



1 **Technical note: „U-Th Analysis” – an open-source software**
2 **dedicated to MCICPMS U-series-data treatment and**
3 **evaluation**

4
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9 **Abstract**

10 We present our standalone data analysis application for ²³⁰Th/U dating on multi-collector inductively coupled
11 plasma mass spectrometers (MC-ICP-MS). The Python-based algorithm is equipped with a graphical user
12 interface (GUI) and comprises raw data treatment, corrections, age calculation, and error estimation. Our
13 underlying measurement protocol employs a combination of Faraday cups (FC) and secondary electron multipliers
14 (SEM), and the software allows for different detector layouts for the measurement of the least abundant isotopes
15 ²³⁴U, ²³⁰Th and ²²⁹Th. We especially focus on features that ensure reproducibility and enable user-friendly
16 reanalysis of measurements such as customized calculation constants with templates. Result files are saved
17 automatically and contain all relevant settings used. Eventually, we demonstrate the relevance of adequate data
18 outlier treatment and generally recommend using the median instead of the mean of calculated ratios. The
19 performance of our evaluation software is demonstrated by a case study from a Puerto Rican stalagmite with
20 growth phases from modern to 40 ka old. The majority of the obtained ages reaches uncertainties in the range of
21 0.3-0.6%, underlining the capability of our measurement protocol.

22 **1 Introduction**

23 The U-series disequilibrium method, ²³⁰Th/U dating, is a precise chronometer covering approximately the last 650
24 kiloyears, and has proven indispensable for the age determination of marine and continental carbonate archives
25 and their applications (Bourdon et al., 2003). The method is based on a complete disequilibrium of ²³⁴U, with its
26 daughter nuclide ²³⁰Th, during the formation of secondary carbonates. It presumes a subsequent closed system
27 evolution of the activity ratio of (²³⁰Th/²³⁴U) and (²³⁴U/²³⁸U) since the time of formation. Ideally, the initial ²³⁰Th
28 activity of the material is presumed zero or can be estimated from the total Th concentration via an initial
29 (²³⁰Th/²³²Th) activity ratio. The dating applications for secondary carbonates and other appropriate materials are
30 manifold in geochemistry, archaeology, and climate science. The applications relying on this dating method are
31 manifold in geochemistry, archaeology and climate science. Further development of this dating method includes
32 both improvements in instrumentation and measurement protocols, as well as reproducible data analysis and age
33 calculation schemes (Pourmand et al., 2014; Andersen et al., 2004; Cheng et al., 2013; Breton et al., 2015; Chiang
34 et al., 2019; Hellstrom, 2003; Hoffmann et al., 2007; Shen et al., 2002; Shen et al., 2012; Kerber et al., 2023; Shao et
35 al., 2019). The presently most sensitive and precise technology for high precision U and Th isotope measurements
36 is multi-collector inductively coupled plasma source mass spectrometry (MC-ICPMS). Recent technological



37 advances of MC-ICP-MS include the implementation of high ohmic amplifiers allowing to enhance the dynamic
38 range of multiple Faraday-collectors (FC) to six orders of magnitude for the simultaneous detection of very large
39 and low isotope abundances, instead of the conventionally used combination of secondary electron multipliers
40 (SEM) and FC (Breton et al., 2015). Measurement protocol updates aim at increasing measurement precision
41 and/or decreasing input sample masses by combining new detector layouts, improving the understanding of
42 correction factors, and ensuring a stable measurement environment (Cheng et al., 2013; Chiang et al., 2019; Shen
43 et al., 2002; Shen et al., 2012; Hellstrom, 2003; Andersen et al., 2004; Hoffmann et al., 2007; Kerber et al.,
44 2023; Shao et al., 2019).

45 We here focus on the third route for the enhancement of $^{230}\text{Th}/\text{U}$ dating, which is clear and reproducible data
46 analysis and age calculation schemes. Up to now, only two $^{230}\text{Th}/\text{U}$ dating data analysis routines have been
47 published (Shao et al., 2019; Pourmand et al., 2014). However, regarding the rising amount of data being produced
48 in MC-ICP-MS laboratories, data management is becoming more and more important. For example, some samples
49 might require later adaptation of the individual corrections of isotope ratios due to residual contamination with
50 non-carbonate material or detection of initial ^{230}Th from the carbonate forming environment.

51 Dating young materials of only a few years to centuries in age is challenging (i) due to the small number of counts
52 on especially ^{230}Th , which implies that all correction factors including “ghost signal” corrections need to be
53 determined very precisely (Zhao et al., 2009; Chiang et al., 2019; Kerber et al., 2023). Regarding the removal of
54 scatter ions on the specific low abundance masses 230 and 229 amu, Kerber et al. (2023) demonstrated an effective
55 correction based on a linear dependence of the scattered ions on the ^{238}U signal. Other authors separate U and Th
56 chemically to reduce or remove the ^{238}U beam from the low abundance Thorium isotope measurements (Chiang
57 et al., 2019), which implies flexibility in the detector arrangement and data treatment protocol. As such scatter
58 peaks may depend on the specific instrument or vary through time, these corrections need to be adaptable constants
59 in the data evaluation routine. The influence on final atomic ratio and accuracy of the ghost signals as well as by
60 typical variation in other individual corrections such as peak tailing, mass fractionation, isobaric interferences are
61 evaluated in detail in Kerber et al. (2023).

62 In addition, the correction for initial Thorium may cause large age corrections and propagated uncertainties, in
63 particular since adequate initial Th values based on the $^{230}\text{Th}/^{232}\text{Th}$ ratio may be variable and difficult to detect
64 (Hellstrom, 2006; Wenz et al., 2016; Wortham et al., 2022). There are different methods to estimate the initial Th
65 isotope ratio: First, isochrons can be used to determine the isotopic composition of the detrital component in the
66 carbonate (Ludwig and Titterton, 1994; Wenz et al., 2016; Stinnesbeck et al., 2020; Töchterle et al., 2022).
67 Secondly, analyses of modern drip waters or recent carbonate deposits allow estimation of the value and sources
68 of initial Th (Wortham et al., 2022; Li et al., 2022). In some cases the “true” age of a stalagmite can be also inferred
69 from other dating methods, such as radiocarbon (Akers et al., 2019; Huang et al., 2024) or the stratigraphic order
70 (Hellstrom, 2006). Also, several approaches can be combined (Warken et al., 2020; Akers et al., 2016; Roy-Barman
71 and Pons-Branchu, 2016).

72 Other aspects are updating half-lives (such as e. g. from Cheng et al. (2000b) to Cheng et al. (2013)), which makes
73 re-evaluation of previously measured data necessary. These tasks are error-prone, in particular when they require
74 copy-and-pasting data in e. g. spreadsheets. Also, a clear and unified documentation of the applied constants and
75 the way of saving data is desirable. Additionally, the statistical methods, for example for outlier correction, should



76 undergo clear documentation. Altogether, this helps to report Th/U ages in a standardized way (Dutton et al.,
77 2017).

