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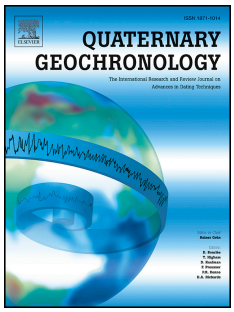

Testing the precision and accuracy of the U-Th chronometer for dating coral mortality events in the last 100 years

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Highlights

Testing the precision and accuracy of the U-Th chronometer for dating coral mortality events in the last 100 years

Tara R. Clark[1*], George Roff[2,3], Jian-xin Zhao[1*], Yue-xing Feng[1], Terence J. Done[3,4], John M. Pandolfi[2,3].

- A high-precision U/Th protocol to date recently-dead, sediment-contaminated coral
- Non-radiogenic ^{230}Th correction schemes trialled to improve ^{230}Th age accuracy
- Two-component mixing scheme successful in correcting non-radiogenic ^{230}Th
- A powerful tool developed for dating recent mortality events in coral communities

Testing the precision and accuracy of the U-Th chronometer for dating coral mortality events in the last 100 years

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ABSTRACT

To assist with our understanding of reef dynamics prior to modern monitoring programs and recent observations of coral decline, a robust dating technique is required to place coral mortality events and historical changes in community structure in an accurate chronological framework. In this study we adopted a refined Uranium-Thorium (U-Th) isotope measurement protocol using multi-collector inductively coupled plasma mass spectrometry (MC-ICP-MS) for rapid, precise and accurate age determination of a large branching *Acropora* coral death assemblage from an inshore reef of the Great Barrier Reef (GBR) where the timing of mortality is independently constrained. To achieve this, we developed a vigorous sample cleaning/treatment procedure to remove most non-carbonate detritus from the coral skeleton, and a correction scheme that accounts for initial ^{230}Th sources in the dead coral skeletons. Using this method, the ^{230}Th ages (with 2σ errors of 1 to 5 years) from 41 individual dead *Acropora* branches precisely bracket the timing of a documented ~100% loss of hard coral cover, primarily *Acropora*, that was caused by increased sea-surface temperatures during the 1997-1998 mass bleaching event. Our results demonstrate the applicability of U-Th dating in accurately determining the timing of previous disturbance events in coral reef communities, as well as identifying potential drivers. This approach provides a powerful tool to researchers and managers in assessing the current status of reefs and identifying areas vulnerable to degradation where long-term monitoring data are absent or too recent.

Keywords: U-Th dating, MC-ICP-MS, coral mortality, *Acropora*, Great Barrier Reef

1. Introduction

Current observations and future predictions of the status of corals reefs appear grim in the face of anthropogenic disturbance and climate change (Hughes et al. 2003; Pandolfi et al. 2011). However, on the Great Barrier Reef (GBR), Australia, there is still considerable debate as to whether inshore reefs are degraded or not (Hughes et al. 2011; Sweatman and Syms, 2011), partly due to a lack of understanding of coral community structure and disturbance history beyond the time period of long-term monitoring (Pandolfi et al. 2003; Roff et al. 2013). Palaeoecological studies provide a means to examine past changes in coral community structure and historical mortality events (Pandolfi and Greenstein, 1997), but a well-established chronology is also required to determine the absolute timing of these events. While a large number of studies have quantitatively described historical changes in coral communities (e.g. Greenstein et al. 1998; Pandolfi and Jackson 2006), only a few studies have isotopically dated samples at high enough resolution and with consistently low uncertainties to be able to link mortality events with specific drivers (e.g. Cramer et al. 2012; Pandolfi et al. 2006; Roff et al. 2013; Yu et al. 2012a, 2012b, 2006).

^{238}U-^{230}Th disequilibrium (U-Th) dating, which utilises the ^{238}U-^{234}U-^{230}Th decay chain, has proven to be a reliable method for determining the age of Pleistocene to Recent carbonate deposits. While the majority of studies have focused on dating samples thousands of years old, recent analytical advancements has led to dating coral samples as young as a few years with a precision of up to ± 1-2 years (Clark et al. 2012; McCulloch and Mortimer, 2008; Roff et al. 2013; Shen et al. 2008; Yu et al. 2012a, 20012b, 2006; Zhao et al. 2009). However, the uncertainty in hydrogenous and detrital sources of ^{230}Th [also termed 'non-radiogenic' ($^{230}Th_{nr}$), initial ($^{230}Th_0$) or secondary ^{230}Th sources that were not generated by the *in situ* decay of U] incorporated into the coral skeleton during skeletogenesis and after death, respectively, can result in inaccurate age estimates [for review see Zhao et al. (2009)]. This is especially true for young coral samples where the proportion

of $^{230}Th_0$ is significantly greater than the radiogenic ^{230}Th component, or for corals from inshore reef settings where non-carbonate terrestrial input is much greater compared to offshore settings; the former having ^{232}Th levels typically a few orders of magnitude higher than the latter (cf. Burley et al. 2012; Clark et al 2012; Roff et al. 2013; Yu et al. 2012a, 2012b). As the $^{230}Th_0$ in a sample cannot be separated from the radiogenic ^{230}Th during measurement, it can only be corrected for using the measured ^{232}Th level in conjunction with the initial $^{230}Th/^{232}Th$ ratio in the sample (expressed here as $^{230}Th/^{232}Th_0$). As $^{230}Th/^{232}Th_0$ may vary between sites or even between samples, a bulk-Earth activity value of 0.82 (atomic value ~4.4×10^{-6}) with a large arbitrarily assigned uncertainty of ± 50-100% has been commonly assumed to correct for the $^{230}Th_0$ contribution. Although this assumption was proven to be acceptable for most coral samples from inshore settings (Clark et al. 2012; Shen et al. 2008), the large associated uncertainty makes the age uncertainty of the corrected ^{230}Th age for young corals too large to be meaningful (see Zhao et al. 2009).

The $^{230}Th/^{232}Th_0$ ratio in a sample with detrital and carbonate components is generally constrained using isochron diagrams from multiple sub-samples of coeval material. This approach works if all the initial/detrital ^{232}Th and ^{230}Th are from the detritus, the $^{234}U/^{238}U$ and $^{230}Th/^{232}Th$ of the detrital component are the same for all sub-samples, and the system has remained closed (Bischoff and Fitzpatrick, 1991; Richards and Dorale, 2003). However, sources of $^{230}Th_0$ in corals can be highly variable (Clark et al. 2012; Cobb et al. 2003; Shen et al. 2008; Yu et al. 2006) and it is likely that corals from coastal environments contain two (or more) sources of $^{230}Th_0$: both detrital particulates and hydrogenous ^{230}Th. In that regard, inshore reef corals would be comparable to lake carbonates (Haase-Schramm et al. 2004; Lin et al. 1996) and deep-sea corals (Cheng et al. 2000a). Where the $^{230}Th/^{232}Th$ values of the detrital and hydrogenous components are dissimilar, the inability to account and correct for both sources can introduce substantial biases to the ^{230}Th age. To improve both the precision and accuracy of corrected ^{230}Th ages, it is necessary to be able to correct for $^{230}Th_0$ based on a well-constrained site- or sample-specific $^{230}Th/^{232}Th_0$ ratio (Clark et al. 2012; Shen et al. 2008).

For this study, the remains of a large number of dead branching *Acropora* corals were used to verify the accuracy of the U-Th dating method and the application of a sample-specific $^{230}Th_0$ correction scheme to determine the time of death. The *Acropora* 'death assemblage' (Pandolfi and Greenstein, 1997) sampled is at Pandora Reef, an inshore reef from the central GBR, Australia, for which both short-term observations (DeVantier et al. 1997) and long-term coral monitoring data are available (Done et al. 2007; Sweatman et al. 2005). These data allowed our estimates of the timing of coral death to be compared with those noted during direct field observations. A close match would facilitate the dating of the time of death of corals on the vast majority of reefs over long temporal scales (viz. pre-1980s) for which direct time-series observations of corals do not exist.

2. Materials and methods

2.1. Study site

Pandora Reef (18°48′S, 146°26′E; 750m long × 200m wide; Fig. 1) lies 17 km from the mainland and is episodically reached by flood plumes from the Burdekin and Herbert Rivers and other smaller tributaries adjacent to the region. Branching *Acropora* colonies were found in high abundance on the fore reef slope (sites P1 and t5) from 1985 to the start of 1998, and lesser abundance on the back reef slope (sites t1-4, V1 and V2) which was characterised by other coral genera including *Goniopora*, *Turbinaria*, and *Porites* (Done et al. 2007)

In 1997-1998 a severe El Niño event created a heat wave causing widespread bleaching along the entire length of the GBR from mid-December to early March (Berkelmans et al. 2004; Marshall and Baird, 2000). In the Palm Islands, bleaching was first reported on the 10 February 1998 and reduced coral cover by more than 50% in *Acropora*-dominated communities and up to 100% at some exposed reef flats (Gralton, 2002; Marshall and Baird, 2000). At Pandora Reef, bleaching was first reported at the beginning of March (Suzuki et al. 2003), affecting approximately 80% of hard corals down to a depth of 10 m. Members of the family Acroporidae were the most

affected, with almost a complete loss reported across the entire fore-reef [(Done et al. 2007; Sweatman et al. 2005); Fig. 5c]. At the time of sampling in May 2008, a large expanse of dead *Acropora* branches, attributable to the 1998 bleaching event, formed a consolidated matrix at the sediment-water interface along the entire south-west flank of Pandora Reef. This is in contrast to the high coral cover (>50%) reported in 1994, when branching *Acropora* colonies were present at all three [shallow (1-3 m), mid-slope (4-6 m) and deep (9-11 m)] depth ranges (De Vantier et al. 1997).

