other sampling
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50 programs as well.
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53 1) Procedural blanks should be collected frequently at sea; for example, one
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55 seawater blank is collected at each station on U.S. GEOTRACES cruises by adding 2 to 3
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3 liters of Milli-Q (or equivalent) water to a sample container and processing it as a sample.
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5 Improved characterization of blanks will better constrain overall uncertainties, even if the
6
7 blanks themselves cannot be reduced significantly. High blanks should only affect
8
9 detection limits, not accuracy, if the blank is well characterized statistically.
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12 2) Concentrations of dissolved ^{230}Th and ^{231}Pa in ~10 liters of surface ocean water
13
14 are close to the detection limits for many labs. Analysis of replicate samples from
15
16 surface waters (the mixed layer) will improve the reliability of results.
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19 3) Concentrations of particulate ^{230}Th and ^{231}Pa in ~100 liters of seawater are
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21 below the limit of detection for many labs. It is recommended that the community work
22
23 together to either lower detection limits or increase sample sizes for particulate material.
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25 Alternatively, it may be necessary to designate a subset of labs with the best detection
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27 limits to analyze samples of particulate material for ^{230}Th and ^{231}Pa collected during
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29 GEOTRACES cruises.
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33 4) The distribution and analysis of the artificial seawater solutions (SW STD
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35 2010-1 and 2010-2) helped constrain the internal consistency among the labs reporting
36
37 results (Figure 13). It is recommended that these solutions, or an equivalent, be analyzed
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39 routinely when determining the concentrations of dissolved ^{230}Th , ^{232}Th and ^{231}Pa in
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41 seawater samples, much like the SAFe seawater standards that are analyzed routinely
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43 during the determination of trace metal concentrations in seawater ([http://www.obs-
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45 vlfr.fr/GEOTRACES/index.php/science/intercalibration/215-safe-standards-and-
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47 reference-materials](http://www.obs-vlfr.fr/GEOTRACES/index.php/science/intercalibration/215-safe-standards-and-reference-materials)).
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51 5) Results from the SW STD solutions (Figures 13 and 14) also illustrate the large
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53 differences in the magnitude of the errors reported among participating labs. This
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3 difference is due in part to the different instruments used to measure atom ratios, and the
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5 different instrumental procedures (e.g., the number of atom ratios averaged), but it also
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7 reflects the differences among participating labs in the protocols used to evaluate and
8
9 report errors. It was beyond the scope of this intercalibration to make a synthesis and
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11 evaluation of the different error reporting protocols, but a comparison of two of the
12
13 participating labs illustrates the nature of the difference. As noted above, investigators at
14
15 LDEO propagate one sigma errors in atom ratios measured by ICP-MS with uncertainties
16
17 in background count rate, tail corrections and spike calibration to report their analytical
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19 uncertainty. By contrast, investigators at WHOI propagate two standard errors (two
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21 sigma divided by the square root of the number of ratios) of the atom ratios determined
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23 by ICP-MS with uncertainties in background count rate and spike calibration to report
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25 analytical uncertainty (Auro et al. this volume). We recommend that a comparison be
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27 made of the methods used to report uncertainties and that the results of that synthesis be
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29 reported in a future publication.
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36 6) A standard protocol for GEOTRACES cruises is to collect replicate samples to
37
38 be analyzed by a lab other than the main lab(s) participating in the cruise. Although this
39
40 recommendation is directed primarily to labs that have not participated in the
41
42 GEOTRACES intercalibration exercise, it is good general advice under all circumstances.
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44 Having a second lab analyze replicate samples will identify inconsistencies and assure
45
46 overall data quality. The authors recommend that replicate samples be collected for
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48 analysis by a collaborating lab as a standard protocol for future work on dissolved ^{230}Th ,
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50 ^{232}Th and ^{231}Pa in seawater.
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Figure Captions

Figure 1. Results from the test of sample storage effects during transport from the ship to the shore-based lab (LDEO). All samples were collected at a depth of 2000 m at the BATS site using the U.S. GEOTRACES carousel equipped with GO-Flo bottles. Water was filtered, and then acidified and homogenized in the 1000-liter SAFe tank system (Section 2.3.1). Isotope yield monitors (^{233}Pa and ^{229}Th) were added to samples 1 - 9 at sea. Isotope yield monitors were added to samples 10 - 12 when they were processed at LDEO. Iron carrier added to samples 1 - 4 was precipitated and concentrated at sea. Iron carrier was added to samples 5 - 12 when they were processed, together, at LDEO. Error bars represent the propagation of one sigma errors based on the mean and standard deviation of five sequences of isotope ratios collected by ICP-MS, the estimated error in the ^{229}Th or ^{233}Pa spike concentrations, and the blank correction of the individual isotopes for each sample batch. Horizontal lines represent mean values for each set.

Figure 2. Concentrations of (A) ^{232}Th , (B) ^{231}Pa and (C) ^{230}Th determined after increasing duration of sample storage. All samples were collected at a depth of 2000 m at the BATS site. Solid symbols indicate samples that were collected using the U.S. GEOTRACES carousel equipped with GO-Flo bottles and homogenized in the 1000-liter SAFe tank. Open symbols indicate samples collected in Niskin bottles deployed on the ship's rosette. In (B) and (C) the black symbols have been corrected for ingrowth of ^{231}Pa and ^{230}Th , respectively, whereas the grey symbols have not been corrected. A high procedural blank during the analysis of the first Niskin sample precluded the determination of a meaningful ^{232}Th concentration. The solid horizontal line indicates the mean concentration of each nuclide (corrected for ingrowth in B and C).

Figure 3. Concentrations of (A) ^{230}Th , (B) ^{232}Th and (C) ^{238}U determined at the University of Oxford as a function of time after a suite of one-liter aliquots of a deep-sea water sample were spiked with isotope yield monitors. The time axis quantifies the interval between spike addition and co-precipitation.

Figure 4. Concentrations of (A) ^{232}Th , (B) ^{231}Pa and (C) ^{230}Th determined by filtering seawater collected in individual Niskin bottles that were all closed at a depth of 2000 m on the same cast at the BATS site. The type of filter used in each case is indicated across the bottom, along with filter diameter (mm) and nominal pore size (μm).

Figure 5. Concentrations of (A) ^{232}Th , (B) ^{231}Pa and (C) ^{230}Th determined by filtering seawater collected in replicate Niskin bottles from a depth of 635 m at a site on the continental slope off the east coast of the U.S. ($37^\circ 2.0' \text{ N}$, $74^\circ 23.87' \text{ W}$). The type of filter used in each case is indicated across the bottom along with filter diameter (mm) and nominal pore size (μm).