78 In this study, we present our user-friendly GUI and the underlying algorithm for data treatment and age
79 calculation. Methods to treat outliers in measurement data are particularly highlighted. As a case study, we present
80 newly obtained ages from a stalagmite from Larga Cave, Puerto Rico, which shows a modern growth phase, as
81 well as continuous deposition during the last Glacial into the deglaciation, thus demonstrating the performance of
82 our method for both very young and older sample materials. Our protocol enables a precise determination of
83 speleothem growth rates, which allows a comparison to a coevally deposited stalagmite from the same cave
84 highlighting the influence of in-cave processes on speleothem growth rates. In particular, this dataset showcases
85 how initial ^{230}Th correction models can be easily tested with our here presented software and GUI, and how those
86 influence speleothem chronologies.

87 2 Methods

88 2.1 Standards and reference materials

89 We use our in-house triple spike solution (TriSpike) with a ^{233}U concentration of (0.038556 ± 0.0000009) ng/g, a
90 ^{236}U concentration of (3.86778 ± 0.00009) ng/g and a ^{229}Th concentration of (0.018055 ± 0.000008) ng/g (2
91 standard error of the mean) (Kerber et al., 2023). For standard bracketing, we employ the Harwell-Uraninite 1
92 (HU-1) as a reference material. Its activity ratios ($^{230}\text{Th}/^{238}\text{U}$) and ($^{234}\text{U}/^{238}\text{U}$) are presumed to be 1, as it is a secular
93 equilibrium material. Abundance sensitivity and hydride correction are determined by measuring CRM-112A U
94 reference solution and an in-house ^{232}Th standard. The CRM-112A gravimetric standard solution has a ^{238}U
95 concentration of (4.3021 ± 0.0015) $\mu\text{g/g}$, while the inhouse ^{232}Th standard calibrated with TriSpike has a ^{232}Th
96 concentration of (505.8 ± 1.02) ng/g (2 σ uncertainties). CRM-112A solution is also used to track the values of
97 the two ghost signal constants, k_{229} and k_{230} (Kerber et al., 2023). For k_{229} , it is measured without addition of
98 TriSpike, while in the case of k_{230} , the spiked CRM-112A solution is employed. For age determination, the ^{230}Th
99 and ^{234}U decay constants determined by Cheng et al. (2013) are used. Ages are reported with 2 σ statistical standard
100 mean error, but do not include half-life uncertainties.

101 2.2 Chemical preparation and instrumentation

102 The chemical preparation of carbonate samples includes sample dissolution in ultra clean nitric acid, spiking with
103 TriSpike and two steps of wet column chromatographic ion exchange separation of U and Th from matrix elements
104 using Eichrom UTEVA resin (Douville et al., 2010; Wefing et al., 2017; Matos et al., 2015). Chemical blanks are
105 commonly below 0.4 fg for ^{234}U and 0.04 fg for ^{230}Th and Ca matrix concentrations are required to be below 10
106 ppm. For the mass-spectrometric measurement, samples are dissolved in 1 % HNO_3 and 0.05 % HF. All samples
107 were measured on a MC-ICP-MS (ThermoFisher Neptune Plus) at the Institute for Environmental Physics,
108 Heidelberg University (Germany). The mass spectrometer is equipped with Faraday cups (FC) and a central
109 secondary electron multiplier (SEM). The central detector can be selected between the SEM and a FC connected
110 to a 10^{13} Ω amplifier. ^{238}U is measured on a 10^{10} Ω amplified resistor. All other FC are connected to 10^{11} Ω
111 amplifiers. The desolvating system CETAC Aridus II is used as inlet. A measurement sequence starts with the
112 determination of abundance sensitivity and tailing on two different solutions for both uranium and thorium. Each
113 sample and standard measurement is preceded by a procedural blank measurement to ensure that the background



114 signal has gone back to a clean state. CRM-112A measurements are carried out to track ghost signal values at the
115 beginning and end of a measurement sequence. Samples are bracketed with HU-1 as a reference material. The
116 GUI is written for this type of measurement protocol. Other adaptations such as fewer procedural blank
117 measurements or else require small changes in the code, but are easily feasible.

118 **2.3 Speleothem sample description**

119 Stalagmite B1 was collected in 2019 in Larga cave, Puerto Rico (18°19'N 66°48'W, 350msl, supplementary Figure
120 S1A) from a passage in the deep part of the cave connected to the “Collapse room”. The host rock overburden at
121 the location of the sample is about 40-60m. It is in total 60 cm long, and has an average diameter of 15 cm
122 (supplementary Figure S1B). The drip site was still active, and was monitored with spot measurements over
123 several years, revealing varying drip intervals between 2 s and >120 s. A total amount of 37.7 ml water from the
124 drip site of stalagmite B1 was analysed for its U and Th activity ratios. Samples for ²³⁰Th/U dating of the
125 speleothem with typical input masses of 100-150 mg have been cut using a diamond wire saw along the growth
126 axis. Chemical preparation, mass-spectrometric measurements, data treatment and evaluation of drip water and
127 the speleothem samples followed the methods described in Kerber et al. (2023) and in this study.

128 Larga Cave is located in the north central karst region of Puerto Rico (supplementary Figure S1A). Previous work
129 including extensive cave air and drip monitoring has demonstrated that the cave is a valuable location to study of
130 the influence of changing climate on past rainfall patterns in the Western tropical Atlantic (Vieten et al.,
131 2018b; Warken et al., 2022a; Vieten et al., 2018a). In particular, the main passage of Larga Cave is subject to a
132 seasonally varying ventilation, which results in pCO₂ values of 600 ppm close to atmospheric values during
133 winter, and higher values up to 1800 ppm in summer (Vieten et al., 2016). In contrast, in the deep part of the cave,
134 where also stalagmite B1 was collected, ventilation is strongly muted, and cave air pCO₂ values are higher with
135 values up to 2300 – 3600 ppm (Vieten et al., 2016). As a result of this ventilation regime, growth rates are expected
136 to vary both seasonally, but also between different locations inside the cave (Vieten and Hernandez, 2021).

137 So far, two speleothem records from Larga Cave are been published, where the most recent covers the past 500
138 years (Vieten et al., 2024), and the second stalagmite grew during the period of 46.2-15.3 ka with a hiatus from
139 41.1 to 35.5 ka (Warken et al., 2020). For ²³⁰Th/U dating of Larga speleothems, high initial Th contents have to
140 be considered - a phenomenon that regularly occurs in speleothem records from the Caribbean and Central
141 American region (Fensterer et al., 2010; Steidle et al., 2021; Moseley et al., 2015; Schorndorf et al.,
142 2023; Stinnesbeck et al., 2020; Beck et al., 2001; Akers et al., 2016; Rivera-Collazo et al., 2015).