2.2. Coral collection and sampling

Dead *Acropora* coral rubble samples were collected from the leeward back reef environment at Pandora Reef at a depth of 4-5 m (Fig. 1). Five grab samples (~5 litres each) of dead coral rubble were collected by hand at the sediment-water interface and placed in calico bags at random points along each of four 20 m transects laid parallel to the reef flat (T1-T4; Table 1); thus totalling 20 grab samples with a spatial coverage of >80 m. The contents of the bags which predominately contained dead *Acropora* branches, were dried and individual branches with an intact branch tip or first-order branch (Bottjer, 1980) representing the most recent growth were selected from each of the 20 grab samples. Approximately 0.5-1 g of material was sampled from the cleanest part along the length of the branch, but within 16 cm (average sampling location was 5.0 cm) of each branch tip, using a diamond blade saw from 41 individual branches (Table 1). This ensured enough high quality material, free from alteration, for U-Th dating. Given high linear extension rates in branching *Acropora* corals typical of turbid, inshore, sheltered environments [from an average ~17.9 cm yr^{-1} (Crabbe and Smith, 2005) up to 33.3 ± 4.2 cm yr^{-1}(Diaz-Pulido et al. 2009)], an average sampling location ~5.0 cm from the branch tip would ensure that the site of skeletogenesis where sampling took place was within 0.3 yrs (or ~4 months) of the time of colony death.

Each sample was crushed to a ~1 mm grain size fraction and soaked overnight in a pre-cleaned glass beaker containing ~10% H_2O_2. It was then rinsed with Milli-Q water and centrifuged for 15 min at 4,000 rpm in a pre-cleaned Teflon beaker containing enough ~10% H_2O_2 to cover the

sample. Samples were again rinsed with Milli-Q water and ultra-sonicated several times until the water was clear. The excess water was then decanted and the sample dried on a hotplate at 40°C. This rigorous cleaning procedure ensures that detrital contaminants containing high concentrations of ^{232}Th are removed from the pore spaces of the skeletal matrix. The quality of each sample was then inspected under a binocular microscope and approximately 500 mg of the cleanest skeletal material selected for U-Th dating.

2.3. U-Th chemistry procedures

All U-Th chemistry and analytical procedures were performed under ultra-clean conditions at the Radiogenic Isotope Laboratory, the University of Queensland. Approximately 0.03 ml of a ^{229}Th-^{233}U mixed tracer (^{229}Th-^{233}U-spike #2) was added to each pre-cleaned Teflon beaker using a pipette and the weight recorded. The spike solution was then dried down completely at 60°C on a hot-plate, after which ~0.5 g of sample material was added to the spiked beaker. The sample/spike was then dissolved in double-distilled 70% HNO_3, co-precipitated with $Fe(OH)_2$ and U and Th separated using the standard ion-exchange column chemistry procedure slightly modified after Edwards et al. (1987). Following separation, the U and Th solutions were dried overnight at 80°C then redissolved using 1ml of 2% HNO_3. The amount of uranium solution to be measured was calculated based on pre-screened signals from a more dilute U-Th solution to ensure that the ^{238}U signal in the final solution did not exceed the capacity of the Faraday cups. For 500 mg of coral sample, approximately 20 µl U solution was taken from the 1 ml stock solution and transferred to a pre-cleaned 3 ml polypropylene tube to achieve a final concentration of ~7 ppb U. The entire 1 ml solution of Th was also added and made up to 3 ml using 2% HNO_3. Tubes were then centrifuged at 4,000 rpm for 20 min to remove any suspended material (mainly a trace amount of leaked resin) from the solution. All 41 samples were subsequently measured for U and Th isotopes using a Nu Plasma Multi-Collector Inductively Coupled Plasma Mass Spectrometer (MC-ICP-MS) to ascertain the timing of mortality, following a protocol first reported by Zhou et al. (2011).

2.4. MC-ICP-MS measurement

The U-Th mixed solution was injected into the MC-ICP-MS through a DSN-100 desolvation nebulizer system with an uptake rate of around 0.12 ml per minute. The U-Th isotopic ratio measurement protocol, which was first reported in Zhou et al. (2011), is similar to that previously described by Hellstrom (2003), but with minor modification to the detector configuration, viz. the U-Th isotopes were measured in two, instead of three sequences as used in Hellstrom (2003) (Table S1).

In our protocol, the two secondary electron multiplier (or SEM) gains are calculated from the SEM and Faraday signal ratios of ^{233}U and ^{229}Th (shaded grey in Table S1) that are collected in the two sequences, respectively. In addition, a deceleration lens behind SEM2 was used to increase the abundance sensitivity by 10 times (to reduce ^{238}U tailing at mass 237 to <0.5 ppm), so that the ^{232}Th tailing effect at mass 230 is small enough for accurate correction (the tailing effect was corrected using the geometric mean of background measurements at masses 229.5 and 230.5), regardless of the size of the ^{232}Th/^{230}Th ratio in the sample. For instance, even for samples with a ^{232}Th/^{230}Th abundance ratio of ~100,000, the ^{232}Th tailing contribution to the ^{230}Th signal is typically less than 3%. Each sample took about 25 minutes to measure. Measurements of samples, standards, and carryover memories were performed fully automatically using a modified Cetac ASX-110 autosampler.

A ‘drift monitoring’ solution was made by adding ^{229}Th and ^{233}U spikes separately into a dilute solution of a uranium oxide impurity standard New Brunswick Laboratory-6 (NBL-6) from the USA. The ‘drift monitoring’ was repeatedly measured after every six unknown samples, and the results were used to correct for long-term drift in a number of parameters such as ion counter gain (gain values were interpolated from bracketing the ‘drift monitors’ if ^{229}Th and/or ^{233}U signals in the samples were too small), and minor bias in the ^{230}Th/^{238}U and ^{234}U/^{238}U ratios during each session (the bias was mainly caused by the imperfect signal peak shapes and alignments). The

working U concentration in the 'drift monitor' when this set of samples were analysed, was ~6 ppb, with ^{238}U, ^{233}U and ^{229}Th signal sizes typically around ~3 volt (V), ~8 mV and ~2 mV at typical machine sensitivities, respectively, and has been precisely calibrated against the secular equilibrium Harwell Uraninite, HU-1 (Stirling et al. 1995; Zhao et al. 2001; Hellstrom, 2003).

2.4.1 Spike measurements

A mixed ^{229}Th-^{233}U tracer designed for dating young coral samples was added to each of the Teflon beakers prior to digestion. The isotopic compositions of this mixed tracer were determined to be: $^{238}U/^{233}U = 4.90 \times 10^{-4} \pm 0.74\%$, $^{234}U/^{233}U = 2.37 \times 10^{-3} \pm 0.15\%$, $^{235}U/^{233}U = 1.25 \times 10^{-4} \pm 0.30\%$, $^{232}Th/^{229}Th = 1.22 \times 10^{-4} \pm 1.6\%$ and $^{230}Th/^{229}Th = 4.78 \times 10^{-5} \pm 0.5\%$. The spike concentrations are $^{233}U = 1.42029 \times 10^{-2}$ nm g^{-1}, $^{238}U = 6.96390 \times 10^{-6}$ nm g^{-1}, $^{229}Th = 1.25476 \times 10^{-3}$ nm g^{-1}, $^{232}Th = 1.53356 \times 10^{-7}$ nm g^{-1}.

2.4.2 Blank correction

For MC-ICP-MS measurements of samples with very young ages, one of the main contributors to age error is the carry-over memory between samples being measured. However, this was alleviated by flushing the system prior to a new sample being measured for 15 min with 5% Aqua regia followed by 2% HNO_3 to prevent any cross contamination or 'memory' effect. In the clean-up stage all isotopes were monitored and raw counts measured on their respective detectors to ensure no carry-over memories from previous samples. Long-term monitoring of carryover memories over 20 months shows that ^{230}Th memory is less than 0.1 counts per second (cps) most of the time, which is negligible for most samples. ^{230}Th signals in the samples range from 20 to 50 cps, about 200-500 larger than the carry-over memory. The memories for all other isotopes are also negligible.

The total procedural ^{230}Th blank was determined to be $1.18 \pm 0.24 \times 10^{-10}$ nmol or 0.27 ± 0.05 fg (N=10); contributing an average 0.09 yr to the ^{230}Th ages of the samples in this study. The

procedural blanks for ^{238}U and ^{232}Th were averaged at $1.4 \pm 0.9 \times 10^{-5}$ nmol (or 3.3 ± 2.2 pg) and $3.0 \pm 1.9 \times 10^{-6}$ nmol (or 0.69 ± 0.41 pg), respectively, which are negligible for coral samples containing typically ~3 ppm U. These values are much lower than the procedural blanks measured using thermal ionisation mass spectrometry (TIMS), where high blank contributions were considered to be a result of more complex column chemistry and the colloidal graphite used to load the sample onto the filaments (Clark et al. 2012). Procedural ^{230}Th and ^{232}Th blanks were extracted from the samples in the Microsoft Excel spreadsheet used for U-Th age calculation.

2.5. Initial ^{230}Th ($^{230}Th_0$) correction

After MC-ICP-MS measurements, U-Th ages were calculated using the Isoplot/Ex version 3.0 program (Ludwig, 2003b). In order to assess whether the presence of $^{230}Th_0$ can be reliably corrected for, four different corrections schemes using likely $^{230}Th/^{232}Th_0$ ratios were tested: a bulk Earth, isochron-derived (detrital), live coral (hydrogenous) and site-specific model $^{230}Th/^{232}Th$ value that accounts for both detrital and hydrogenous sources. The corrected ^{230}Th ages were then compared with the 'true', independently constrained age of the coral death assemblage to evaluate which scheme returns the most accurate ^{230}Th age.

2.5.1. Bulk Earth-based correction scheme

The ^{230}Th age data was corrected using the bulk-Earth activity value of 0.82 (atomic ratio ~4.4×10^{-6}) with an arbitrarily assigned uncertainty of ± 50-100% (Richards and Dorale, 2003).