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3 Figure 6. Concentrations of (A) ^{232}Th , (B) ^{231}Pa and (C) ^{230}Th determined by analyzing
4 aliquots of homogenized siliceous ooze from the Southern Ocean. Participating
5 laboratories are identified anonymously by Lab Number across the bottom. The hollow
6 grey symbol for Lab 15 in (A) represents the result from reanalysis of the sample by ICP-
7 MS subsequent to the original analysis by alpha spectrometry.
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10 Figure 7. Concentrations of (A) ^{232}Th , (B) ^{231}Pa and (C) ^{230}Th determined by analyzing
11 aliquots of homogenized seawater collected from a depth of 2000 m at the BATS site
12 during the first intercalibration cruise. Samples were drawn from the 1000-liter SAFE
13 tank system that had been filled using repeated casts of the U.S. GEOTRACES carousel,
14 equipped with GO-Flo bottles. Water was filtered as it was transferred from the GO-Flo
15 bottles to the tanks and acidified to a pH ~ 1.7 . Hollow grey points for Lab 7 in (A) are
16 from 1-liter samples that were archived separately during the cruise and analyzed by Lab
17 7 subsequent to the discovery of Th contamination in their larger samples (Section 5.7).
18 Circled results in (C) indicate samples that were flagged by Lab 7 as being contaminated,
19 corresponding to their high ^{232}Th values (but not circled in A). Lab numbers are
20 unchanged from Figure 6.
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24 Figure 8. Concentrations of (A) ^{232}Th , (B) ^{231}Pa and (C) ^{230}Th determined by analyzing
25 aliquots of homogenized seawater collected from a depth of ~ 15 m at the BATS site.
26 Samples were drawn from the 1000-liter SAFE tank system that had been filled using
27 water pumped from a towed trace metal-clean fish (Section 2.3.1). Water was filtered as
28 it was transferred from the fish to the tanks (Section 2.3.1) and acidified to a pH ~ 1.7 .
29 Circled results in (A) and (C) indicate values that were flagged by Lab 7 at the time of
30 data submission as being contaminated. Grey points in (A) and (C) are from 1-liter
31 samples that were archived separately during the cruise and analyzed by Lab 7
32 subsequent to the discovery of Th contamination in their larger samples (Section 5.7).
33 Lab numbers are unchanged from Figure 6.
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37 Figure 9. Concentrations of (A) ^{232}Th , (B) ^{231}Pa and (C) ^{230}Th determined by analyzing
38 aliquots of homogenized seawater collected from a depth of 3000 m at the SAFE site.
39 Samples were drawn from the 1000-liter SAFE tank system that had been filled using
40 repeated casts of the U.S. GEOTRACES carousel, equipped with GO-Flo bottles. Water
41 was filtered as it was transferred from the GO-Flo bottles to the tanks and acidified to a
42 pH ~ 1.7 . Circled results in (A) indicate values that were flagged by Lab 9 at the time of
43 data submission as being contaminated. Grey symbols represent results from a second set
44 of samples that was distributed to Lab 8 after they reported that blanks had been
45 improved substantially. Lab numbers are unchanged from Figure 6.
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49 Figure 10. Concentrations of (A) ^{232}Th , (B) ^{231}Pa and (C) ^{230}Th determined by analyzing
50 aliquots of homogenized seawater collected from a depth of ~ 15 m at the SAFE site.
51 Samples were drawn from the 1000-liter SAFE tank system that had been filled using
52 water pumped from a towed trace metal-clean fish (Section 2.3.1). Water was filtered as
53 it was transferred from the fish to the tanks and acidified to a pH ~ 1.7 . Lab numbers are
54 unchanged from Figure 6.
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3 Figure 11. Concentrations of (A) ^{230}Th and (C) ^{231}Pa in particulate matter collected by in
4 situ filtration at a depth of 2000 m at the BATS site. (B) Particulate $^{232}\text{Th}/^{230}\text{Th}$ activity
5 ratio. Lab numbers are unchanged from Figure 6.
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8 Figure 12. Concentrations of (A) ^{230}Th and (C) ^{231}Pa in particulate matter collected by in
9 situ filtration at a depth of 80 m at a site over the continental slope off the east coast of
10 the U.S. ($37^\circ 2.0' \text{N}$, $74^\circ 23.87' \text{W}$). (B) Particulate $^{232}\text{Th}/^{230}\text{Th}$ activity ratio. Note the
11 position of “zero” on the concentration axis in (C). Lab numbers are unchanged from
12 Figure 6.
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15 Figure 13. Concentrations of (A) ^{230}Th and (B) ^{232}Th determined by analyzing aliquots of
16 SW STD 2010-1 (Section 5.9) as an unknown. Units have been changed here to reflect
17 the units used by participants. The following conversion factors are used at LDEO:
18 ^{232}Th : $2.43438 \times 10^{-7} \text{ dpm/pg}$; ^{230}Th 0.04558 dpm/pg . An insufficient number of results
19 for ^{231}Pa concentration have been reported to present them here. Labs are renamed here,
20 and the sequence of labs differs from that in previous figures.
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23 Figure 14. Atom ratios reported by participating labs for SW STD 2010-2, the solution
24 that had been spiked with ^{229}Th at LDEO (Section 5.9). Labs identifications (A - H) are
25 the same as in Figure 13. Lab I reported Th atom ratios but not Th concentrations (Figure
26 13).
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29 Figure 15. Concentrations of ^{10}Be determined by analyzing aliquots of homogenized
30 seawater collected from a depth of 2000 m at the BATS site. Samples were drawn from
31 the 1000-liter SAFE tank system that had been filled using repeated casts of the U.S.
32 GEOTRACES carousel, equipped with GO-Flo bottles. Water was filtered as it was
33 transferred from the GO-Flo bottles to the tanks and acidified to a pH ~ 1.7 .
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36 Figure 16. Baseline concentration profiles for ^{230}Th (A) and ^{232}Th (B) at the BATS site
37 (roughly 75 km southeast of Bermuda at $31^\circ 50' \text{N}$, $64^\circ 10' \text{W}$). Unfiltered samples
38 collected in 1993 were analyzed at the University of Minnesota (Section 4). Unfiltered
39 samples collected in 2009 were analyzed at the Woods Hole Oceanographic Institution
40 following methods described in Auro et al. (this volume). Duplicate samples of filtered
41 seawater were collected during KN193-6 in 2008 aboard the first GEOTRACES
42 intercalibration cruise and analyzed at LDEO (Section 3).
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45 Figure 17. (A) Concentration profiles for ^{230}Th collected at the Hawaii Ocean Time
46 Series (HOT) Aloha station ($22^\circ 45' \text{N}$, 158°W). Samples collected in 1994 were acidified
47 ($\sim 2 \text{ ml}$ doubly-distilled HCl per liter of seawater) without filtration (Roy-Barman et al.
48 1996). Samples collected in 2002 were processed as described by Choi et al. (2001). (B)
49 Results for ^{230}Th at the SAFE Station (40°N , 140°W), based on samples collected during
50 the second GEOTRACES intercalibration cruise, are compared with the September 1994
51 results from the HOT site. SAFE baseline profile samples were collected in Niskin
52 bottles and gravity filtered through Pall AcropakTM 500 filters, which contain a paired,
53 pleated filter arrangement containing a $0.8 \mu\text{m}$ pre-filter followed by a $0.45 \mu\text{m}$ filter.
54 Filtered samples were acidified with 4 ml/liter of ultraclean 6N HCl to bring the solution
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3 to a pH of ~ 1.7. SAFe tank results are from Figure 9. Triplicate results from SAFe tank
4 samples processed at LDEO are shown as individual points. Results from all other labs
5 (Figure 9) are presented as a grand average (SAFe Tank All) and one standard deviation
6 of the mean values reported by labs that reported results. The depth of the grand average
7 is offset from the LDEO results for clarity. Triplicate results from the baseline profile at
8 3042 m represent the full procedural analysis of three individual samples.
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11 Figure 18. Concentrations of (A) ^{232}Th , (B) ^{231}Pa and (C) ^{230}Th determined by analyzing
12 samples of seawater collected from the Angola Basin and from the Arctic Ocean (see
13 Section 5.13 for collection details). Unfiltered samples were acidified at sea. Samples
14 from the Angola Basin and those labeled “Arctic normal” were collected using a standard
15 rosette. Samples labeled “Arctic UC” were collected using the trace metal-clean
16 sampling system of the Netherlands Institute of Sea Research (De Baar et al. 2008). Lab
17 identifications in this figure differ from those in previous figures, but Labs W - Z here are
18 represented among Labs 1 - 12 in Figures 6 - 9.
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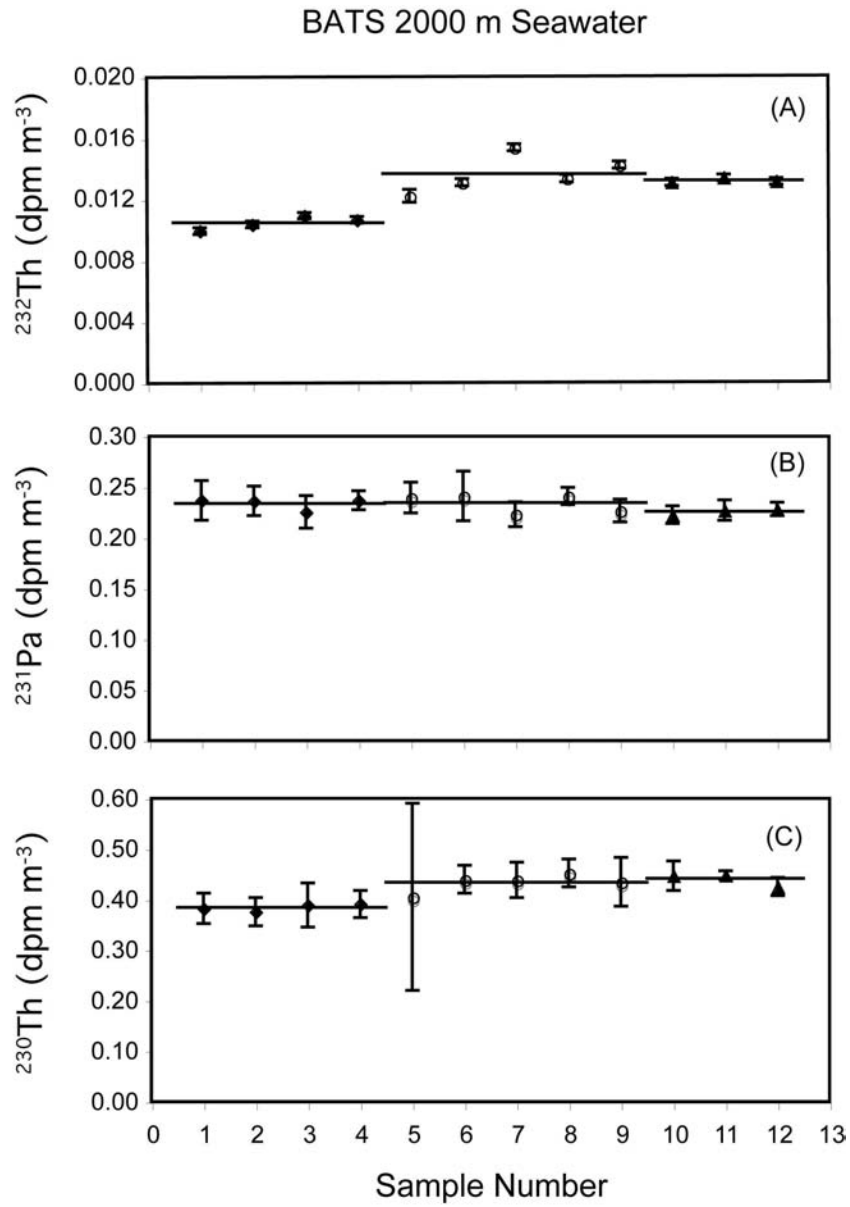