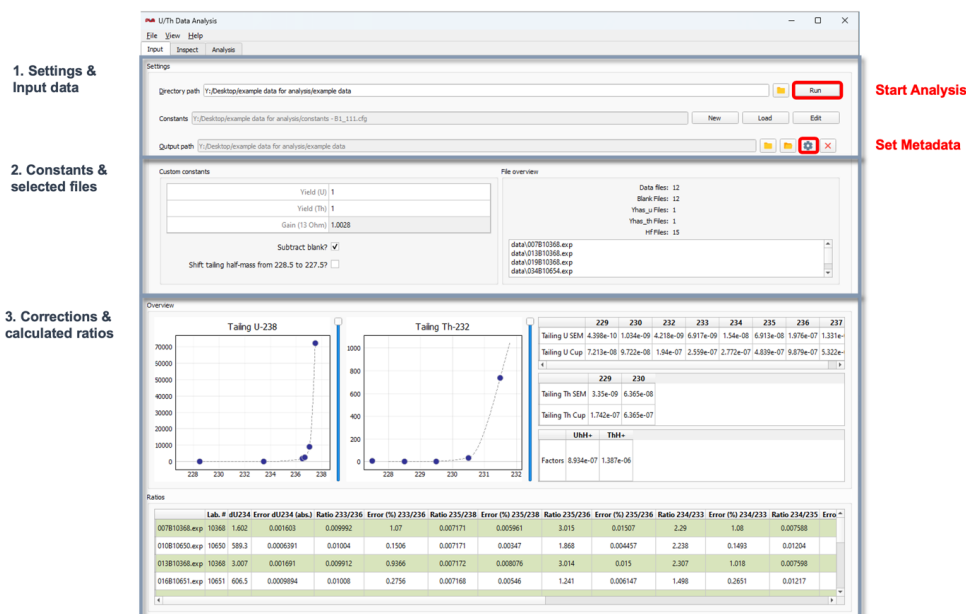
143 **3 Data treatment and analysis procedures**

144 The whole analysis procedure from raw data treatment to age calculation is conducted in one GUI featuring three
145 tabs: ‘Input’ for isotopic ratio calculations, ‘Inspect’ for outlier correction of the signal and ‘Analysis’ for age
146 calculation. The source code is accessible at https://github.com/EnvArchivesHD/UTh_Analysis. It is based on
147 the open source PyQt5 Python library (<https://pypi.org/project/PyQt5/>). The folder
148 https://github.com/EnvArchivesHD/UTh_Analysis/tree/main/dist contains the compiled .exe file for the GUI
149 (“UTh Data Analysis.exe”) as well as default configuration files (“constants – coral.cfg” and “constants –
150 stalag.cfg”). Input and output format of files are .csv or .xlsx. The GUI consists of three consecutive tabs, for which
151 the functionalities and the underlying calculations and processes will be described in the following.



152 **3.1 Input tab**

153 In ‘Input’, as presented in Figure 1, the user can navigate to the folder containing the raw mass spectrometer data
 154 and start the calculation of corrected isotopic ratios (Box 1 in Figure 1). All tab screenshots present data from
 155 stalagmite B1. Prior to the calculations, a configuration file containing all necessary constants used in the
 156 calculations needs to be loaded (same Box 1). This file contains constants and correction factors used for
 157 evaluation of the activity ratios and ages, such as mass fractionation coefficients, decay constants, the exact masses
 158 of the isotopes and the values applied for initial ²³⁰Th correction model. All constants can be edited manually
 159 either in the configuration file directly, or within the GUI using the button “edit”. An exemplary configuration
 160 table is also provided in the supplementary material (Figure S2). To apply a ²³⁰Th correction model a value can
 161 be set for the activity ratio and uncertainty of the contaminating material (“A230Th232Th Init. ”). The standard
 162 value would be the bulk earth mean activity ratio of 0.75 ± 0.38. Exemplary templates for corals and speleothems
 163 with conventionally used correction models are provided. For speleothems, a typical activity ratio of
 164 (²³⁰Th/²³²Th)_{ini/detr} of detritus is estimated to 0.75 ± 0.38, which is derived from a bulk earth Th/U weight ratio of
 165 4.1 ± 2.05 (Wedepohl, 1995) and the assumption of ²³⁰Th, ²³⁴U, and ²³⁸U being in secular equilibrium (Cheng et
 166 al., 2013). Nevertheless, this ratio may require adjustment according to local conditions. The coral template
 167 assumes as default value an activity ratio of 8 ± 4, which is estimated for corals dwelling in waters of the northeast
 168 Atlantic upper thermocline (Wefing et al., 2017). For one data series, only one correction constant, the
 169 (²³⁰Th/²³²Th) activity ratio of the contamination, can be added to the calculation. Hence, in case several factors
 170 need to be explored, the data series requires repeated treatment.



171
 172 **Figure 1.** Input tab: (1) In the top part the data folder is selected (“directory path”), the constants file (“constants”) can be
 173 loaded (“load”), edited (“edit”), or created (“new”). In addition, it is possible to set an “output path”. Red boxes show the
 174 “settings” button to enter metadata for saving, as well as the “run” button to start the analysis. Box (2) shows the custom
 175 constants box as well as the file overview for the selected folder. In box (3), the plots on the top left show the interpolated
 176 tailing. On the top right, numerical values of U and Th tailing and hydride correction are presented. The calculated ratios are
 177 shown in the bottom panel.



178 Figure 1 shows the layout of the GUI ‘Input’ tab. Once the constants are implemented and the input data are
179 selected, it is optional to choose an output path to store the analysis output (Box 1). If no path is specified, the
180 results will be stored in the raw data folder. When clicking the settings button next to the output path (highlighted
181 in red in Box 1), a menu opens in which the following parameters about the sample can be noted: denomination,
182 type of archive, lab numbers, geographic origin, and a general description. The first and last laboratory number
183 are automatically read out from the raw data. The final output result files will then be saved in a newly created
184 folder under the name $[_{labnumber_1-labnumber_n}]$ denomination in the directory chosen before. The metadata
185 information transferred through the GUI dialogue window is stored in a .json file in the respective folder. In the
186 ‘custom constants’ panel (Box 2), some settings can be selected, for example, if the blank has already been
187 subtracted in the mass spectrometric software or not. Next to this panel, an overview over the files read in from
188 the folder is shown. After running the evaluation script with the loaded data and adjusted settings (Button “run”,
189 highlighted in red in Box 1), the results of tailing and hydride correction, respectively as well as the calculated
190 ratios are displayed in the tab (Box 3). In addition, four excel .xlsx output files are created by the software at this
191 stage and stored in the directory path folder: Ratios.xlsx, Tailing.xlsx, PrBlank.xlsx and Intensities.xlsx.
192 Ratios.xlsx contains all calculated ratios and their errors as also presented in the GUI (10). Tailing.xlsx
193 summarizes the U and Th tailing values (in cps/V ^{238}U) for each mass. In PrBlank.xlsx, the average values for
194 each mass of the procedural blank measurements before each standard and sample are presented. Intensities.xlsx
195 contains the full data tables, with the signals in cps or V for each mass over all cycles. Every standard or sample
196 has its own sheet.
197 The algorithm of the ‘Input’ tab starts by reading in the '.exp' measurement files for sample and standard
198 measurements, process blank (=instrumental background) and Uranium and Thorium abundance sensitivity
199 measurements. The lines for all cycles for all isotopes are imported into a pandas data frame. Firstly, matrices for
200 tailing, hydride and process blank correction are produced that are later subtracted from the isotopic masses used
201 for ratio building. The individual steps are carried out as follows:

- 202 • Tailing: Uranium tailing is determined by measuring the off-masses 228.5, 233.5, 236.5, 236.7, 237.05
203 and 237.5 before a measurement sequence starts. The first half-mass can be changed between 228.5 and
204 227.5 as we observed a scatter peak around this mass that switched its exact position every few months.
205 Tailing off-masses are 227.5, 228.5, 229.5, 230.5 and 231.5. For interpolation to full masses, we use
206 piecewise cubic Hermite interpolating polynomial fits (Kerber et al. 2023). The masses that undergo ^{238}U
207 tailing correction are ^{233}U , ^{234}U , ^{235}U , ^{236}U , ^{229}Th , ^{230}Th and ^{232}Th , while ^{232}Th correction is applied to
208 ^{229}Th and ^{230}Th .
- 209 • Hydride isobaric interference: Hydride correction is determined by measuring 239 amu for UH^+ and 233
210 amu for ThH^+ during the abundance sensitivity measurements. The instrumental background (or
211 memory) is here referred to as process blank. It is measured between all sample and standard
212 measurements for 70 s. Typical blank levels afterwards are 0.5 cps for ^{230}Th and 6 cps for ^{234}U . The
213 matrices from these three corrections are then used for data reduction of each isotope.
- 214 • Detector setting: Three main different detector layouts are possible and are detected automatically by the
215 software: 1) all isotopes on cup, 2) ^{234}U , ^{230}Th and ^{229}Th on SEM and 3) ^{234}U on FC, ^{230}Th and ^{229}Th on
216 SEM. In normal operation, option 2) and 3) are used, depending on the ^{234}U concentration of the



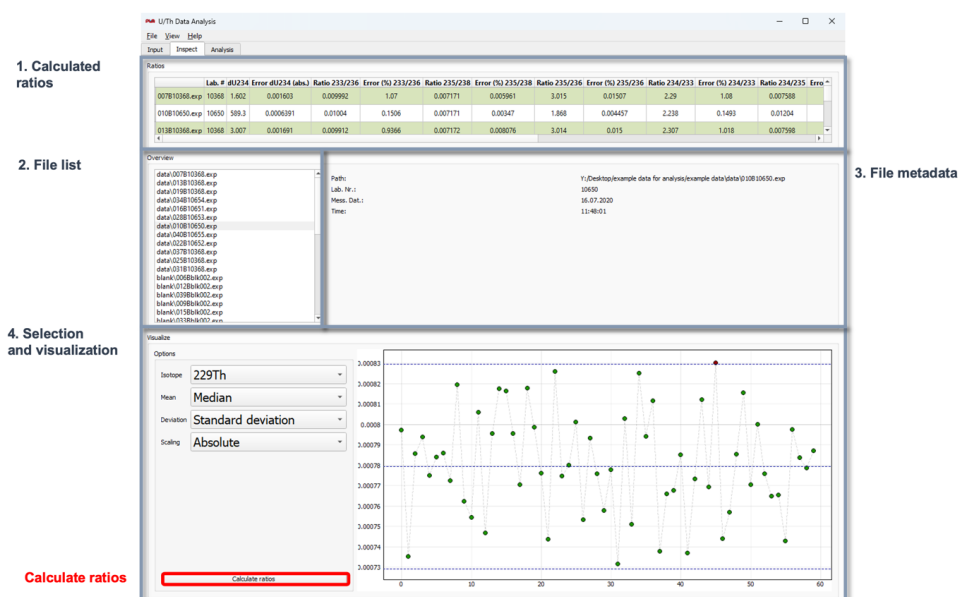
217 respective samples. ²³⁴U signals above 2 mV are measured on the centre FC which is the case for the
 218 absolute majority of samples.

219 The treated data are now used for the calculation of all relevant isotopic ratios followed by subsequent outlier
 220 tests, as described in Section 3.2.

221 3.2 Inspect Tab

222 Following the initial raw data treatment in the Input Tab, the ‘Inspect’ tab (presented in Figure 2) allows to
 223 visualize and retreat the data prior to final age calculation. In particular, the settings for the outlier test can be
 224 adapted.

225 The Inspect tab allows the user to plot the signal datapoints over the measurement cycle number for all isotopes
 226 in the individual measurement files of the sequence. In the top of the tab (1), the ratio results table from the ‘Input’
 227 tab is presented. On the left (2), the list of measurement files (.exp) is shown. By clicking on a specific file, the
 228 metadata and the signal plotted over the measurement cycle number are presented (3). On the bottom left (4), four
 229 dropdown menus are available: The first one, “Isotope”, allows to select one isotope from all of the isotope species
 230 measured. “Mean” offers to switch between mean and median of the signal. The “Deviation” menu provides three
 231 options for the assessment of data dispersion: standard deviation, median absolute deviation and interquartile
 232 range. By setting “Scaling” to absolute or relative, the y-axis of the plot on the right can be changed between
 233 signal intensities in V or cps and relative values. Any selection in the dropdown menus leads to an automatic
 234 update of the plot on the right. Mean resp. median, as well as the dispersion ranges are presented as blue dashed
 235 lines. Data points outside of the dispersion range are marked in red as outliers.



236

237 **Figure 2:** Inspect tab. (1) Ratio results table (from Input tab), (2) Overview of measurement files in folder, (3) metadata of a
 238 selected file and signal over measurement cycle number for one isotope (which can be selected in (4)), (4) option selection
 239 panel for the signal plotting.



240 The “Calculate ratios” button (5) provides the option to recalculate the ratios using the updated mean and deviation
241 selection for all isotopes. The default settings are median and standard deviation. However, these updated options
242 are then used to exclude outliers from the ratio arrays, not the signal intensity arrays themselves. This means that
243 not necessarily exactly the same data points are marked as outliers in the signal intensity plots and will be
244 excluded, but the ones where signal ratios of two isotopes are outside of the accepted deviation range. The option
245 selected in “Mean” will then also be used to calculate the average of the isotope ratios.

246 The method of calculating the uncertainty of outlier-corrected isotopic ratios, however, is fixed:

$$247 \quad \text{err}_{ratio} = 2 \frac{s}{\sqrt{n}} \quad (1)$$

248 with s being the standard deviation and n the number of data points (after outlier correction). The arithmetic mean
249 ($\bar{x} = \frac{x_1 + \dots + x_n}{n}$) and the median \bar{M} (central value of all values) are different ways of determining the average of a
250 distribution. The three different options for dispersion are defined as follows:

251 The standard deviation s is defined as

$$252 \quad s = \sqrt{\frac{1}{n-1} \sum_{i=1}^n (x_i - \bar{x})^2} \quad (2)$$

253 with \bar{x} being the mean. The interquartile range in turn is defined as the range containing the “middle” 50 % of
254 data points (Tukey, 1977). The median absolute deviation MAD is the median of absolute deviations from the
255 median, expressed as:

$$256 \quad MAD = k_i M_i(|x_i - \bar{M}|) \quad (3)$$

257 with M being the medians and x_i the original data (Leys et al., 2013; Huber, 2004; Rousseeuw and Croux, 1993).