2.5.2. Live coral (hydrogenous)-based correction scheme

The live coral (hydrogenous) correction was based on 12 $^{230}Th/^{232}Th_0$ ratios obtained from live *Porites* colonies collected from Pandora, Havannah and Fantome Island in the Palm Islands region, central GBR (Clark et al. 2012), which have a weighted mean activity ratio of 1.08 ± 0.08 corresponding to an atomic ratio of $5.85 \pm 0.52 \times 10^{-6}$. This value, with a more conservative

uncertainty of ± 20% (to encompass the full range of variation in the 12 $^{230}Th/^{232}Th_0$ ratios), was subsequently used to correct for $^{230}Th_0$ in the 41 *Acropora* samples in the present study.

2.5.3. Sediment (detrital)-based correction scheme

Pandora Reef is periodically reached by plumes from the Burdekin River and nearby streams (Done et al. 2007; McCulloch et al. 2003), and as a result, the detrital $^{230}Th/^{232}Th$ component in the *Acropora* samples is best represented by a mean Th/U ratio of 4.8 ± 0.9 based on 44 sediment samples from the Burdekin River catchment area measured in our laboratory (Cooper et al. 2006). Assuming secular equilibrium, this corresponds to an activity ratio of 0.65 ± 20% (atomic ratio of $3.53 \pm 0.71 \times 10^{-6}$). This is further supported by isochron-inferred $^{230}Th/^{232}Th$ ratios determined using local dead *Porites* corals. U-Th analyses of five *Porites* samples of coeval material (i.e. material that was formed at the same time but with different $^{232}Th/^{238}U$) defined $^{238}U/^{232}Th$ vs $^{230}Th/^{232}Th$ isochrons with intercepts on the $^{230}Th/^{232}Th$ axis giving a weighted mean initial $^{230}Th/^{232}Th$ activity ratio of 0.61 ± 0.02 (2σ), which is within error of the Burdekin sediment value (Fig. 3). This isochron-derived mean initial $^{230}Th/^{232}Th$ ratio, with a more conservative uncertainty of ± 20% as reflected by the Burdekin sediments, is considered to approximate the value of the detrital component and is therefore used in the age correction.

2.5.4. Two-component mixing correction scheme

A new $^{230}Th/^{232}Th_0$ correction ratio [$(^{230}Th/^{232}Th)_{mix}$] was developed to account for two major isotopically distinctive components (or end-members) contributing $^{230}Th_0$ to the ^{230}Th age of the coral sample: 1) an insoluble Th component adsorbed to terrestrially derived sediments or particulates that were incorporated into the skeleton either post-mortem (major) or during coral growth (minor), and 2) a soluble or hydrogenous Th component present in the water column that was incorporated into the skeleton during growth. As both insoluble (detrital) and soluble (hydrogenous) Th components could be incorporated into the skeletal matrix during growth, initial

Th in live coral skeletons itself is also a mixture of both components; therefore live corals should theoretically fall onto the binary mixing line between the two above-mentioned end-members. Because of this, the $(^{230}Th/^{232}Th)_{mix}$ ratio can be calculated for each sample using the following mixing equation:

$$\left(\frac{^{230}Th}{^{232}Th}\right)_{mix} = \left(\left(\frac{^{232}Th_{live}}{^{232}Th_{dead}}\right)\times\left(\frac{^{230}Th}{^{232}Th}\right)_{live}\right) + \left(\left(\frac{^{232}Th_{dead} - ^{232}Th_{live}}{^{232}Th_{dead}}\right)\times\left(\frac{^{230}Th}{^{232}Th}\right)_{sed}\right) \quad \text{Eqn. 1}$$

where $^{232}Th_{dead}$ is the measured ^{232}Th value (ppb) in the individual dead coral sample of interest. $^{232}Th_{live}$ is 0.95 ppb, being the mean ^{232}Th value of live *Porites* coral samples in and near the study area (N = 12) with a corresponding $^{230}Th/^{232}Th_{live}$ activity ratio of 1.08 ± 20% (atomic ratio of 5.85 × 10^{-6} ± 20%). This value is representative of the isotopic composition of the mixture of the detrital/hydrogenous components incorporated in the live coral skeleton during growth, and is isotopically closer to the soluble Th end-member. $^{230}Th/^{232}Th_{sed}$ activity ratio is representative of the terrestrially-derived insoluble Th component incorporated either post-mortem or as particulates during coral growth. This ratio is calculated to be 0.61 ± 20% (atomic ratio 3.53 × 10^{-6} ± 20%), based on y-intercept values of $^{230}Th/^{232}Th$ vs $^{238}U/^{232}Th$ isochrons defined by local dead *Porites* corals, with a conservative uncertainty of 20% to account for the variability in the region. The influence of detrital $^{234}U/^{238}U$ was considered totally negligible in our samples and was therefore not incorporated in our age calculations (See Supplementary Fig. S1).

3. Results and discussion

U-Th dating of inshore-reef coral mortality events that have occurred over relatively recent timescales (less than 200 yrs) is extremely challenging due to the presence of high and variable levels of $^{230}Th_0$ present both in the water column and adsorbed to fine sediments or particulates (Clark et al. 2012; Cobb et al. 2003; Robinson et al. 2004; Shen et al. 2008; Yu et al. 2006).

Physical separation of extraneous sources of Th from that produced by the *in situ* decay of ^{238}U is virtually impossible. Despite using a rigorous H_2O_2 cleaning method to help remove the bulk of sediments adhered to the coral skeleton (which is reflected by a reduction in ^{232}Th levels), ^{232}Th concentrations in the measured samples still averaged 1.5 ± 1.1 ppb, which is significantly higher than concentrations found in live coral skeletons (Fig. 4) and on average 50-100 times higher than branching corals from the central Pacific (e.g. Burley et al. 2012; Weisler et al. 2006). The presence of high levels of ^{232}Th is an indication of high $^{230}Th_0$ still present in the cleaned samples. For such young coral samples (< 200 years old), high levels of $^{230}Th_0$ can cause the measured ^{230}Th age to be highly inaccurate, if not meaningless. The effect high concentrations of detrital Th (as reflected by elevated ^{232}Th levels) can have on the U-Th data can be seen in Figure 5a. By comparing ^{232}Th concentrations obtained from each sample with their respective uncorrected ^{230}Th age, a negative correlation exists between increasing ^{232}Th and increasing uncorrected age of the sample (Fig. 5a). As expected, where ^{232}Th concentrations are high, the uncorrected ^{230}Th ages are much older than the 'true' ages of the samples, indicating that there is a significant contribution of $^{230}Th_0$ to their ^{230}Th ages. Using the skeletons of 41 *Acropora* corals obtained from the death assemblage whose time of death was independently constrained by long-term observations, we were able to assess the effectiveness of four different $^{230}Th_0$ correction schemes in accounting for the presence of $^{230}Th_0$ in the coral samples. How close the corrected ^{230}Th ages matched the timing of the 1998 bleaching event was the measure of success.

The conservative bulk-Earth $^{230}Th/^{232}Th$ activity ratio of 0.82 (atomic ratio 4.4×10^{-6}) (Richards and Dorale, 2003) is considered suitable for the correction of the $^{230}Th_0$ component in corals where the dominant source of detrital Th is terrestrially derived [e.g. Clark et al. (2012); and Shen et al. (2008)]. However, the large arbitrarily assigned uncertainty of 50-100% can result in excessively large age uncertainties, which, although acceptable for dating events/processes thousands of years ago, will render the corrected ages meaningless. The positive correlation

326 between the corrected ^{230}Th age and ^{232}Th concentration (Fig. 5b), suggests that the ^{230}Th age data
327 are overcorrected. The resulting poor age precision precludes the identification of a specific episode
328 of mortality (Fig. 6; Table 2).
329
330 Using live corals of known age (Clark et al. 2012; Cobb et al. 2003; Shen et al. 2008) and
331 ambient seawater thorium isotopic measurements (Shen et al. 2008), site-specific ^{230}Th/^{232}Th$_0$ ratios
332 can be constrained by comparing the ^{230}Th ages of the corals with their 'true' or absolute ages and
333 can be considered close to representing the hydrogenous ^{230}Th/^{232}Th, as initial Th in live corals is
334 mainly derived from (but not limited to) seawater during growth (Shen et al. 2008). Using this
335 method, 12 samples of known age obtained from *Porites* corals provided an alternative correction
336 for ^{230}Th$_0$ (Clark et al. 2012). Yet despite using the average live *Porites* value, the corrected ^{230}Th
337 ages still show a positive trend with ^{232}Th, suggesting that this correction scheme again over-
338 corrected the ^{230}Th$_0$ component in the dead coral skeletons (Fig. 5c; Table 2). The significantly
339 higher ^{232}Th concentrations in the dead *Acropora* coral samples [1.5 ± 1.1 ppb (N = 41)] compared
340 to their living counterparts [0.15 ± 0.18 ppb (N = 7)], suggests that much of the Th in these corals
341 was incorporated after death (Fig. 4). A live coral ^{230}Th/^{232}Th$_0$ ratio is thus not suitable for ^{230}Th$_0$
342 correction, as it does not take into consideration the contribution of an insoluble Th component with
343 a contrasting ^{230}Th/^{232}Th$_0$ ratio.
344
345 The isochron-derived ^{230}Th/^{232}Th$_0$ activity ratios [weighted mean 0.61 ± 0.02 (2σ)] obtained
346 from multiple individual *Porites* coral samples from Pandora Reef and adjacent islands, most likely
347 reflect ambient sediment values and the primary source of detrital ^{230}Th in our samples (Fig. 3).
348 These low ^{230}Th/^{232}Th ratios are within error of those calculated from 44 trapped sediment samples
349 from the Burdekin River catchment (0.65 ± 20%; Cooper et al. 2006) whose plumes periodically
350 reach Pandora Reef (Done et al. 2007). Moreover, when all the U and Th isotopic data are plotted in
351 ^{230}Th/^{232}Th-^{238}U/^{232}Th space (Fig. 7), all values fall on a straight line, with the intercept at the y-axis

corresponding to an $^{230}Th/^{232}Th_0$ activity ratio of 0.64 ± 0.04 which is analytically indistinguishable from the values constrained using the other two independent methods described above. However, the use of the isochron derived $^{230}Th/^{232}Th$ ratio of 0.61 ± 20% for correction yielded ^{230}Th ages that centre around a peak of ~1994 (Fig. 6d; Fig. S2b; Table 2; Table 3); which does not match the timing of the observed 1997/1998 mortality event. Although coral mortality did occur as a result of multiple flood plumes and elevated SSTs in 1994 at Pandora Reef, overall mortality at this time was reported to be less than 1% of the total cover for this genus (DeVantier et al. 1997). The 1994 disturbance event is therefore unlikely to have been responsible for the normally distributed age population produced by 41 coral fragments collected over a distance of more than 80 m. The failure of this correction scheme is due to the fact that the hydrogenous ^{230}Th component with higher $^{230}Th/^{232}Th$ was ignored. Similar results have also been shown for Lake Lahontan carbonates, where samples containing hydrogenous ^{230}Th and corrected using isochron derived $^{230}Th/^{232}Th_0$ ratios were also too old (Lin et al. 1996).