Figure 1
145x207mm (300 x 300 DPI)

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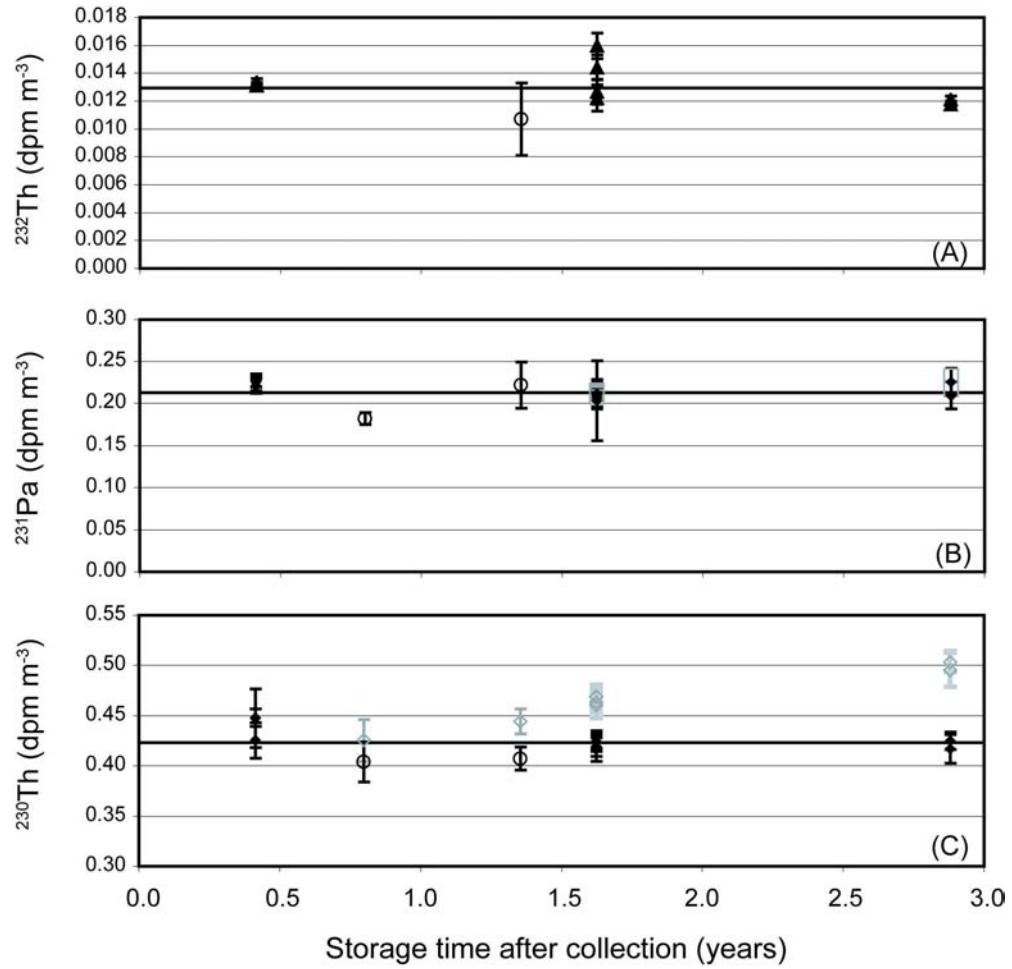


Figure 2
169x163mm (300 x 300 DPI)



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Spike Equilibration

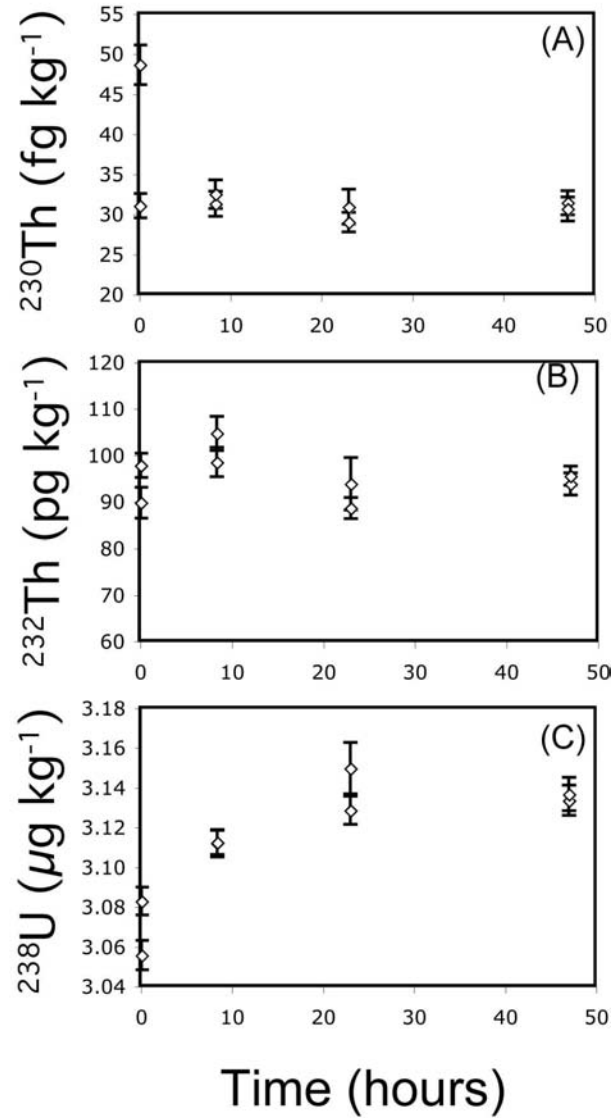


Figure 3
84x165mm (300 x 300 DPI)

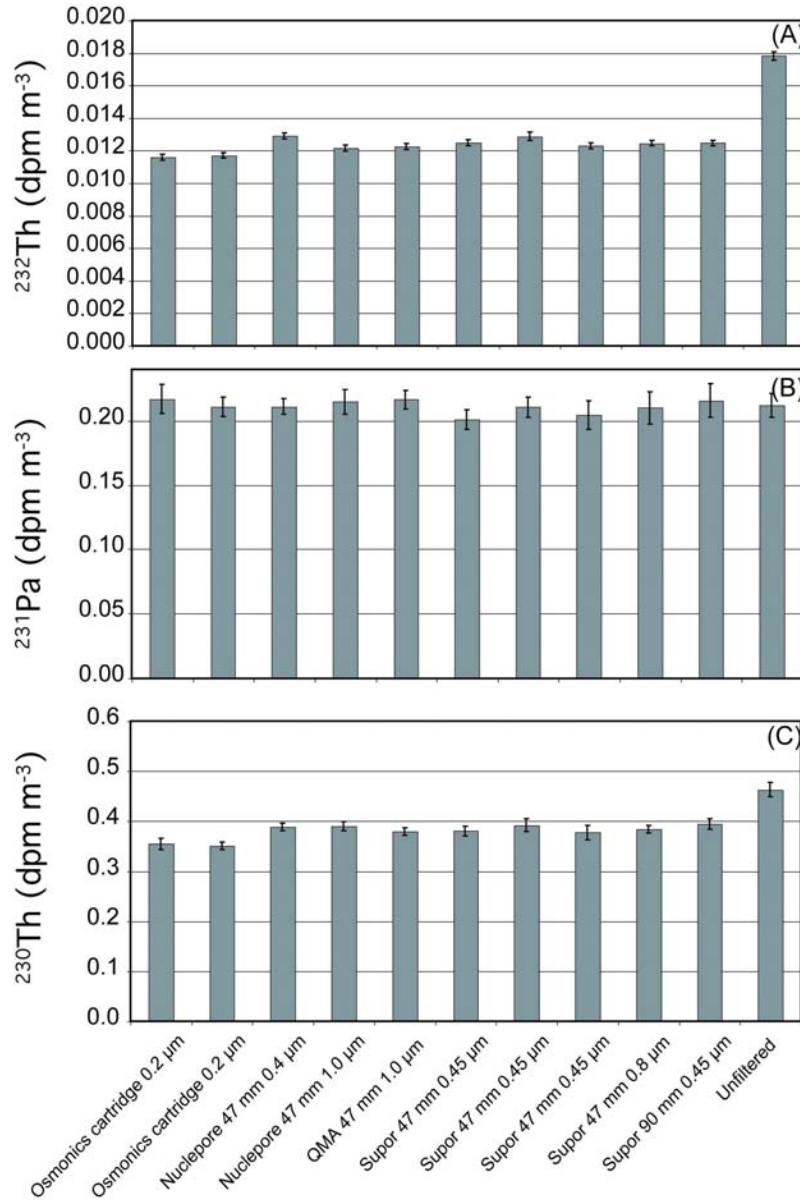


Figure 4
144x215mm (300 x 300 DPI)