258 For the calculation of the uncertainty MAD we assume normal distributed data, thus $k = 1.4286$.

259 3.3 Analysis tab

260 In a last step, age calculation is carried out in the ‘Analysis’ tab presented in Figure 3. Here, additional input data
261 is necessary from the sample weight tables (1). There are several ways to import these tables: Either by clicking
262 “Load” and navigating to the respective folder, or by manually creating the table directly in the GUI (“Create”).
263 An exemplary weight table is provided in the supplementary data (Figure 1). In the panel “Metadata history”, the
264 previously loaded sample weight tables in the directory path folder are shown, and can be directly imported (2).
265 “Start Analysis” calculates the ages (3). Outputs are both presented in the GUI (4) and stored in an Results.xlsx
266 file. In case an output path was specified, Results.xlsx is created both in the output and in the directory path folder.
267 If the output path is missing, the file is only saved in the directory path folder. If an output directory has been
268 created for specific lab numbers, all following analysis of these same files will be written to the same output
269 directory, but not overwrite earlier Results.xlsx. The Results.xlsx has five sheets: *Inputs*, *Calc*, *Results*, *Constants*
270 and *Options*. *Inputs* presents sample weight and metadata as well as the calculated ratios. In *Calc*, all steps of the
271 age calculation such as concentrations and activity ratios are shown. *Results* is a summary of the most important
272 calculation steps and final age values and the same table as is presented in the GUI as (4). *Constants* contains the
273 whole list of values from the (potentially edited) ‘.cfg’ file. In *Options* the average and dispersion measure option
274 are stored.



275 The equations for activity ratios to calculate ages are implemented according to Ivanovich and Harmon (1992),
 276 with:

$$277 \left(\frac{^{234}\text{U}}{^{238}\text{U}}\right) (t) = \left(\left(\frac{^{234}\text{U}}{^{238}\text{U}}\right)_{\text{init}} - 1\right) \cdot e^{-\lambda_{234}t} + 1 \quad (4)$$

$$278 \left(\frac{^{230}\text{Th}}{^{238}\text{U}}\right) = 1 - e^{-\lambda_{230}t} + \frac{\delta^{234}\text{U}}{1000} \cdot \left(\frac{\lambda_{230}}{\lambda_{230}-\lambda_{234}}\right) \cdot \left(1 - e^{-(\lambda_{230}-\lambda_{234})t}\right) \quad (5)$$

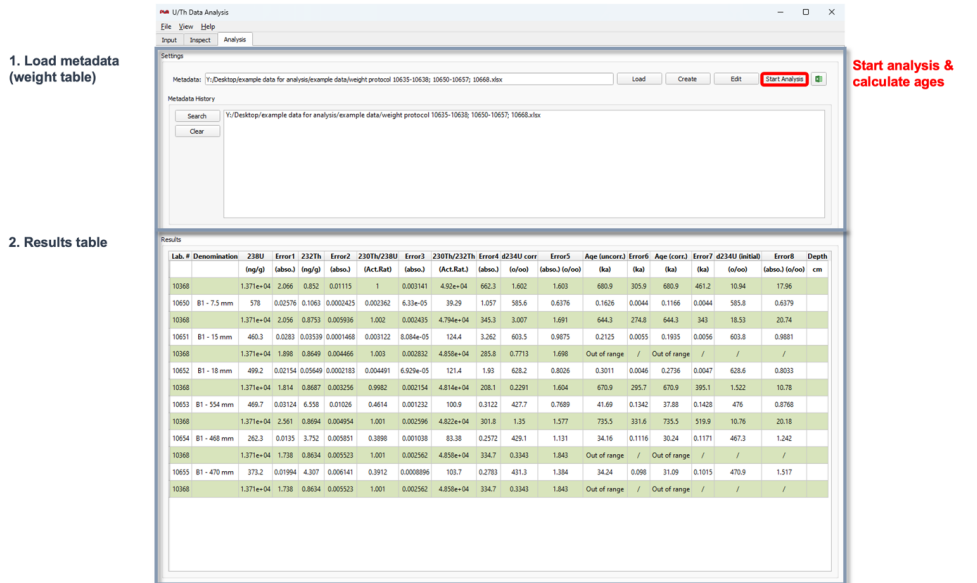
279 with

$$280 \delta^{234}\text{U} = \left(\left(\frac{^{234}\text{U}}{^{238}\text{U}}\right)_{\text{meas}} - 1\right) \cdot 1000 \text{ (‰)} \quad (6)$$

281 To obtain ages corrected for initial/detrital ^{230}Th , the $^{230}\text{Th}/^{238}\text{U}$ activity ratio used in eq. 5 is corrected using the
 282 initial $(^{230}\text{Th}/^{232}\text{Th})_{\text{ini/detr}}$ ratio and

$$283 \left(\frac{^{230}\text{Th}}{^{238}\text{U}}\right)_{\text{corr}} = \left(\frac{^{230}\text{Th}}{^{238}\text{U}}\right)_{\text{meas}} - \left(\frac{^{232}\text{Th}}{^{238}\text{U}}\right)_{\text{meas}} \cdot \left(\frac{^{230}\text{Th}}{^{232}\text{Th}}\right)_{\text{ini/detr}} \cdot \left(\frac{\lambda_{230}}{\lambda_{230}-\lambda_{234}}\right) \cdot e^{-\lambda_{230}t} \quad (7)$$

284 These equations need to be solved numerically. For the determination of age uncertainty, the usual approach is to
 285 repeat the numerical determination of the age for several thousand runs in a Monte-Carlo simulation while random
 286 sampling the input ratios from a normal distribution with μ corresponding to the ratio's value and σ corresponding
 287 to the uncertainty on this parameter.

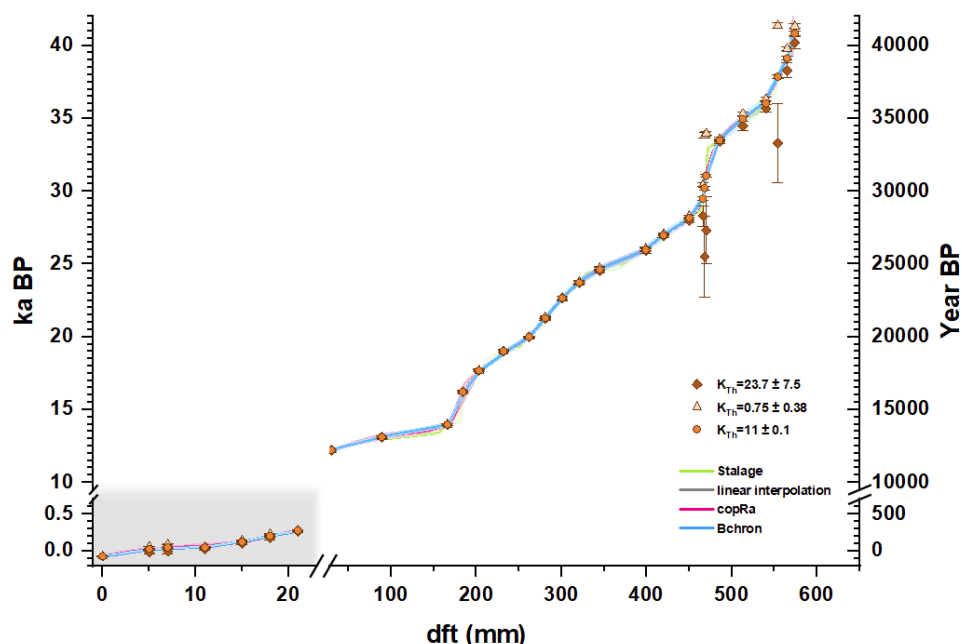


288 **Figure 3:** Analysis tab. (1) Load sample weight tables (metadata files). The bottom panel lists the history of previously loaded
 289 tables. The button highlighted in red starts the analysis to calculate ages ("Start Analysis") button. The panel in box (2)
 290 displays the results table.
 291