The most accurate correction scheme accounted for the two isotopically contrasting components contributing $^{230}Th_0$: 1) detrital ^{230}Th from terrestrially-derived insoluble Th components incorporated into the coral skeleton either during growth (minor) or post-mortem (major), and 2) hydrogenous ^{230}Th adsorbed on detritus or present in the water column and directly incorporated into the aragonite matrix (Haase-Schramm et al. 2004; Lin et al. 1996). For living reef corals, the primary source of $^{230}Th_0$ is mainly from the dissolved fraction in seawater (Shen et al. 2008), although the presence of a minor detrital component in seawater cannot be ruled out at inshore settings. Live corals may scavenge a small amount of the detrital component present in the seawater column in particulate/colloidal forms as it switches from a passive autotroph to active heterotroph. It is likely that the minor detrital component in live corals is variable among different species and in different environmental settings: higher in *Porites* compared to *Acropora* (Clark et al. 2012; this study), and higher in inshore compared to offshore settings (cf. Burley et al. 2012;

Roff et al. 2013; Yu et al. 2012a, 2012b). In this regard, the mean $^{230}Th/^{232}Th$ ratios obtained from both live *Porites* and *Acropora* should theoretically fall on a binary mixing line between the hydrogenous (seawater) and detrital (sediment) end-members. Post-mortem, corals can no longer actively exclude sediments which become adsorbed into the porous skeletal matrix of the coral (Lasker, 1980). As a result, both sources of Th_0 need to be corrected independently using the binary mixing model in order to achieve accurate U-Th ages.

In this study, live *Porites* and isochron-derived $^{230}Th/^{232}Th_0$ ratios were used to approximate the isotopically distinctive hydrogenous and detrital end-membersin the dead *Acropora* corals, respectively. When the mean $^{230}Th/^{232}Th$ ratios for the live *Porites*, live *Acropora* and isochron-derived detrital values from dead *Porites* are plotted in a $^{230}Th/^{232}Th$ versus $1/^{232}Th$ diagram, all four types of samples fall on a binary mixing line between two end-members: a hypothetical seawater (hydrogenous) component and a terrestrial (detrital) component (Fig. 4). Interestingly, the live *Acropora* coral $^{230}Th/^{232}Th$ ratios fall between those determined for the dissolved and particulate fraction of seawater analysed from continental shelf settings in the western Pacific and eastern Indian Ocean (Shen et al. 2008), suggesting that this value may be an accurate representation of seawater values (although this is yet to be confirmed). While values obtained from live *Acropora* corals would better reflect the hydrogenous end-member in the dead *Acropora* samples dated in this study, they were not used in the equation for two reasons: 1) concentrations of ^{232}Th in the live *Acropora* samples are extremely low (0.15 ± 0.18 ppb) and difficult to measure accurately; 2) it is also difficult to independently constrain the 'true' age of a sample from an *Acropora* branch in order to determine $^{230}Th/^{232}Th_0$ without cross-referencing with annual growth bands (from X-rays) or elemental cycles (using ICP-MS). The assumption that we are sampling from within one year of growth is based on a few observational studies of annual extension rates for *Acropora* colonies from the inshore GBR and other turbid reef environments (Crabbe and Smith 2005; Diaz-Pulido et al. 2009). Moreover, directly determining seawater $^{230}Th/^{232}Th$ ratios may also be an inaccurate estimate of the hydrogenous $^{230}Th/^{232}Th_0$ component in the dead coral skeleton as

seawater ^{232}Th concentrations are highly variable over spatial and short-term temporal scales. For example, dissolved ^{232}Th concentrations measured from coastal environments by Shen et al. (2008) were on average ~80 times higher than the bulk ^{232}Th concentrations (excluding organics) in seawater from the Bahamas (0.00828 versus 0.00010 ppb, respectively). Shen et al. (2008) also found higher concentrations of ^{232}Th at high tide compared to low tide. Thus an understanding of site-specific seawater ^{232}Th concentrations is needed. In addition, it is difficult to know the ^{232}Th concentrations of the local seawater at the time when the coral died.

When the measured ^{230}Th/^{232}Th data for the dead *Acropora* corals are corrected using the two-component mixing equation (Eqn. 1), the isotopic variations shift towards this mixing line, with most of them falling between the detrital end-member and the live *Porites* ^{230}Th/^{232}Th ratio. The ^{230}Th age population corrected using the two-component mixing scheme also becomes normally distributed, as can be seen when the data are plotted as a relative probability plot [that incorporates both the mean and 2-sigma uncertainties of the individual dates using the Isoplot Program (Ludwig, 2003)]. The peak value of this distribution is ~1998 AD (weighted mean 1998.2 ± 0.3 AD, MSWD = 1.1) (Fig. 6d and Table 3), which is within error of the timing of the 1998 mass bleaching event (Fig. 6; Fig. S2a; Table 2). At this time, long-term monitoring datasets from sites similar in community composition less than 500 m away (Fig. 1) and regional scale observations documented almost 100% mortality in *Acropora* (Done et al. 2007; Maynard et al. 2008; Sweatman et al. 2005). In contrast, the probability distribution patterns of corrected ^{230}Th ages derived from all the other three correction schemes are all skewed to some extent, suggesting those schemes are insufficient to correct for $^{230}Th_0$ components in the samples.

Had the death assemblage been derived from a number of events (e.g. cyclones, predators or freshwater inundation during previous decades), we would not have expected such a tightly constrained age estimate (Edinger et al. 2001). For example, if the death assemblage included the skeletal material from mortality events spanning 5-10 generations, using the mean age of

reproductive maturity for *Acropora* species as the generation time (Van Oppen et al. 2000) [i.e. 3 to 8 years (Csaszar et al. 2010; Wallace, 1999)], a random sample of skeletons would be between 15 to 80 years old (Van Oppen et al. 2000). This is a much greater range than the two-component corrected 2σ age range observed for our ^{230}Th ages (11.6 ± 1.9 to 16.4 ± 5.8 years, or 1999.3 ± 1.9 to 1994.3 ± 5.8 AD). Having said that, there appears to be a slight tailing in the U-Th data in the relative probability plot of Figure 6d and the individual age plot of Figure S3, implying minor mortality that may have occurred during the 1994 bleaching/flood event. However, this has little impact on the well-defined 1998 AD age peak recorded in Figure 6d.

4. Conclusions

The congruence between the ^{230}Th age data corrected using the two-component equation and the documented catastrophic loss of *Acropora* both at Pandora Reef and over a much broader scale as a result of the 1998 bleaching event affirms that it is possible to use the U-Th method to accurately date recently dead coral skeletons from the death assemblage. For dead corals (including both massive and branching growth forms) from inshore reef settings, it is necessary to correct for both hydrogenous (dissolved) and detrital ^{230}Th incorporated during growth as well as post-mortem (see also Clark et al. 2014). This approach can then be used as a powerful tool for researchers and managers to identify mortality events and estimate rates of recovery in a historical context. For example, following the 1997-1998 bleaching event, it was predicted that it would not be until 2008-2010 that coral on the shallow fore-reef could recover to its 1981 status (Done et al. 2007). Our observations in 2008 and 2009 suggest that recovery severely lags behind this prediction due to an apparent failure of coral recruitment at Pandora Reef. However, for the vast majority of coral reefs there are no such long-term ecological data. On those reefs, high-precision U-Th dating with high sample throughput can now be used with surety on degraded coral reefs to determine when the reefs were damaged and hence ascertaining not only the drivers but also the time that has been available for their recovery post-disturbance; critical issues that have received insufficient attention in coral

reef science (Hughes et al. 2010). 'Time for recovery' is a key variable in evaluating a damaged reef's recovery performance against established benchmarks (Done et al. 2010). As much as it provides the timescale, high-precision U-Th dating thus has important applications that extend beyond scientific understanding and into the realm of coral reef policy and management.

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Supplementary Material

Supplementary material accompanies this paper

Figure captions

Figure 1. Map showing location of the Palm Islands in the central region of the Great Barrier Reef and an enlarged map of Pandora Reef. Sites marked as V1, V2, t1-4 and t5 (green lines) are the locations of video transects surveyed by AIMS from 1992 onwards. Site P1 (blue circle) is a photo transect monitored by T. Done from 1980-2005. Red bar represents the location where dead branching *Acropora* were sampled in this study. This site was also surveyed by DeVantier et al. (1997) in February and April 1994, however, quantitative data are unavailable.

Figure 2. a) Photograph of *Acropora* death assemblage overgrown by macroalgae taken in May 2008 from the south-west flank of Pandora Reef; b) Example of a dead arborescent *Acropora* branch collected for U-Th dating; c) Example of coral skeletal sample after pressurised cleaning with H_2O_2. Note the successful removal of detrital material. Scale represented by 5 mm grid paper.

Figure 3. $^{230}Th/^{232}Th$ versus $^{238}U/^{232}Th$ isochron for five coeval sets of sub-samples obtained from annual growth bands of dead *Porites* skeletons collected from the Palm Islands region, central Great Barrier Reef. Inset shows the isochron-inferred $^{230}Th/^{232}Th_0$ ratios (y-intercepts with 2σ errors) of the detrital component (average 0.61 ± 0.01 (1σ)).