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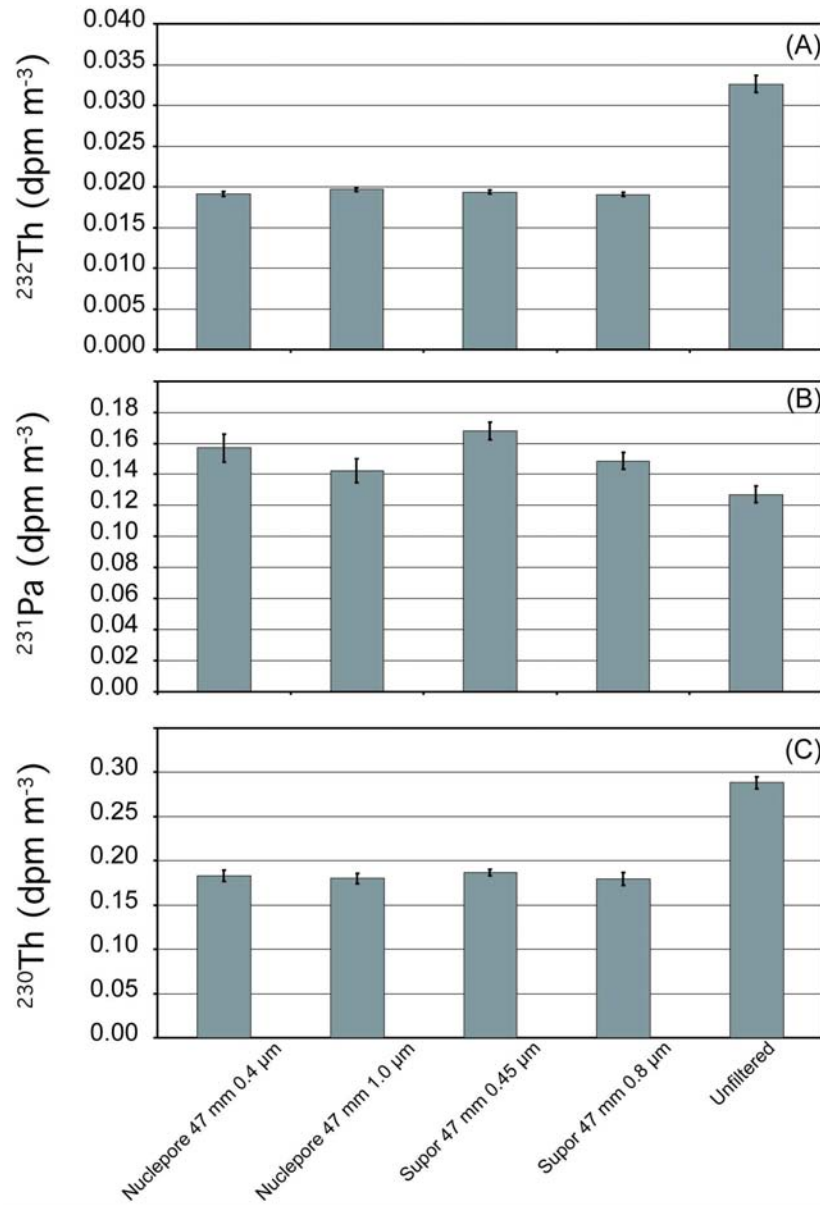


Figure 5
155x203mm (300 x 300 DPI)

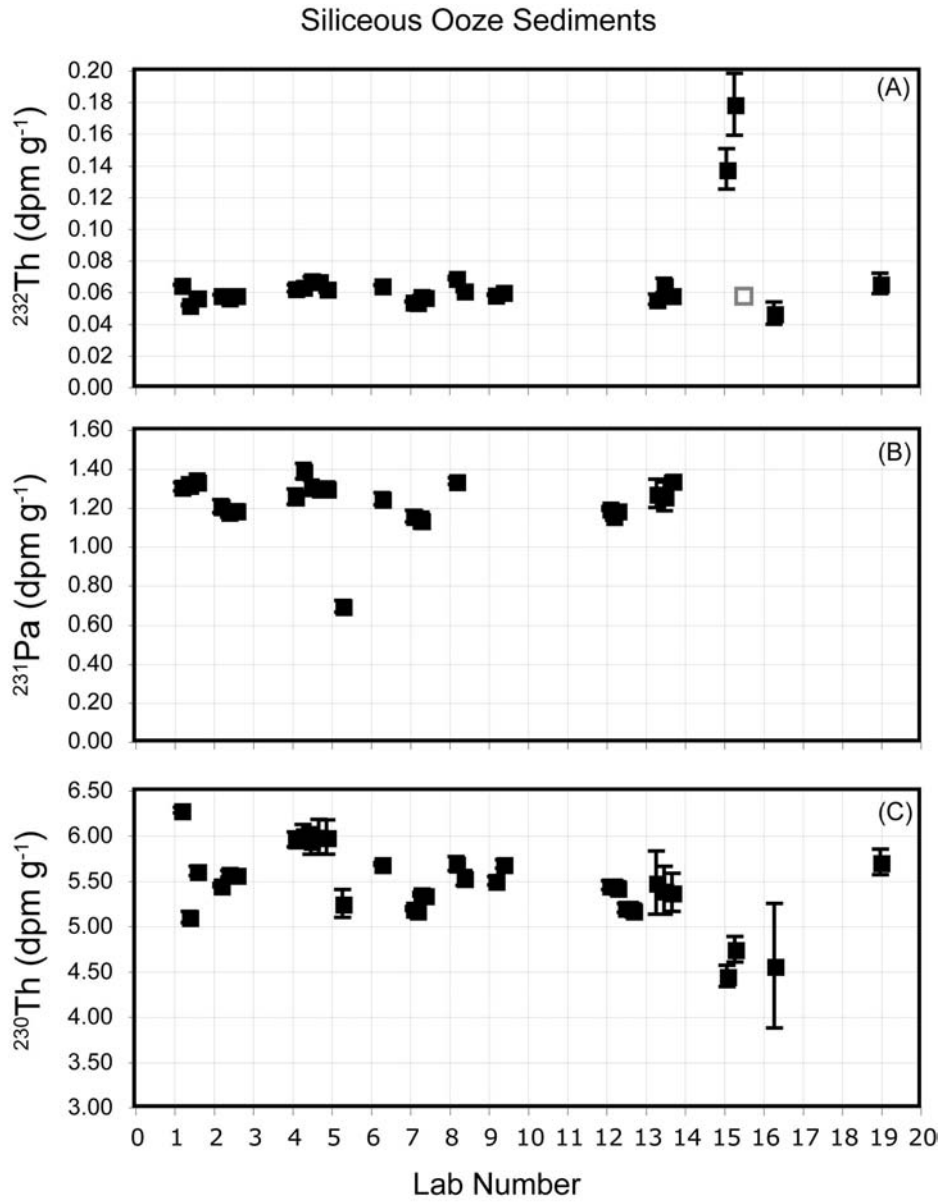


Figure 6
171x218mm (300 x 300 DPI)

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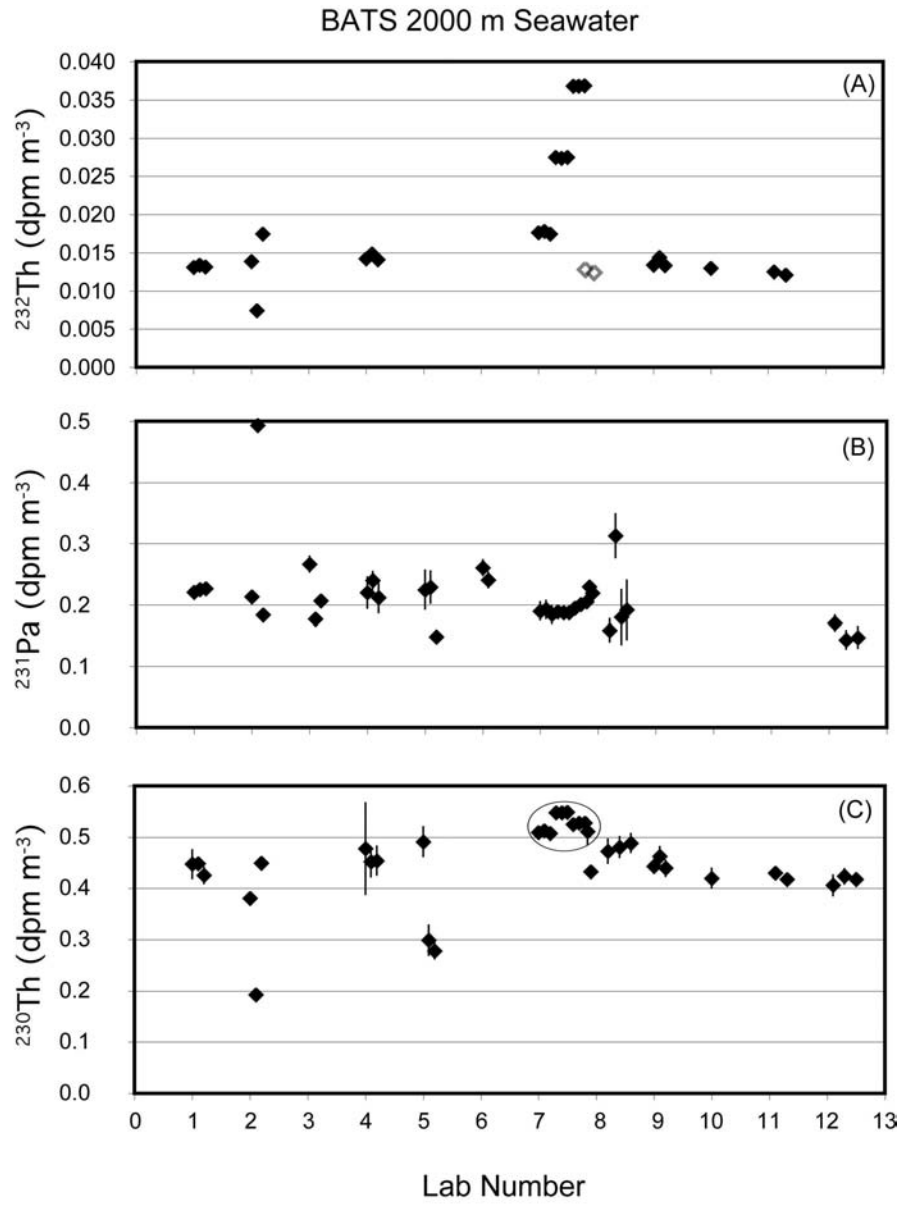