292 **4 Example dataset: Stalagmite B1**

293 To demonstrate our data evaluation tool, we here present newly obtained ages of stalagmite B1 from Larga Cave,
 294 Puerto Rico. The results of activity ratios and calculated ages can be accessed in the supplementary table S1.
 295 Analysis of the speleothem samples reveals moderate U concentrations in the range between 300 and 600 ng/g,
 296 and minor detrital ²³²Th contamination with (²³⁰Th/²³²Th) activity ratios of typically >300. However, in both the
 297 top 20 mm and around 450 mm dft lower (²³⁰Th/²³²Th) activity ratios of c. 40 – 125 are measured. U isotopic
 298 composition varies between 450 and 640 ‰ of δ²³⁴U values. Uncertainties of the uncorrected ages are typically
 299 in the range of 0.2 to 0.6 ‰ (Table S1). Drip water shows high U concentration of 0.825 ng/g and elevated initial
 300 Th concentrations, with an activity ratio of K=(²³⁰Th/²³²Th)=11.1 ± 0.1. We have used the software to test how
 301 the chronology changes to assess the influence of a varying initial Th activity ratio. For this, we used three different
 302 correction models, including the measured initial Th ratio of the drip water (K=11.1 ± 0.1), the detrital correction
 303 value of K=0.75 ± 0.38 derived from the bulk Earth crust chemical composition, as well as a value of K=23.7 ±
 304 7.5 as previously determined using isochrons on speleothem PR-LA-1 from the same cave (Warken et al. 2020).
 305 Figure 4 shows the ages corrected for initial ²³⁰Th using the three different models. Only the initial ²³⁰Th value
 306 measured in the drip water yields a stratigraphic order of the corrected ages supporting the use of this value.
 307 Residual variability around the mean chronology increases and age inversions appear in the record when using a
 308 different value of K.

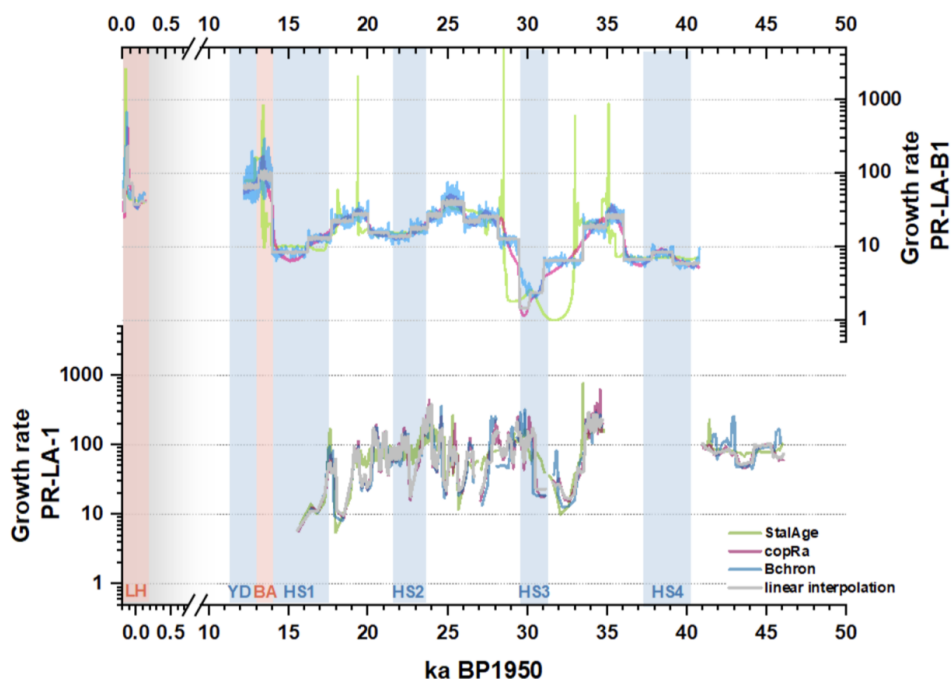


309

310 **Figure 4:** ²³⁰Th/U ages and different age-depth simulations for stalagmite B1 using linear interpolation, as well as the
 311 algorithms Stalage, copRa and bchron linear interpolation, as well as the algorithms StalAge (Scholz and Hoffmann, 2011),
 312 CopRa (Breitenbach et al., 2012) and Bchron (Haslett and Parnell, 2008). Note that the axes are split at the position of the
 313 growth stop at 23mm dft to visualize the age-depth relationship also during the short growth phase during the latest Holocene
 314 after 0.3 ka BP.



315 A Rosholt - Isochron determined for the section between 466 and 470 mm dft similar to the approach of Warken
 316 et al. (2020) further supports the chosen correction value (supplementary Figure S3). This result highlights the
 317 relevance to obtain such measures for better age correction either by studying the drip water ^{230}Th and ^{232}Th
 318 isotope composition and using isochron approaches. The resulting corrected ages suggest a speleothem growth
 319 between 0.060 ± 0.013 ka BP₁₉₅₀ and 40.81 ± 0.16 ka with a hiatus between 12.22 ± 0.043 and 0.277 ± 0.008 ka
 320 BP at 23mm dft. Hence, speleothem B1 extends the existing speleothem record from Larga Cave into the
 321 deglaciation covering Heinrich Stadial 1 (HS1), the so called Bølling/Allerød warming (BA), as well as the
 322 beginning of the Younger Dryas cold reversal (YD).
 323 Figure 6 shows the growth models obtained by linear interpolation, as well as the algorithms StalAge (Scholz and
 324 Hoffmann, 2011), CopRa (Breitenbach et al., 2012) and Bchron (Haslett and Parnell, 2008) as implemented by
 325 Roesch and Rehfeld (2020) (code accessed at <https://github.com/paleovar/SISAL.AM>, codes licensed by the right
 326 holder(s) under a GPL-3). Growth rates of stalagmite PR-LA-B1 vary between c. 10 and 150 $\mu\text{m/a}$, with highest
 327 values during the warm Bølling–Allerød period c. 13.97 ± 0.051 and 13.114 ± 0.073 ka BP as well as the late
 328 Holocene growth phase after 0.277 ± 0.008 ka BP. Lowest growth rates occur during the final stage of Heinrich
 329 Stadial (HS) 1 (16.23 ± 0.082 to 13.97 ± 0.051 ka BP), HS3 (31.02 ± 0.10 to 29.38 ± 0.12 ka BP), and HS4 (40.81
 330 ± 0.16 to 39.12 ± 0.12 ka BP).



331
 332 **Figure 5:** Growth rates of speleothems B1 (top panel, this study) and PR-LA-1 (Warken et al., 2020). Vertical red (blue) bars
 333 indicate the timing of warm (cold) phases in Puerto Rico, including the growth phase of B1 during the latest Holocene (0.3 ka
 334 BP to present), the Younger Dryas (YD), Bølling/Allerød warming (BA), and Heinrich stadials (HS) 1 to 4.