Figure 4. Mixing diagram of $1/^{232}Th$ (ppb) plotted against $^{230}Th/^{232}Th$ (± 2σ) activity ratios using mean values obtained from live *Porites* ($^{230}Th/^{232}Th$ = 1.08 ± 0.19, ^{232}Th = 0.95 ppb; green line), live *Acropora* ($^{230}Th/^{232}Th$ = 3.5 ± 0.8, ^{232}Th = 0.15 ppb; blue line), and isochron-derived detrital ratio (similar to Burdekin River sediments) obtained from dead *Porites* ($^{230}Th/^{232}Th$ = 0.61 ± 0.02, $1/^{232}Th$ = 0 ppb; orange line). Isotopic data from all components show a negative correlation between two end-members: a detrital (sediment) phase and a hypothetical hydrogenous (seawater) phase (large open circle with a question mark) that is yet to be constrained. Following correction for $^{230}Th_0$ using a two-component mixing model, the measured $^{230}Th/^{232}Th$ ratios obtained from the dead *Acropora* samples (grey circles) shift towards and fall on the mixing line (grey triangles with error bars not shown for ease of interpretation), with most lying close to the sediment and live *Porites* values (see enlargement).

Figure 5. The effects of each correction scheme on ^{230}Th ages. a) Uncorrected b) bulk Earth c) live coral d) isochron-derived detrital component from dead *Porites* corals, and e) two-component corrected ^{230}Th ages versus ^{232}Th (ppb). If the bulk Earth- and live coral-based $^{230}Th/^{232}Th_0$ values were used for correction, the corrected ages show a positive relationship with measured ^{232}Th (r =

0.8670, P = <0.0001 and r = 0.9702, P = <0.0001, respectively). However, if the isochron-derived detrital $^{230}Th/^{232}Th_0$ value and our two-component mixing model $^{230}Th/^{232}Th_0$ value are used there is no correlation: (r = -0.2288 P = 0.1502 and r = -0.2361 P = 0.1372, respectively), suggesting that the presence of ^{232}Th (a proxy for the amount of initial ^{230}Th in a sample) has been appropriately corrected for. Dashed line represents the timing of the 1997-1998 bleaching event. Grey line represents an uncertainty of 0.3 years in order to account for systematic bias towards slightly older ages due to the material used for dating being taken from areas of the coral skeleton that was deposited in the months leading up to the coral colony's death. This value was based on average linear extension rates in branching *Acropora* corals typical of turbid, inshore, sheltered environments (~17.9 cm yr^{-1}; Crabbe and Smith, 2005) and taking into consideration the average sampling location with respect to the distance from the tip of the branch which was determined to be ~5.0 cm.

Figure 6. a) Annual Burdekin River discharge in mega litres for the years 1981-2005 measured at the Burdekin River station at Clare site 120006B (source: Queensland Department of the Environment and Resource Management); b) Maximum annual sea surface temperatures for 1×1 grid at 146.5°E, 18.5°S (Source: NOAA Reyn and SmithOlv2); c) Percent coral cover of the genus *Acropora*. Video footage was collected by the Australian Institute of Marine Science at sites V1, V2, t1-4 and t5. Photographs were taken by Done et al. (2007) at site P1 (see Fig. 1) d) Relative probability plot of 41 U-Th ages obtained from dead *Acropora* corals corrected for $^{230}Th/^{232}Th_0$ using a two-component (blue), isochron-derived detrital component (similar to Burdekin River sediments) (red), live coral (green) and Bulk Earth (yellow) value. The height and width of the curve represents the number of samples that date to the same time period and associated error, respectively. Orange vertical bars represent bleaching years, light blue bars represent years of major flooding and grey bars represent cyclone events

Figure 7. U and Th isotope measurements for 41 dead *Acropora* samples that reportedly died as a result of the 1998 bleaching event show a linear relationship in ($^{230}Th/^{232}Th$)-($^{238}U/^{232}Th$) space. This plot likely reflects a mixing line and is not a true isochron (due to multiple sources of $^{230}Th_0$). The y-intercept is equivalent to the $^{230}Th/^{232}Th_0$ in the detrital phase. This was determined to be 0.64 ± 0.04 (1σ), which is similar to isochron derived $^{230}Th/^{232}Th_{nr}$ values from dead *Porites* colonies (0.61 ± 0.01 (1σ)), as well as ICP-MS measurements of Burdekin River sediments (Th/U = 4.8 ± 1.0 or $^{230}Th/^{232}Th$ = 0.65 ± 0.2).

541 **Tables**

542 **Table 1.** MC-ICP-MS ^{230}Th ages for dead branching *Acropora* samples collected from the death assemblage at Pandora Reef, central Great Barrier
543 Reef**.**
544

Sample Name	Sampling range (cm)[b]	Sample wt.(g)	U (ppm)	^{232}Th (ppb)	(^{230}Th/^{232}Th)$_{meas}$	(^{230}Th/^{238}U)	$\delta^{234}U^{c}$	Uncorr. ^{230}Th age (AD)	Time of chemistry
PanS1T1.2	5.9	0.52861	3.1264 ± 0.0024	2.3013 ± 0.0048	1.336 ± 0.030	0.0003241 ± 0.0000069	147.0 ± 1.0	1979.8 ± 0.7	2010.7
PanS1T1.3	13.4	0.59958	3.0211 ± 0.0013	0.9677 ± 0.0023	2.362 ± 0.042	0.0002494 ± 0.0000040	146.7 ± 1.2	1986.9 ± 0.4	2010.7
PanS1T1.4	7.5	0.55553	3.1885 ± 0.0015	3.1981 ± 0.0039	1.271 ± 0.021	0.0004202 ± 0.0000066	147.2 ± 1.2	1970.7 ± 0.6	2010.7
PanS1T1.5	13.0	0.50465	3.0345 ± 0.0011	1.7522 ± 0.0023	1.607 ± 0.027	0.0003059 ± 0.0000049	147.0 ± 1.2	1981.6 ± 0.5	2010.7
PanS1T1.7	9.7	0.64273	3.1385 ± 0.0020	0.26074 ± 0.00038	6.75 ± 0.19	0.0001848 ± 0.0000054	145.4 ± 1.2	1993.1 ± 0.5	2010.7
PanS1T1.8	16.5	0.53981	3.0306 ± 0.0019	1.3201 ± 0.0018	1.841 ± 0.032	0.0002643 ± 0.0000045	145.9 ± 1.1	1985.5 ± 0.4	2010.7
PanS1T1.9	11.7	0.61789	3.2140 ± 0.0018	3.8856 ± 0.0051	1.050 ± 0.017	0.0004185 ± 0.0000065	146.8 ± 1.0	1970.8 ± 0.6	2010.7
PanS1T1.10	12.1	0.56444	3.1802 ± 0.0013	0.9354 ± 0.0014	2.407 ± 0.043	0.0002333 ± 0.0000040	145.5 ± 1.0	1988.5 ± 0.4	2010.7
PanS1T1.11	13.8	0.73242	3.1275 ± 0.0015	3.4069 ± 0.0031	1.108 ± 0.017	0.0003977 ± 0.0000059	145.0 ± 0.9	1972.8 ± 0.6	2010.7
PanS1T1.12	9.5	0.51359	3.1434 ± 0.0023	3.3456 ± 0.0062	1.192 ± 0.021	0.0004181 ± 0.0000070	146.1 ± 0.9	1970.9 ± 0.7	2010.7
PanS1T2.1	8.0	0.61479	3.1976 ± 0.0014	0.6177 ± 0.0011	3.890 ± 0.060	0.0002477 ± 0.0000036	146.8 ± 1.2	1987.1 ± 0.3	2010.7
PanS1T2.3	5.4	0.58241	3.1133 ± 0.0016	3.1317 ± 0.0032	1.169 ± 0.019	0.0003876 ± 0.0000063	146.5 ± 1.2	1973.8± 0.6	2010.7
PanS1T2.4	10.8	0.51142	3.2295 ± 0.0016	1.5958 ± 0.0015	1.707 ± 0.031	0.0002779 ± 0.0000050	146.5 ± 1.2	1984.2± 0.5	2010.7
PanS1T2.5	12.4	0.54179	2.9378 ± 0.0017	0.45549 ± 0.00067	4.055 ± 0.086	0.0002072 ± 0.0000043	147.5 ± 1.0	1991.0 ± 0.4	2010.7
PanS1T2.6	8.0	0.57222	2.9336 ± 0.0015	0.43871 ± 0.00060	4.079 ± 0.098	0.0002010 ± 0.0000048	146.7 ± 1.1	1991.6 ± 0.5	2010.7
PanS1T2.7	8.9	0.50702	3.3822 ± 0.0014	1.7674 ± 0.0016	1.616 ± 0.035	0.0002784 ± 0.0000060	146.0 ± 0.9	1984.2 ± 0.6	2010.7
PanS1T2.8	6.8	0.53453	2.9511 ± 0.0015	3.6596 ± 0.0031	1.155 ± 0.018	0.0004719 ± 0.0000073	146.3 ± 0.8	1965.7 ± 0.7	2010.7
PanS1T2.9	5.9	0.55237	3.2718 ± 0.0015	0.9260 ± 0.0013	2.730 ± 0.062	0.0002546 ± 0.0000056	145.7 ± 1.2	1986.4 ± 0.5	2010.7
PanS1T2.10	7.4	0.57864	2.9315 ± 0.0020	1.1244 ± 0.0012	2.029 ± 0.047	0.0002565 ± 0.0000060	145.9 ± 1.4	1986.3 ± 0.6	2010.7
PanS1T2.11	8.0	0.51666	3.1868 ± 0.0016	1.9400 ± 0.0020	1.495 ± 0.028	0.0002999 ± 0.0000056	146.3 ± 1.0	1982.1 ± 0.5	2010.7
PanS1T2.12	6.4	0.65912	3.2470 ± 0.0016	2.6622 ± 0.0026	1.283 ± 0.019	0.0003466 ± 0.0000050	146.0 ± 1.2	1977.7 ± 0.5	2010.7
PanS1T3.3	11.8	0.53734	3.2765 ± 0.0016	0.49495 ± 0.00057	4.211 ± 0.078	0.0002096 ± 0.0000039	147.0 ± 1.2	1990.9 ± 0.4	2010.9
PanS1T3.4	14.5	0.55105	3.1239 ± 0.0015	1.5117 ± 0.0036	1.768 ± 0.027	0.0002819 ± 0.0000039	146.6 ± 0.8	1984.1 ± 0.4	2010.9
PanS1T3.5	8.0	0.53053	3.1027 ± 0.0014	0.30429 ± 0.00060	6.51 ± 0.16	0.0002104 ± 0.0000050	146.7 ± 0.8	1990.9 ± 0.5	2010.9
PanS1T3.6	5.2	0.51268	3.2460 ± 0.0020	0.50127 ± 0.00076	4.378 ± 0.083	0.0002228 ± 0.0000041	146.2 ± 0.9	1989.7 ± 0.4	2010.9
PanS1T3.7	9.2	0.56672	3.2631 ± 0.0020	0.81785 ± 0.00090	2.641 ± 0.058	0.0002182 ± 0.0000048	147.0 ± 1.2	1990.1± 0.5	2010.9
PanS1T3.8	15.5	0.52514	3.2931 ± 0.0016	0.60433 ± 0.00049	3.512 ± 0.065	0.0002124 ± 0.0000040	147.0 ± 0.8	1990.7 ± 0.4	2010.9
PanS1T3.11	14.5	0.53172	3.3683 ± 0.0016	2.3049 ± 0.0017	1.423 ± 0.020	0.0003210 ± 0.0000046	147.2 ± 1.0	1980.3 ± 0.4	2010.9
PanS1T3.13	5.7	0.51982	3.2520 ± 0.0018	3.2679 ± 0.0035	1.168 ± 0.021	0.0003867 ± 0.0000069	146.2 ± 0.7	1974.1 ± 0.7	2010.9