Figure 7
167x225mm (300 x 300 DPI)

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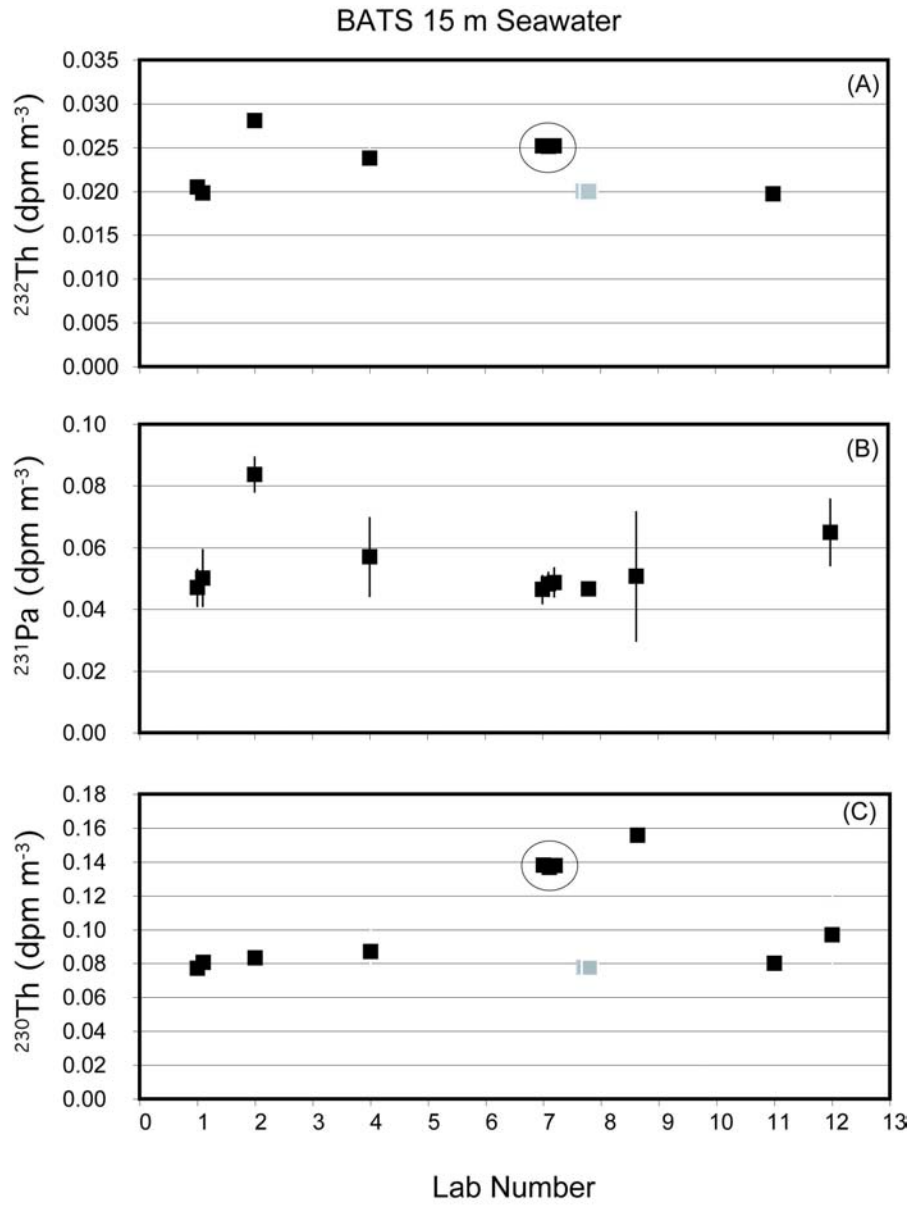


Figure 8
169x224mm (300 x 300 DPI)

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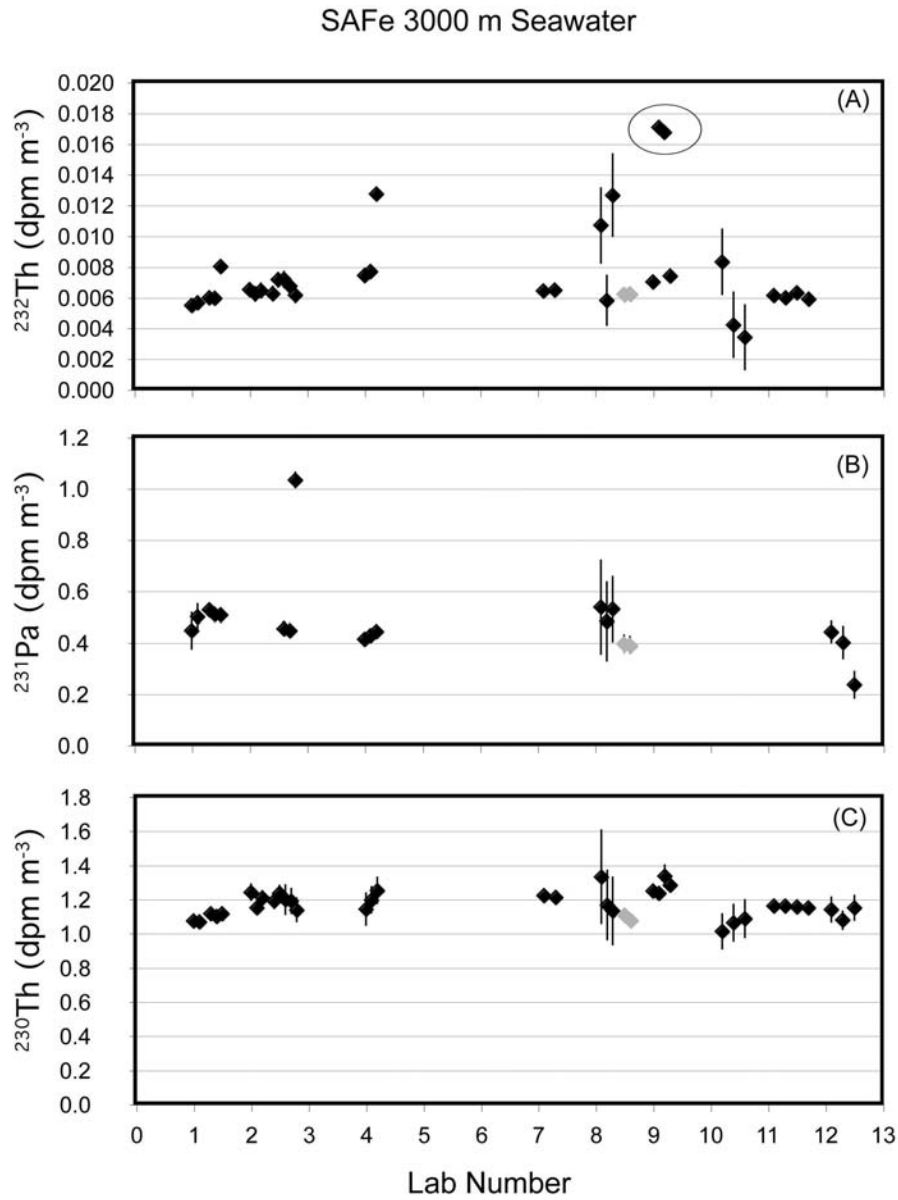


Figure 9
187x223mm (300 x 300 DPI)