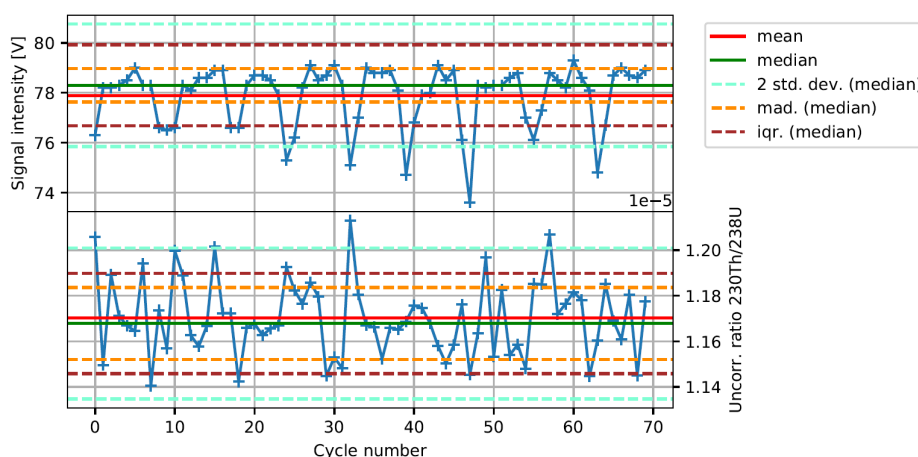


335 **5 Discussion**

336 **5.1 Outlier correction**

337 Outlier correction is carried out automatically by the software adapting the dispersion measure of the raw data
 338 and in the following we argue that generally means should be replaced by medians. Shao et al. (2019) had
 339 addressed this problem by implementing manual outlier removal by comparison to boxplots based on interquartile
 340 ranges. We opted for the automatic version as this is more time efficient for large datasets. The different dispersion
 341 measure options described in Section 3.2 are relevant because measurements are not always ideal cases with
 342 normally distributed data and thus outliers. During measurements, short-term system instabilities occur for a
 343 variety of reasons, such as varying gas flow in the inlet system, plasma instabilities, and varying size of sample
 344 aerosols causing outliers in the signal intensities. Even though only the ratios between the different isotopes are
 345 of interest, strong changes in signal intensity may lead to varying isotope ratios, as a result of changing variance.
 346 Such difference may be amplified by the use of different detectors or with respect to different magnetic field
 347 settings, which are not necessarily responding at exactly same amplitude. Moreover, signal decreases (detuning
 348 events and temporal clocking) cause the statistical variance to increase locally.

349 Figure 6 shows an example: The upper panel displays periodic dips in the ^{238}U signal intensity during a
 350 measurement. In the lower panel of Fig. 4, the uncorrected ($^{230}\text{Th}/^{238}\text{U}$) activity ratio for the same measurement is
 351 plotted. For both curves, the different measures to calculate dispersion are shown. The default method (2 standard
 352 deviation) does not remove all the systematic outliers. Also, it is clearly visible that the median agrees much better
 353 with the majority of signal intensity values than the mean, which is much stronger influenced by the periodic dips
 354 due to the asymmetry in the statistical distribution. Such an obvious difference is not visible in the isotope ratio,
 355 but within the resulting uncertainty. Consequently, we propose to generally use the median instead of mean by
 356 default. This is more accurate in the case of asymmetric small-scale oscillations inside the non-outlier interval and
 357 has no disadvantages.



358

359 **Figure 6:** Upper panel: ^{238}U signal intensities in Volt over measurement cycles for a carbonate sample during routine lab
 360 measurements. Lower panel: Corresponding uncorrected $^{230}\text{Th}/^{238}\text{U}$ ratio. Mean and median as well as the three different
 361 dispersion measures are plotted.



362 Applying standard deviation as a dispersion measure in Figure 6 does not cover most of these outliers due to their
363 large number and relatively small deviation. Thus, applying another dispersion measure for outlier removal is
364 necessary here, and in addition more robust and easier to accomplish than manual deletion of all of the outliers. It
365 is important, however, to stress that the outlier correction using the selected dispersion option is run on the
366 calculated ratios after correction, not on the signal intensities themselves. This implies that when all isotopes are
367 affected in the same way, they pass the outlier test. This is, however, unlikely at least for ratios of isotopes
368 measured in different magnetic field settings. The dispersion measure of the outlier corrected ratio array is the
369 same in every case, as described by Equation 1.

370 5.2 Detrital thorium correction

371 Thorium correction is often crucial for studying carbonates where the correction is significant, but the initial ^{230}Th
372 value is unknown, potentially variable, or when studying “dirty” carbonates such as tufa and travertine (Mallick
373 and Frank, 2002; Hellstrom, 2006; Wenz et al., 2016). Several studies have shown that this correction particularly
374 important for speleothem records from the Caribbean and Central American region, where values were found
375 including 2 ± 1 (Schorndorf et al., 2023) or 14 ± 4 (Moseley et al., 2015). In Larga Cave, initial ($^{230}\text{Th}/^{232}\text{Th}$)
376 ratios are presumably even higher (Vieten et al., 2024; Warken et al., 2020). Besides the terrestrial regime, this
377 aspect is also relevant for marine archive such as corals, where studies propose a large range of seawater
378 ($^{230}\text{Th}/^{232}\text{Th}$) activity ratios. While Cheng et al. (2000a) set the range to 80 ± 80 for deep-sea solitary corals, and
379 Frank et al. (2004) calculated 10 ± 4 from seawater in the Eastern North Atlantic deep sea, values between 0.4 –
380 3.1 were determined for tropical corals (Shen et al., 2008). The range of both absolute values and uncertainties
381 for these widely studied archives is hence enormous, and the choice of the appropriate correction model becomes
382 particularly important, when (i) samples are very young and have generated only small amounts of ^{230}Th from U-
383 decay, or (ii) when ultra-high precision is at play since any possible correction of the data contribute to the final
384 age uncertainty. In our case study, we have run the correction of the ages of stalagmite B1 using three different
385 correction models (Table S1). The resulting differences are visualized in Figure 4, and demonstrate the significant
386 impact not only on absolute corrected ages, but also their uncertainties. For the young age at 7 mm dft
387 (0.0466 ± 0.0045 ka BP), the difference in the absolute corrected age when using another correction factor than the
388 drip water value of $K = 11.1 \pm 0.1$ is c. ± 50 years, which corresponds to a relative difference in the order of 100%
389 (compare Table S1). Another example is the sample at 554 mm dft (37.81 ± 0.14 ka BP for $K = 11.1 \pm 0.1$), for which
390 the other correction models also lead to substantially different ages of 41.37 ± 0.19 ka BP ($K = 0.75 \pm 0.38$) and
391 33.3 ± 2.7 ka BP ($K = 23.7 \pm 7.5$), hence the differences are still in the range of c. 10%. Notably, the low relative
392 error of the initial ($^{230}\text{Th}/^{232}\text{Th}$) activity ratio of the drip water results in equally low uncertainties of the corrected
393 age in the range of 0.4%. In contrast, the relative uncertainty of the age corrected with $K = 23.7 \pm 7.5$ increases to
394 8%. Our GUI permits an easy adjustment of the initial ($^{230}\text{Th}/^{232}\text{Th}$) activity ratio for Th correction, which allows
395 a direct assessment of the resulting corrected ages and uncertainties, and provides thus a convenient basis for
396 further comparisons of the data. The use of a standardized software instead of handmade tuning reduces the
397 susceptibility to potential errors, e.g., from copy-pasting, and ensures reproducibility in case a re-evaluation of
398 the data is required to a later stage.