PanS1T3.15	7.5	0.63710	3.3315 ± 0.0014	2.7712 ± 0.0032	1.229 ± 0.018	0.0003368 ± 0.0000048	146.4 ± 0.8	1978.8 ± 0.5	2010.9
PanS1T4.1	12.7	0.54322	3.2816 ± 0.0015	0.54822 ± 0.00055	4.136 ± 0.064	0.0002277 ± 0.0000035	147.7 ± 0.9	1989.2± 0.3	2010.9
PanS1T4.3	12.0	0.54173	3.1571 ± 0.0013	1.4899 ± 0.0013	1.746 ± 0.032	0.0002716 ± 0.0000050	147.5 ± 0.9	1985.1 ± 0.5	2010.9
PanS1T4.4	9.7	0.51410	3.2284 ± 0.0010	0.85858 ± 0.00081	2.718 ± 0.047	0.0002382 ± 0.0000041	147.9 ± 0.8	1988.2 ± 0.4	2010.9
PanS1T4.5	13.5	0.52256	3.1620 ± 0.0013	1.0616 ± 0.0021	2.207 ± 0.042	0.0002442 ± 0.0000044	146.7 ± 0.9	1987.6 ± 0.4	2010.9
PanS1T4.6	6.2	0.54909	3.1463 ± 0.0016	0.34359 ± 0.00039	5.93 ± 0.10	0.0002133 ± 0.0000036	147.8 ± 1.2	1990.6 ± 0.3	2010.9
PanS1T4.7	14.0	0.54121	3.0927 ± 0.0021	0.81946 ± 0.00083	3.059 ± 0.058	0.0002671 ± 0.0000051	148.9 ± 0.9	1985.5 ± 0.5	2010.9
PanS1T4.9	10.8	0.79178	3.2502 ± 0.0019	0.8433 ± 0.0014	2.792 ± 0.044	0.0002387 ± 0.0000036	146.8 ± 1.0	1988.2 ± 0.3	2010.9
PanS1T4.10	7.5	0.70195	3.2810 ± 0.0015	1.5303 ± 0.0013	1.792 ± 0.028	0.0002755 ± 0.0000042	146.9 ± 1.1	1984.7 ± 0.4	2010.9
PanS1T4.12	9.0	0.59781	3.2461 ± 0.0015	0.63972 ± 0.00085	3.424 ± 0.059	0.0002224 ± 0.0000037	146.9 ± 0.9	1989.7 ± 0.4	2010.9
PanS1T4.13	11.0	0.65207	3.2151 ± 0.0014	0.58939 ± 0.00080	3.560 ± 0.059	0.0002151 ± 0.0000035	146.5 ± 1.1	1990.4 ± 0.3	2010.9
PanS1T4.17	8.3	0.54801	3.2038 ± 0.0015	2.0043 ± 0.0035	1.479 ± 0.028	0.0003050 ± 0.0000054	145.9 ± 1.0	1981.8 ± 0.5	2010.9

Ratios in parentheses are activity ratios calculated from atomic ratios using decay constants of Cheng et al. (2000b). All values have been corrected for laboratory procedural blanks. All errors reported as 2σ. Uncorrected ^{230}Th age (a) was calculated using Isoplot/EX 3.0 program (Ludwig, 2003b), where *a* denotes year.

[a]For the sample nomenclature, S1 refers to Site 1 where the samples were collected at Pandora Reef. T1-T4 refers to transects 1-4 which were each 20 m in length. The number after the decimal refers to the individual *Acropora* branch dated from that particular transect.

[b] Sampling range where material for U-Th dating was collected with respect to distance from the tip of the branch, or end of the branch where the tip has broken off, in centimetres. To ensure a 0.5 – 1.0 g sample size free from alteration, it was not possible to sample from a single location.

[c]$\delta^{234}U = [(^{234}U/^{238}U)-1]\times1000$

Table 2. MC-ICP-MS ^{230}Th ages for dead *Acropora* coral samples collected in May 2008 corrected using the bulk Earth, live coral, sediment and two-component $^{230}Th/^{232}Th_0$ correction value

Sample Name	Uncorr. ^{230}Th Age (AD)	Bulk Earth (AD)[a]	Live coral (AD)[b]	Sediment (AD)[c]	Two-component (AD)[d]
PanS1T1.2	1979.8 ± 0.7	1998.9 ± 9.6	2004.9 ± 5.0	1993.9 ± 2.9	1998.4 ± 3.8
PanS1T1.3	1986.9 ± 0.4	1995.2 ± 4.2	1997.8 ± 2.2	1993.1 ± 1.3	1997.7 ± 2.2
PanS1T1.4	1970.7 ± 0.6	1997 ± 13	2004.8 ± 6.8	1989.9 ± 3.9	1994.3 ± 4.8
PanS1T1.5	1981.6 ± 0.5	1996.5 ± 7.5	2001.2 ± 4.0	1992.6 ± 2.3	1997.3 ± 3.2
PanS1T1.7	1993.1 ± 0.5	1995.2 ± 1.2	1995.9 ± 0.8	1994.7 ± 0.6	1999.2 ± 1.3
PanS1T1.8	1985.5 ± 0.4	1996.8 ± 5.7	2000.3 ± 3.0	1993.9 ± 1.7	1998.5 ± 2.6
PanS1T1.9	1970.8 ± 0.6	2002 ± 15	-ve age	1994.0 ± 4.7	1998.4 ± 5.5
PanS1T1.10	1988.5 ± 0.4	1996.1 ± 3.8	1998.5 ± 2.0	1994.1 ± 1.2	1998.5 ± 2.1
PanS1T1.11	1972.8 ± 0.6	2001± 14	2009.9 ± 7.4	1993.7 ± 4.2	1998.2 ± 5.1
PanS1T1.12	1970.9 ± 0.7	1998 ± 14	2007.1 ± 7.3	1991.2 ± 4.1	1995.7 ± 5.0
PanS1T2.1	1987.1 ± 0.3	1992.1 ± 2.5	1993.7 ± 1.4	1990.8 ± 0.8	1995.2 ± 1.7
PanS1T2.3	1973.8± 0.6	2000 ± 13	2008.0 ± 6.9	1993.0 ± 3.9	1997.6 ± 4.8
PanS1T2.4	1984.2± 0.5	1997.0 ± 6.4	2001.0 ± 3.4	1993.7 ± 2.0	1998.1 ± 2.8
PanS1T2.5	1991.0 ± 0.4	1995.0± 2.0	1996.2 ± 1.1	1993.9± 0.7	1998.7 ± 1.6
PanS1T2.6	1991.6 ± 0.5	1995.4 ± 2.0	1996.6 ± 1.1	1994.4 ± 0.7	1999.2 ± 1.6
PanS1T2.7	1984.2 ± 0.6	1997.7 ± 6.8	2001.9 ± 3.6	1994.2 ± 2.1	1998.4 ± 2.9
PanS1T2.8	1965.7 ± 0.7	1998 ± 16	2007.9 ± 8.5	1989.5 ± 4.8	1994.3 ± 5.7
PanS1T2.9	1986.4 ± 0.5	1993.8 ± 3.7	1996.1 ± 2.0	1991.9 ± 1.2	1996.2 ± 2.0
PanS1T2.10	1986.3 ± 0.6	1996.2 ± 5.0	1999.3 ± 2.7	1993.6 ± 1.6	1998.4 ± 2.5
PanS1T2.11	1982.1 ± 0.5	1997.9 ± 7.9	2002.8 ± 4.2	1993.8 ± 2.4	1998.2 ± 3.3
PanS1T2.12	1977.7 ± 0.5	1999 ± 11	2005.6 ± 5.6	1993.4 ± 3.2	1997.7 ± 4.0
PanS1T3.3	1990.9 ± 0.4	1994.9 ± 2.0	1996.1 ± 1.1	1993.8 ± 0.7	1998.1 ± 1.5
PanS1T3.4	1984.1 ± 0.4	1996.6 ± 6.3	2000.5 ± 3.3	1993.3 ± 1.9	1997.8 ± 2.8
PanS1T3.5	1990.9 ± 0.5	1993.4 ± 1.4	1994.2 ± 0.8	1992.7 ± 0.6	1997.3 ± 1.4
PanS1T3.6	1989.7 ± 0.4	1993.7 ± 2.0	1994.9 ± 1.1	1992.6 ± 0.7	1997.0 ± 1.5
PanS1T3.7	1990.1± 0.5	1996.6 ± 3.3	1998.6 ± 1.8	1994.9 ± 1.1	1999.2 ± 1.9
PanS1T3.8	1990.7 ± 0.4	1995.4 ± 2.4	1996.9 ± 1.3	1994.2 ± 0.8	1998.5 ± 1.6
PanS1T3.11	1980.3 ± 0.4	1998.1 ± 8.9	2003.6 ± 4.7	1993.4 ± 2.7	1997.6 ± 3.5
PanS1T3.13	1974.1 ± 0.7	2000 ± 13	2008.2 ± 6.9	1993.3 ± 3.9	1997.6 ± 4.8
PanS1T3.15	1978.8 ± 0.5	2000 ± 11	2007.1 ± 5.7	1994.7 ± 3.2	1999.0 ± 4.1
PanS1T4.1	1989.2± 0.3	1993.6 ± 2.2	1994.9± 1.2	1992.4 ± 0.7	1996.7 ± 1.5
PanS1T4.3	1985.1 ± 0.5	1997.3 ± 6.1	2001.1 ± 3.2	1994.1 ± 1.9	1998.5± 2.7
PanS1T4.4	1988.2 ± 0.4	1995.1 ± 3.5	1997.3 ± 1.8	1993.3 ± 1.1	1997.7 ± 1.9
PanS1T4.5	1987.6 ± 0.4	1996.3± 4.4	1999.1 ± 2.3	1994.1 ± 1.4	1998.5 ± 2.2
PanS1T4.6	1990.6 ± 0.3	1993.4 ± 1.5	1994.3 ± 0.8	1992.7 ± 0.5	1997.2 ± 1.4
PanS1T4.7	1985.5 ± 0.5	1992.4 ± 3.5	1994.5 ± 1.9	1990.6 ± 1.1	1995.1 ± 2.0
PanS1T4.9	1988.2 ± 0.3	1994.9 ± 3.4	1997.0 ± 1.8	1993.1 ± 1.1	1997.5 ± 2.0
PanS1T4.10	1984.7 ± 0.4	1996.7 ± 6.1	2000.5 ± 3.2	1993.6 ± 1.8	1997.9 ± 2.7
PanS1T4.12	1989.7 ± 0.4	1994.8 ± 2.6	1996.4 ± 1.4	1993.5 ± 0.8	1997.8 ± 1.7
PanS1T4.13	1990.4 ± 0.3	1995.2 ± 2.4	1996.6 ± 1.3	1993.9 ± 0.8	1998.3 ± 1.6
PanS1T4.17	1981.8 ± 0.5	1998.0 ± 8.1	2003.1 ± 4.3	1993.8 ± 2.5	1998.2 ± 3.3