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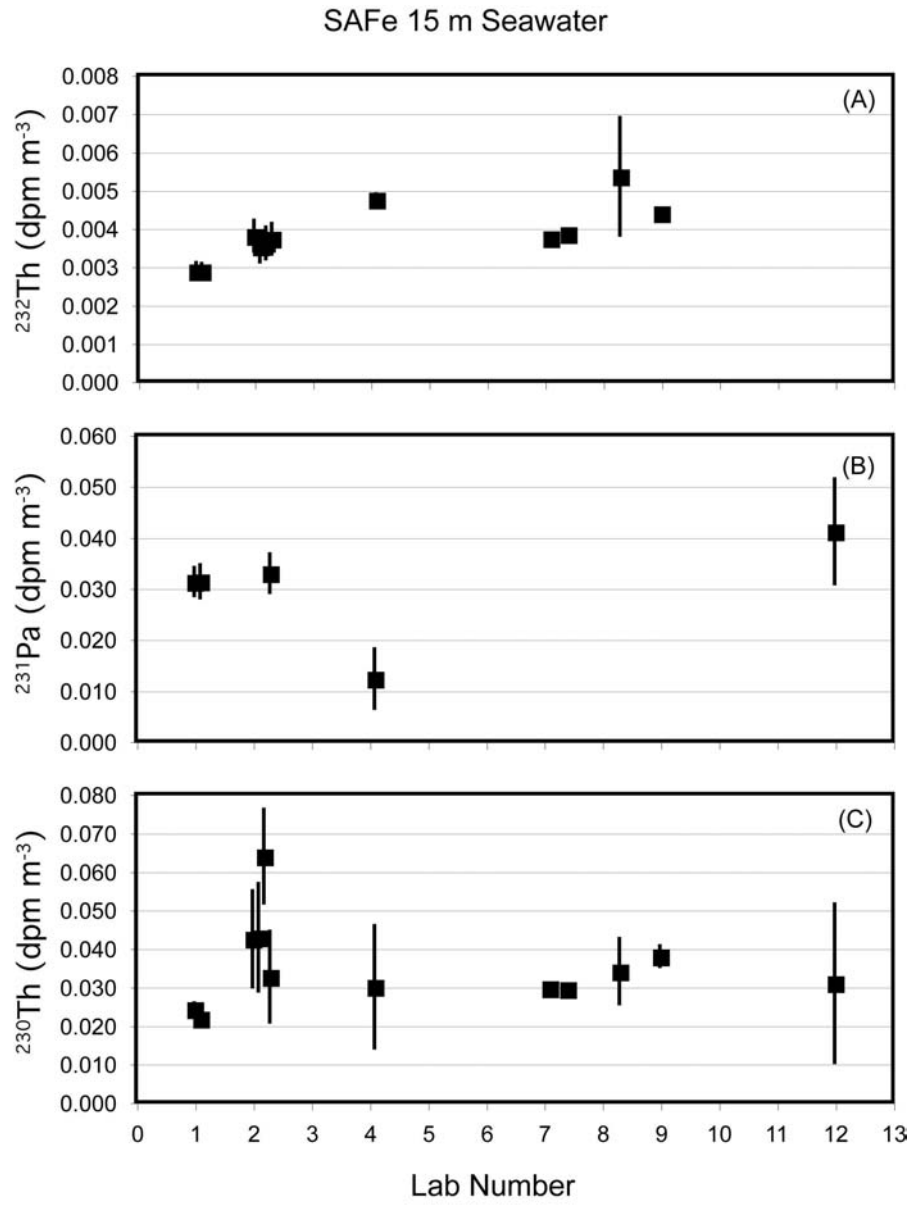


Figure 10
168x223mm (300 x 300 DPI)

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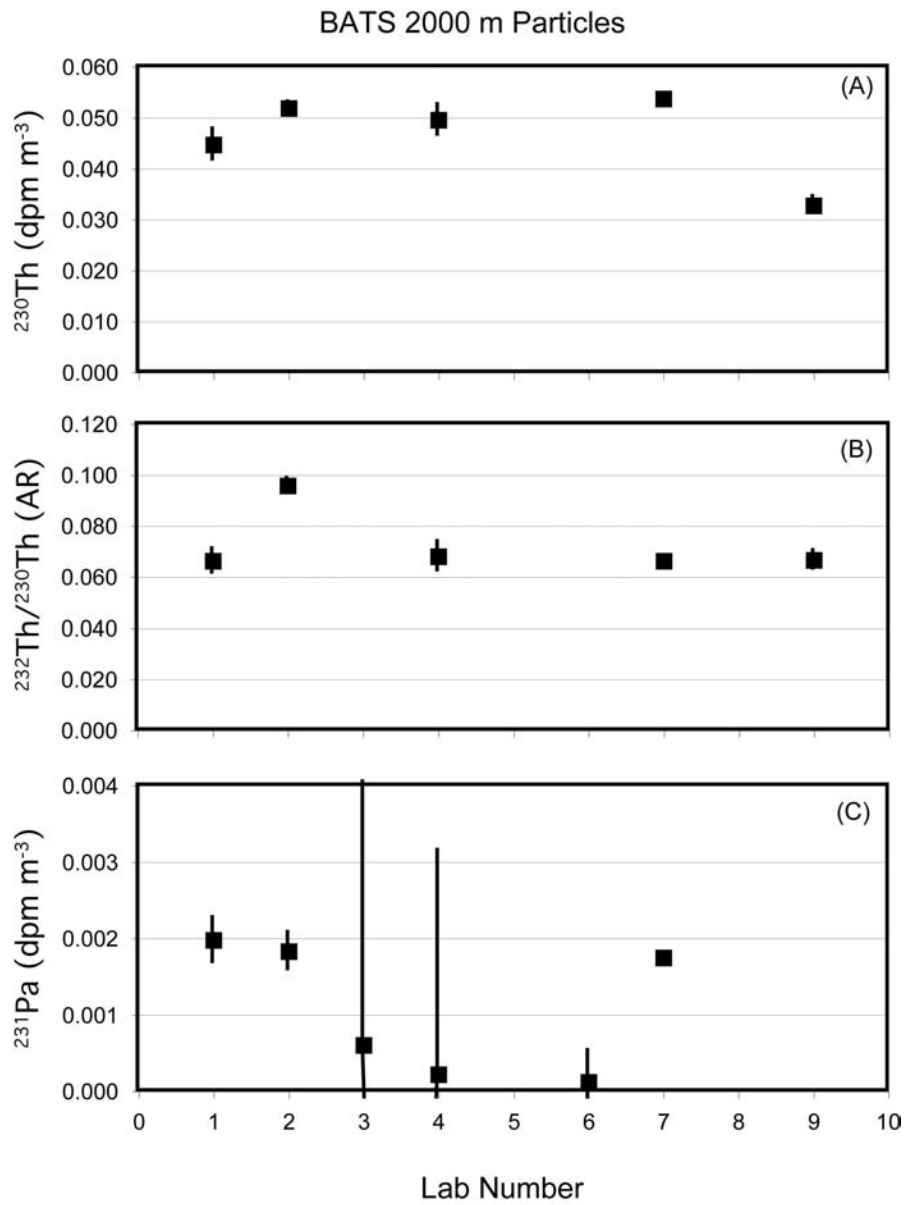


Figure 11
168x225mm (300 x 300 DPI)

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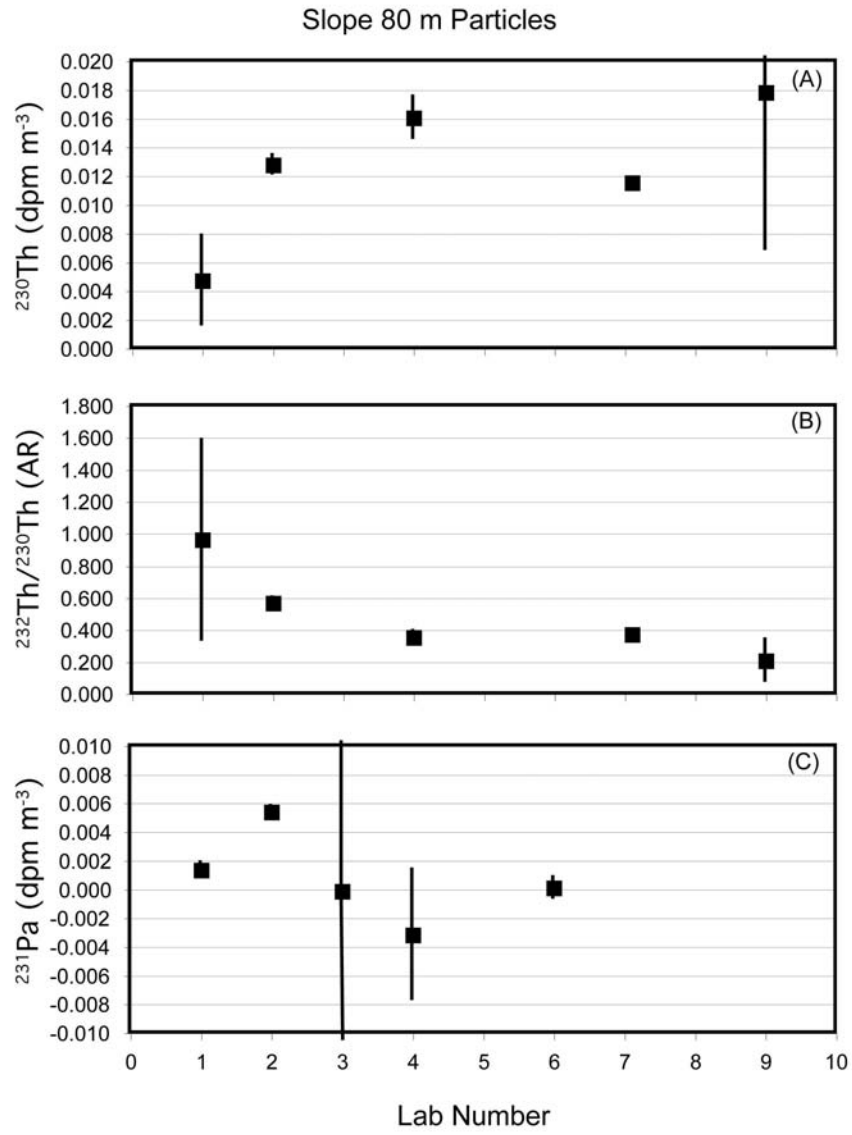