399 **5.3 In-cave comparison of speleothem growth rates**

400 The high number and precision of $^{230}\text{Th}/\text{U}$ ages of speleothem B1 allows investigation of growth rates changes.
401 Comparison with northern hemispheric climatic changes suggests, that speleothem B1 growth is sensitive to
402 prominent millennial-scale temperature variability, with higher growth rate during warmer phases and vice versa.
403 In particular, during the cooler and drier Heinrich stadials (Warken et al., 2022b), growth rates are reduced.
404 In addition, the results allow a comparison of the two coeval stalagmites from Larga Cave as shown in Figure 5.
405 Overall, GRs of PR-LA-B1 are about 5 times lower than observed for PR-LA-1, where average annual growth
406 rates are up to several mm/a. The difference in mean GR is also reflected in the shape of both speleothems, with
407 PR-LA-1 exhibiting a large and variable diameter between c. 15 and 35 cm (Warken et al., 2020), while B1 is
408 thinner with a diameter of 10-15 cm (supplementary Figure S1B). Differences in speleothem growth rates and the
409 shape of a stalagmite may result from temperature, carbonate saturation, drip rate, and carbon dioxide contrast
410 between cave air and saturation concentration of drip water (Merz et al., 2022; Skiba and Fohlmeister,
411 2023; Kaufmann, 2003; Dreybrodt, 1999). Ca concentrations in Larga Cave show no significant differences
412 between drip sites (Vieten et al., 2018a, Vieten et al., 2018b, Warken et al., 2022), and a single Ca concentration
413 measurement at site B1 is also within the same range (Vieten et al., 2018a, Vieten et al., 2018b).
414 Therefore, the amplified GR and generally larger diameter of PR-LA-1 could be the result of the considerably
415 lower pCO_2 values in the main passage (600 and 1800 ppm) than compared to the back part of the cave (2300
416 and 3600 ppm), which facilitates enhanced oversaturation of the drip water with respect to calcite, and hence,
417 stronger degassing of CO_2 and speleothem growth (Merz et al., 2022). In addition, variations in drip rates influence
418 the GR. Spot observations suggest a multi-annual variability of drip rates at the location of stalagmite PR-LA-B1.
419 For PR-LA-1, no modern drip site is available, which precludes direct comparison of the role of the drip rate. The
420 factor five amplified GR of specimen PR-LA-1, however, gives rise to presume a higher drip rate in addition to
421 lower atmospheric cave air pCO_2 values, in particular during centennial increases of the GR to values as high as
422 mm/a at this site. Further note that the timing of the GR increases above the mean GR or paused growth did not
423 occur synchronously, which precludes a common mechanism stimulating the GR. Hence, the two stalagmites
424 reveal growth differences potentially related to ventilation conditions.
425 Other differences of the two sites are visible in both speleothems geochemistry, which, however, cannot be directly
426 related to drip rate or cave air pCO_2 concentration. The Uranium concentration [U] of PR-LA-B1 is systematically
427 higher than for the one of PR-LA-1 (c. 90 – 400 ng/g) and the initial $\delta^{234}\text{U}$ is with values ranging between 450 to
428 640 ‰ strongly elevated compared to the $\delta^{234}\text{U}$ from PR-LA-1, which varies between values of c. 70 - 100 ‰.
429 This demonstrates a reduced flux of excess ^{234}U from the host rock at the drip site of PR-LA-1, potentially resulting
430 from varying release of ^{234}U through alpha-recoil of the decay of ^{238}U at the two sites. A likely explanation may
431 be the difference in local host rock overburden of PR-LA-B1 with c. 40-60 m to PR-LA-1 with c. 20-40 m, and
432 thus moderately longer residence times of the karst water at site B1. Consequently, given the sum of observations
433 it seems most likely that the GR of PR-LA-1 in the better ventilated region with less rock overburden responds to
434 drip rate more sensitively than PR-LA-B1, which in contrast seems more sensitive to cave ventilation, i.e., cave
435 air pCO_2 .



436 **6 Conclusion**

437 We here provide an algorithm combined with a user-friendly GUI application for $^{230}\text{Th}/\text{U}$ MC-ICP-MS data
438 treatment and age calculation. The two so far published programs explicitly aimed at $^{230}\text{Th}/\text{U}$ dating data reduction
439 and age calculation are both written for ThermoFisher Neptune instruments as well. Pourmand et al. (2014)
440 described a Mathematica routine, distributed as a Computable Document Format (.cdf) file, while Shao et al.
441 (2019) had published a Matlab algorithm with GUI. We here have chosen to use Python for our algorithm and
442 GUI to keep it open-source. The advanced user might want to change settings, which makes an opensource
443 language and libraries a major advantage. However, the stand-alone executable .exe format of the GUI allows
444 user-friendly handling also for non-programming experts. Our program supports multiple types of detector
445 configurations: the FC-FC based approach as well as FC-SEM combining protocols. It is however adapted for
446 combined Th and U measurements in three magnetic field lines (compare Kerber et al. (2023)), but other methods
447 (such as separate solutions for Th and U) can be implemented with small changes in the code. Furthermore, we
448 offer the first order Taylor derivation as a time-saving option for uncertainty calculation of final ages. Our
449 application is especially designed to take reproducible and clear data management into account by a collection of
450 methods: This includes that automatic creation of folders containing the results files and information on the sample
451 metadata is possible and that .xlsx output files automatically contain all constants used for calculation, as well as
452 the settings for outlier correction. Manually changing input constants, e. g. correction, of initial/detrital Th does
453 not require to go to the code directly. So, the whole analysis scheme does not require any copy-and-pasting from
454 one excel table to the other, and the constants used for calculation are easy to update.
455 Lastly, we demonstrated our protocols and data analysis scheme by accurately measuring and evaluating 30
456 speleothem ages from Larga Cave, Puerto Rico. Analyses of the growth rates and comparison with a coevally
457 growing stalagmite from the same cave highlights the importance of in-cave processes for speleothem deposition
458 rates.

459 **Author contributions**

460 IK - conceptualized the work, created and tested the implementation and operation of the code, co - supervised
461 FK, who developed the code for the GUI and tested rigorously all corrections. NF - conceptualized the project,
462 supervised IK, and FK and quality controlled the Th U isotope measurements of PR-LA-B1. SW - conceptualized
463 the project, provided guidance on sample selection, verified the code and conceptualized the application. SW
464 further evaluated the resulting age data on PR-LA-B1 and supervised a student project during which these and
465 other data had been collected.

466 **Code availability**

467 The source code of “UTh Data Analysis” is accessible at https://github.com/EnvArchivesHD/UTh_Analysis .
468 The folder https://github.com/EnvArchivesHD/UTh_Analysis/tree/main/dist contains the compiled .exe file for
469 the GUI (“UTh Data Analysis.exe”) as well as default configuration files (“constants – coral.cfg” and “constants
470 – stalag.cfg”).



471 **Data availability**

472 Results of speleothem B1 ²³⁰Th/U dating are available in the online supplementary material.

473 **Sample availability**

474 Sample material is available on request to swarken@iup.uni-heidelberg.de

475 **Competing interests**

476 At least one of the (co-)authors is a member of the editorial board of Geochronology.

477 **Disclaimer**

478

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