Uncorrected ^{230}Th age (AD) was calculated using Isoplot/EX 3.0 program (Ludwig, 2003b).
Corrected ^{230}Th ages were calculated using:
[a] Bulk Earth value = 0.82 ± 50% (atomic value ~4.4×10^{-6} ± 50%).
[b] Region specific $^{230}Th/^{232}Th_0$ value for the Palm Islands derived from live *Porites* of known age = 1.083 ± 20% (atomic value of 5.7×10^{-6} ± 20%)
[c] Burdekin River sediment value derived from 40 ICP-MS measurements =0.61 ± 20% (atomic value 3.53×10^{-6} ± 20%)
[d] Two-component correction value calculated using Equation (1).

Table 3. Summary statistics of ^{230}Th ages derived from 41 dead branching *Acropora* corals.

Correction used	Mean age (A.D.)	Median age (A.D.)	S.D.	Age range (A.D.)	Weighted mean age ± 2σ (A.D.)[a]	MSWD[b]
Uncorrected	1983.9	1985.5	7.0	1965.7-1993.1	1985.8 ± 1.8	671
Bulk Earth	1996.5	1996.5	2.3	1992.1-2002.1	1994.7 ± 0.5	0.8
Live coral	2000.1	1999.2	4.6	1993.7-	1996.2 ± 0.7	7.4
Sediment	1993.2	1993.6	1.3	1989.5-1994.9	1993.3 ± 0.3	3.8
Two-component	1997.7	1997.9	1.2	1994.3-1999.2	1997.8 ± 0.3	1.1

[a] Weighted mean calculated using Isoplot/Ex (Ludwig, 2003b).
[b] MSWD = Mean Square of Weighted Deviates. The MSWD is the sum of squares of weighted residuals divided by the degrees of freedom. MSWD values greater than unity (i.e. >1) indicate either underestimated analytical errors, or the presence of non-analytical 'geological' scatter (Ludwig, 2003b).