Figure 12
173x239mm (300 x 300 DPI)

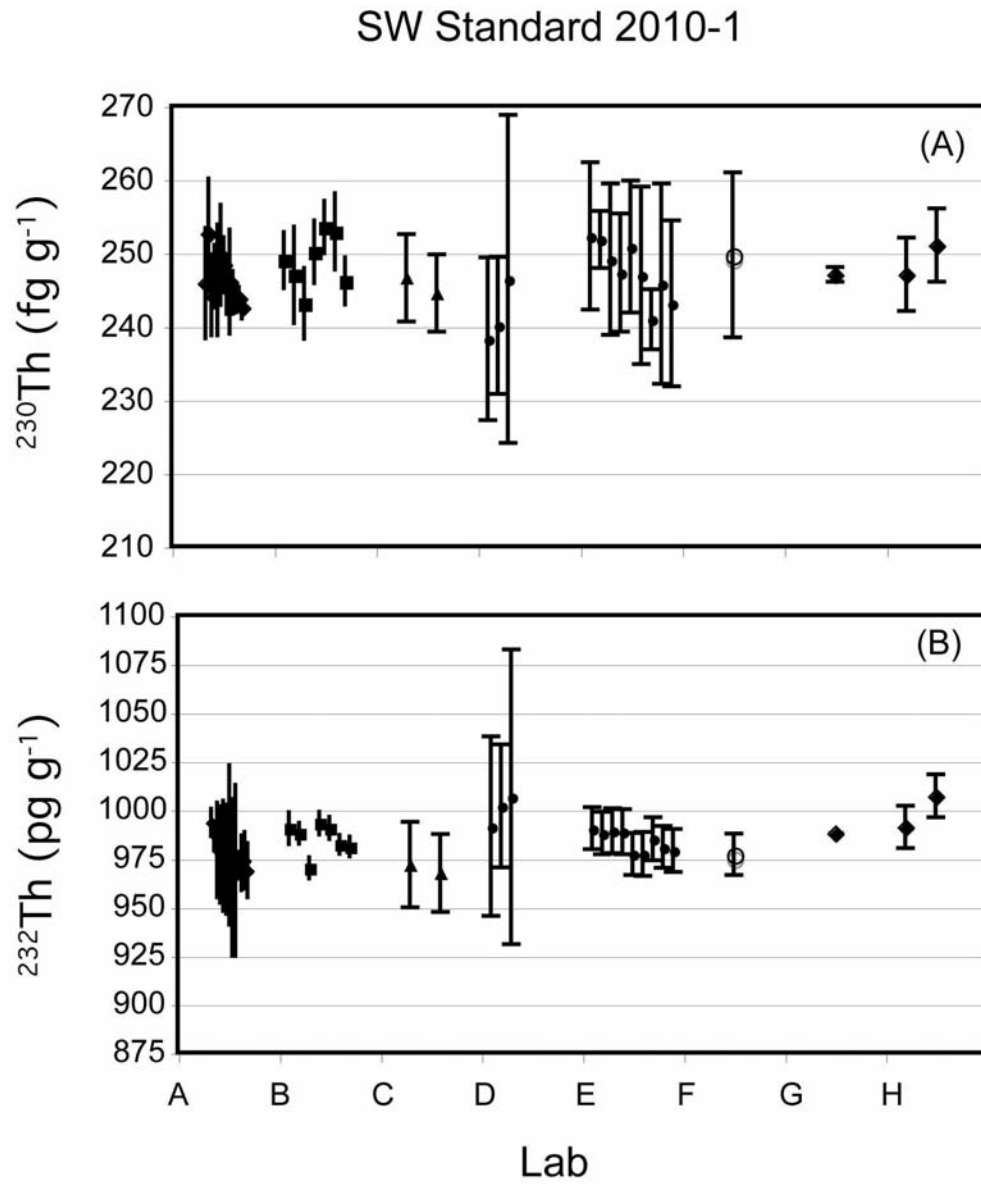


Figure 13
133x158mm (300 x 300 DPI)

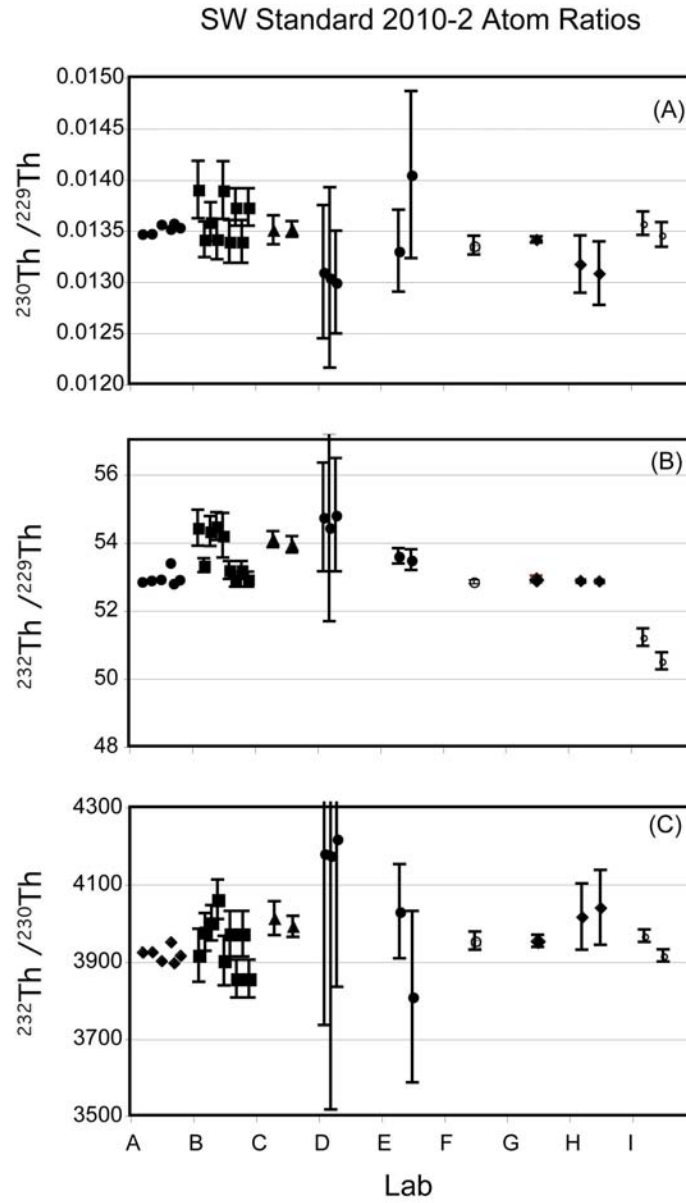


Figure 14
139x228mm (300 x 300 DPI)

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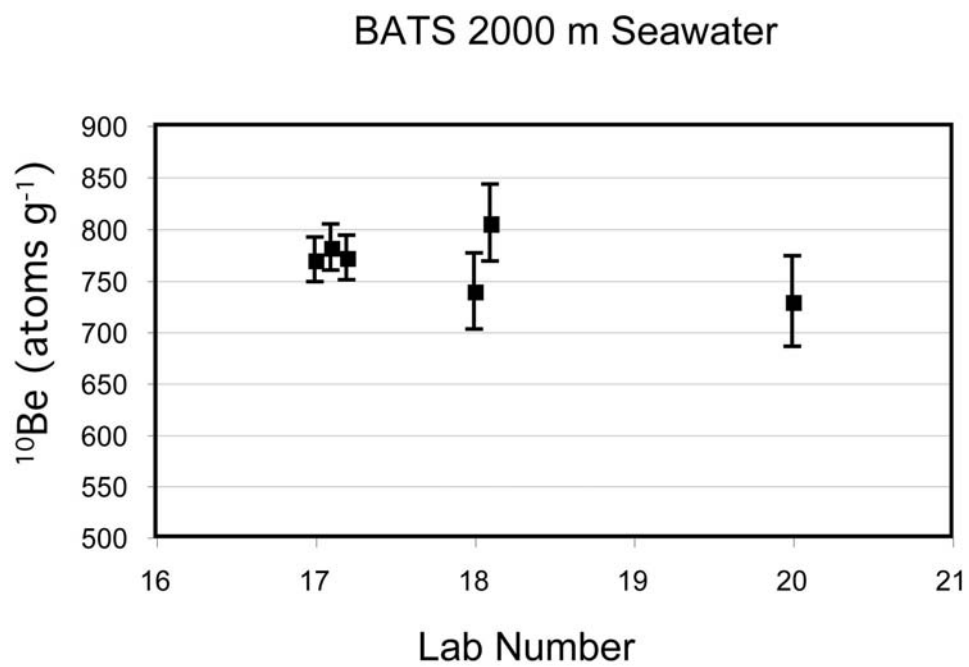


Figure 15
176x116mm (300 x 300 DPI)

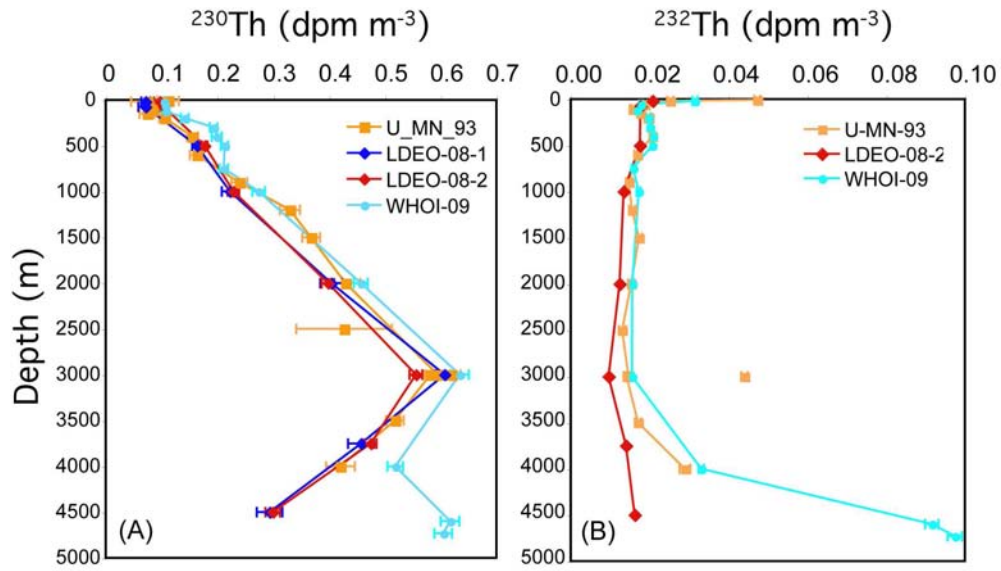


Figure 16
175x106mm (300 x 300 DPI)

View Only

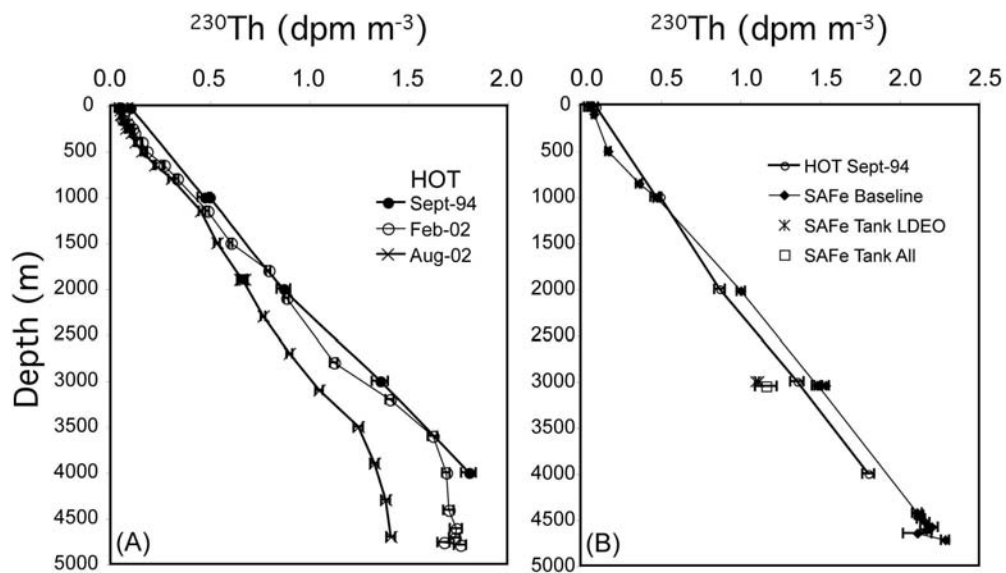


Figure 17
176x115mm (300 x 300 DPI)

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Angola (Atlantic) and Makarov (Arctic) Basin Seawater

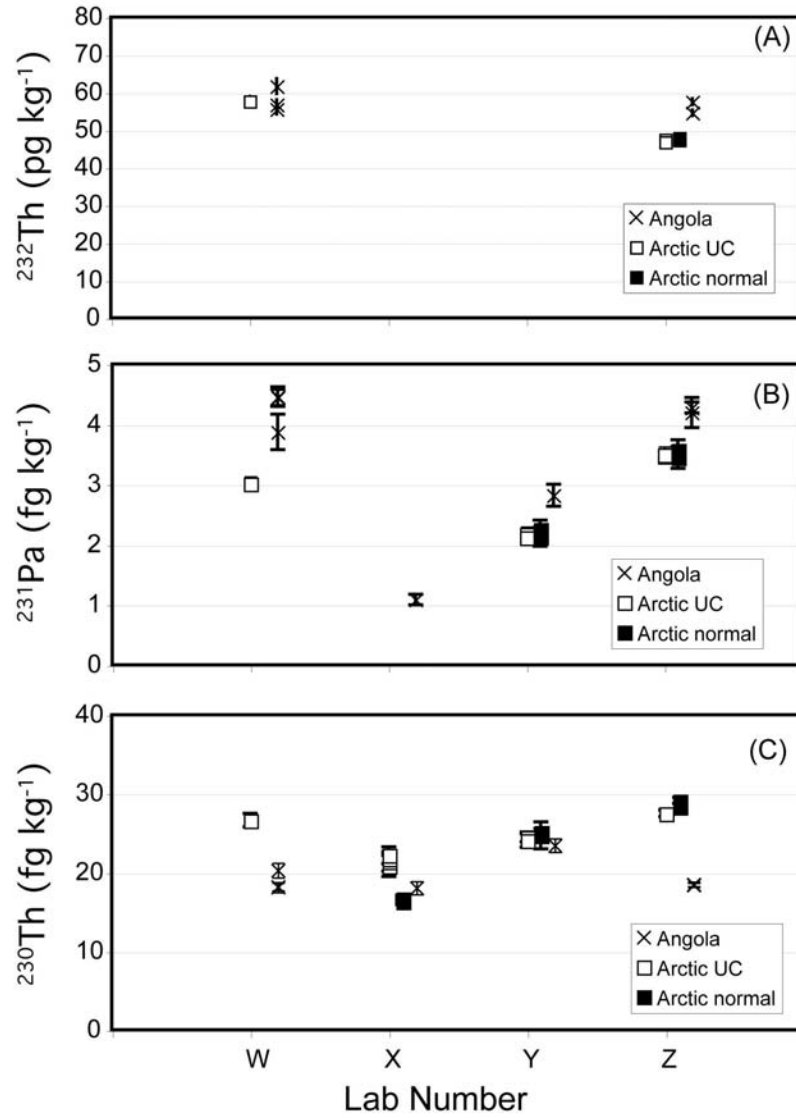


Figure 18
141x199mm (300 x 300 DPI)

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Table 1. Labs participating in the Th, Pa, Be intercalibration

Principal Investigator	Lab	Nation	Sediments Particles	Diss. Th, Pa	¹⁰ Be	Methods
Anderson, R.	LDEO	USA	X	X		1a, 2a, 3b
Andersson, P.	LIG/NRM	Sweden	X	X		1a, 2b, 2c, 3a
Chase, Z.	Oregon St. U	USA	X			2a, 3b
Edwards, RL/Moran SB	Minnesota/Rhode Island	USA	X	X		1a, 2a, 3c
Francois, R.	U British Columbia	Canada	X	X		1a, 2a, 3b
Geibert, W.	U Edinburgh	UK			X	1a, 2a, 2d, 3d
Frank, M.	IFM-GEOMAR	Germany			x	1a, 2a, 2e, 3d
Henderson, G.	Oxford U	UK	X	X		1a, 2a, 3c
Hong, G.	KORDI	Korea	X			2a, 3b
Jeandel, C.	LEGOS	France	X	X		1a, 2a, 3c
Marcantonio, F.	Texas A&M U	USA	X	x		1a, 2a, 3b
Nagai, H.	Nihon U	Japan			X	1a, 2a, 2d, 3d
Okubo, A.	U Tokyo	Japan	X	X		1a, 2a, 3b
Pichat, S.	ENS-Lyon	France	X			2a, 3c
Robinson, L.	WHOI	USA	X	X		1a, 2a, 3c
Roy-Barman, M.	LSCE	France		X		1a, 2a, 3a, 3c
Rutgers vd Loeff, M.	AWI	Germany	X	X		1a, 2a, 3b
Scholten, J.	IAEA	Monaco	X	X		1c, 2a, 3c
Yamada, Y.	NIRS	Japan	X			2a, 3b

Methods footnotes

1) Preconcentration: 1a Fe(OH)₃, 1b MnO₂, 1c Mg(OH)₂

2) Purification: 2a AG1X8, 2b UTEVA Eichrom, 2c TRU Eichrom, 2d AG50X8,

2e Coprecipitation in the presence of EDTA

3) Detection: 3a TIMS, 3b SC-ICP-MS, 3c MC-ICP-MS, 3d accelerator mass spectrometry

Table 1
177x121mm (300 x 300 DPI)