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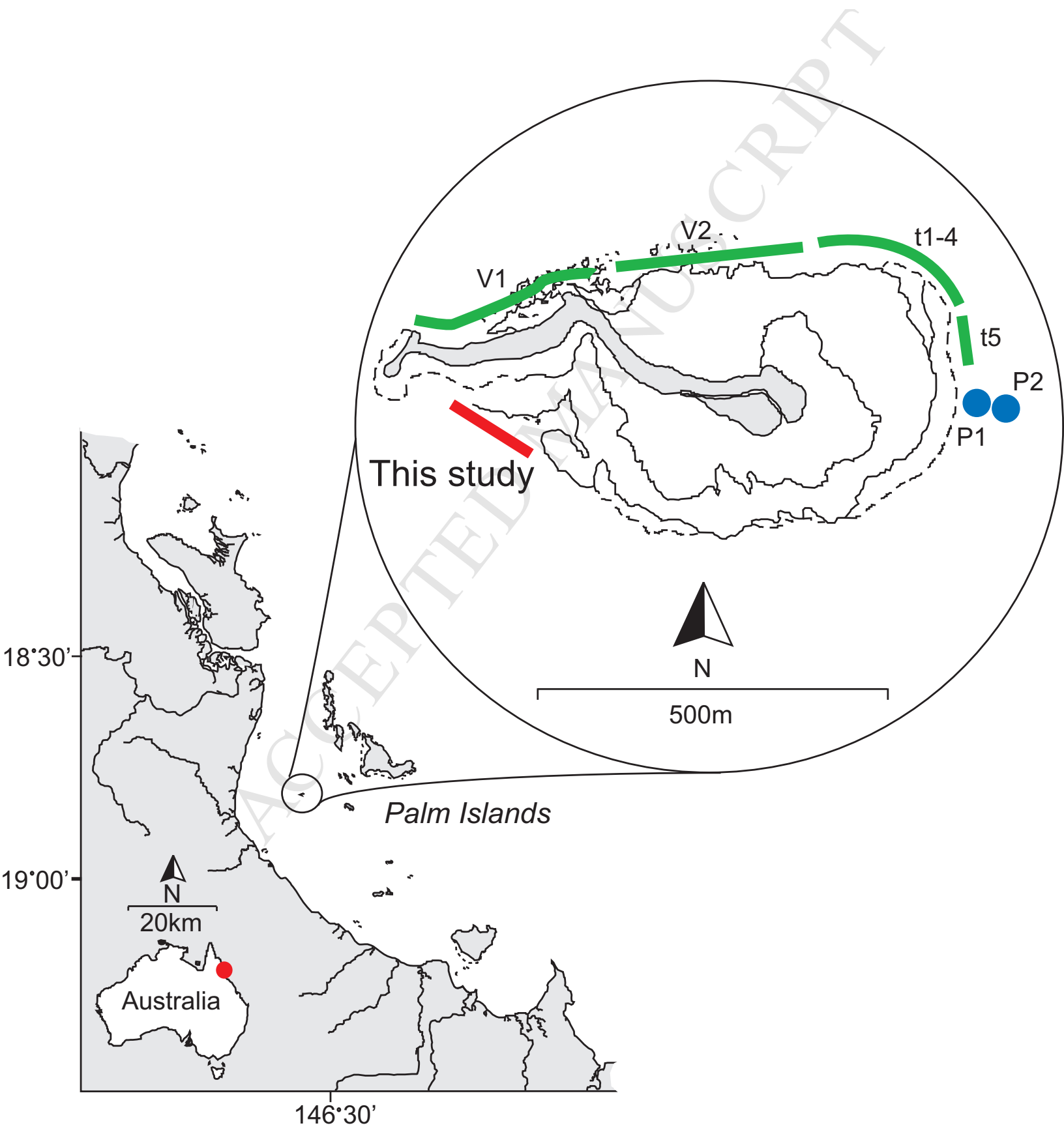
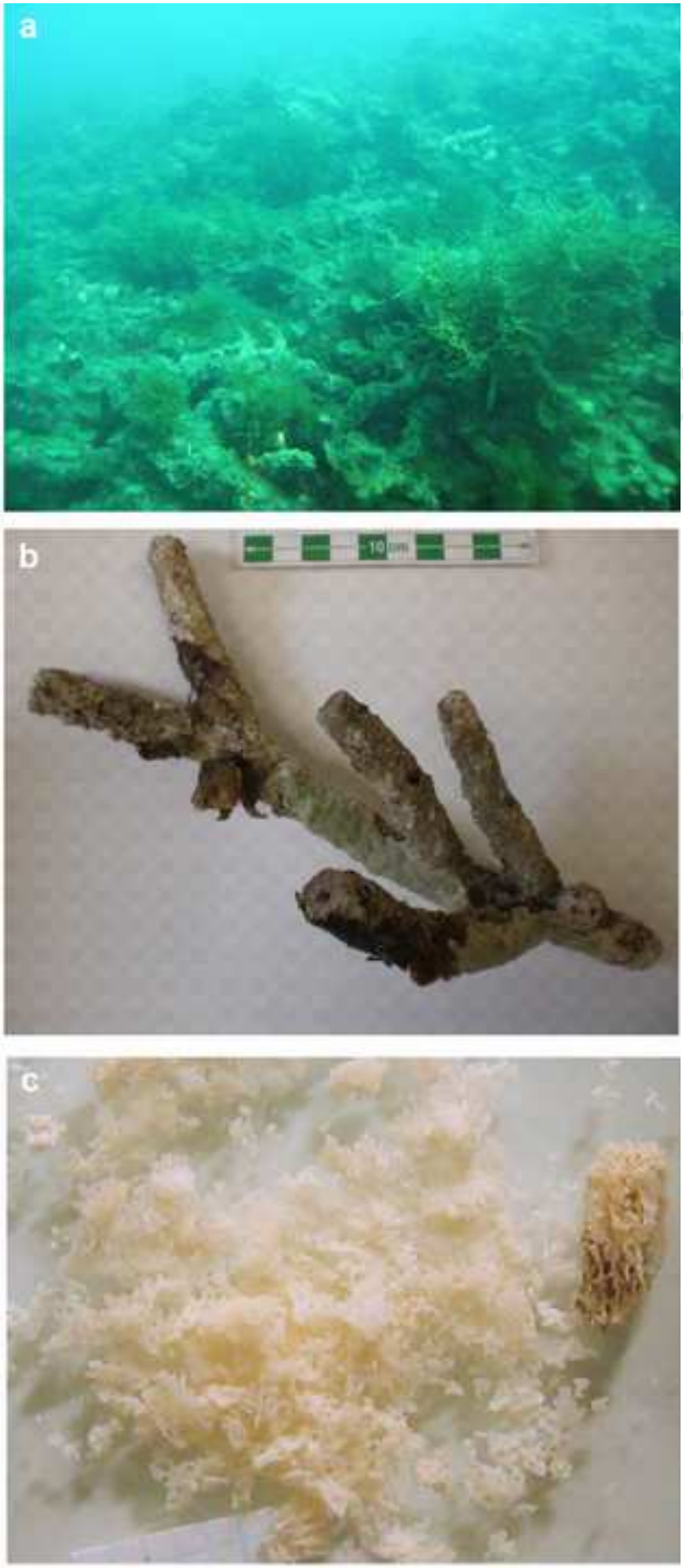
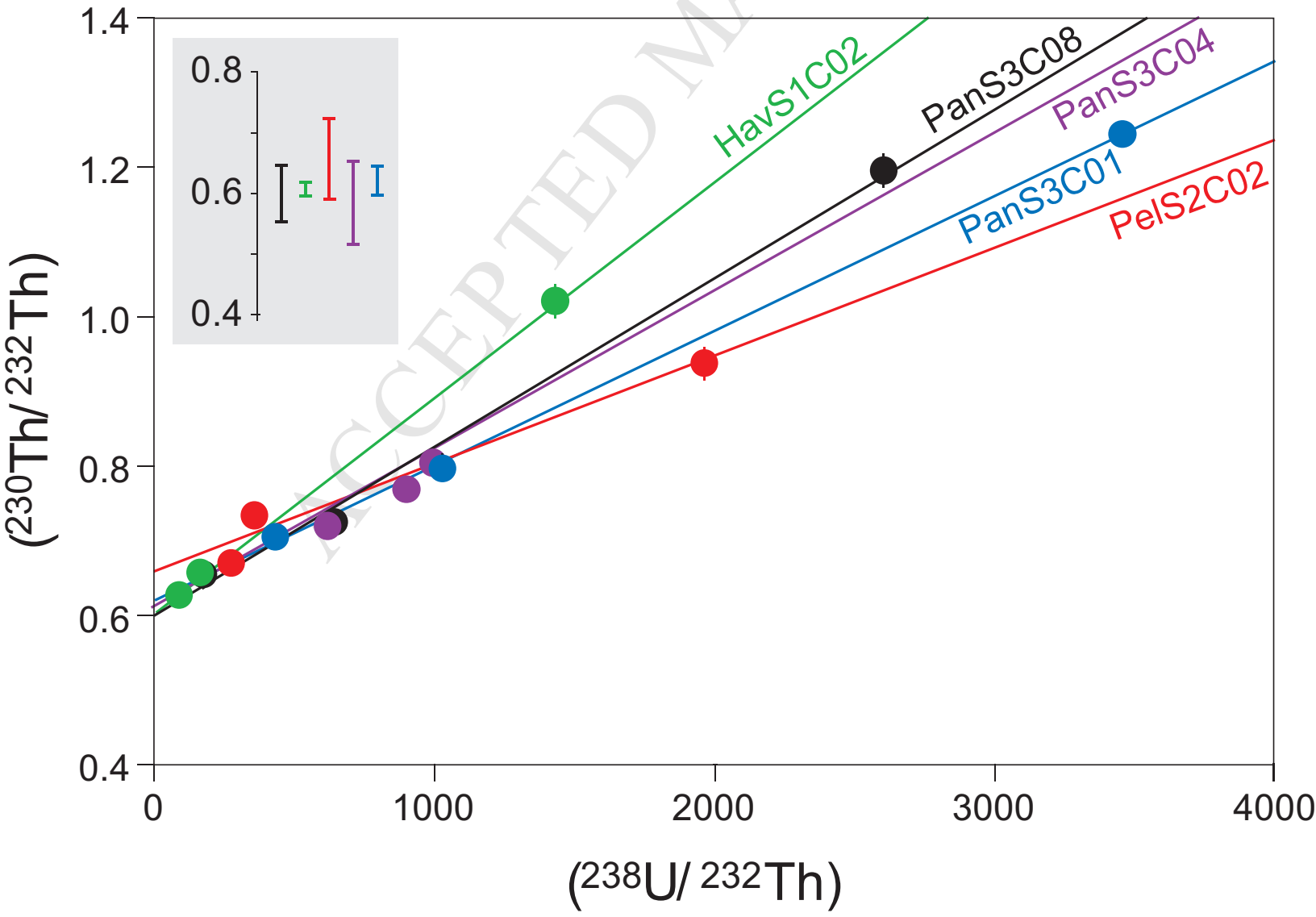
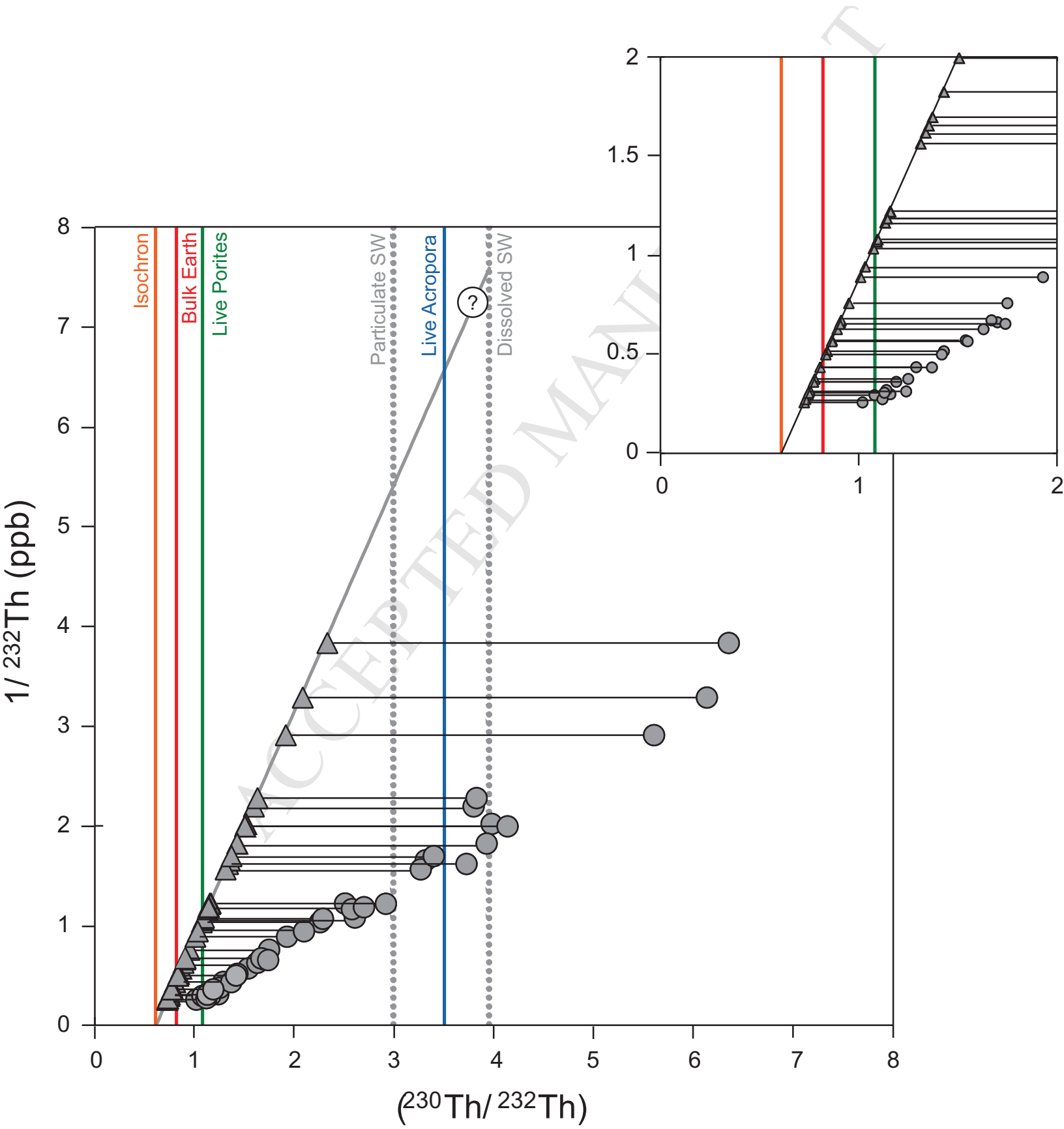




Figure



Figure



Figure