



## Evidence for preindustrial variations in the marine surface water carbonate system from coralline sponges

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[1] Coralline sponge skeletons are excellent tools for reconstructing the carbon isotope history of dissolved inorganic carbon (DIC) in tropical surface waters. Carbon isotope records from coralline sponges clearly reflect the industrial  $^{12}\text{C}$  increase in atmospheric  $\text{CO}_2$  with a precision that permits quantitative interpretations. We find from a set of  $\delta^{13}\text{C}$  records of four Caribbean sponge specimens that the isotopic response of surface water DIC to the changing isotopic composition of atmospheric  $\text{CO}_2$  varied dynamically during the last century, depending on the rate of atmospheric change. Three of our sponges provide 600 year long  $\delta^{13}\text{C}$  records. For the first time, we can reconstruct surface water  $\delta^{13}\text{C}_{\text{DIC}}$  for the full history of the industrial  $\text{CO}_2$  release as well as the preceding preindustrial period back to the beginning of the Little Ice Age. This provides a well-founded estimate of the anthropogenically uninfluenced, preindustrial background level of surface water  $^{13}\text{C}/^{12}\text{C}$  ratios. Our records show small but systematic variations that appear to be linked to the climate fluctuations of the Little Ice Age.

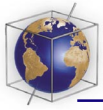
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## 1. Introduction

[2] We investigate stable carbon isotope records of aragonitic skeletons of the coralline sponge *Ceratoporella nicholsoni* (Figure 1), which lives in surface and upper thermocline waters of the Caribbean. These sponges precipitate aragonite very close to isotopic equilibrium with ambient sea water [Druffel and Benavides, 1986; Böhm *et al.*, 1996; Reitner, 1992]. Carbon isotope histories recorded by the sponges display the anthropogenic carbon isotope shift of the 19th and 20th century [Druffel and Benavides, 1986; Böhm *et al.*, 1996; Swart *et al.*, 1998; Lazareth *et al.*, 2000; Moore *et al.*, 2000; Nozaki *et al.*, 1978] as well as variations prior to the 19th century. These preindustrial variations provide evidence for natural variations in the atmosphere/surface ocean CO<sub>2</sub> inventory.

[3] On the basis of ice core analyses [Etheridge *et al.*, 1996], it has been shown that the atmospheric CO<sub>2</sub> mixing ratio ( $p\text{CO}_2$ ) varied by 6 ppm before the onset of industrial influences. Carbon isotope data of these ice core inclusions vary in concert with  $p\text{CO}_2$  [Francey *et al.*, 1999]. These variations have been linked to climatic variations known as “Medieval Warm Period” and “Little Ice Age.” Combined with the ice core record, our data provide further constraints on the modeling of global climate changes during this period and support the notion that the Little Ice Age had a measurable, global impact on the carbon cycle.

## 2. Material and Setting

[4] We collected living sponges (Figure 1) in 1993 and 1996 at Jamaica (Rio Bueno and Montego Bay) in reef caves in ~20 m of water and at the northwestern slope of Pedro Bank in 125 m of water (78°57'W 17°32'N). The sites are well exposed to the open sea. The Jamaican sites are situated within the mixed layer of the “Caribbean Surface Water,” while the Pedro Bank site is positioned below a barrier layer [Sprintall and Tomczak, 1992] near the top of the “Subtropical Underwater” [Morrison and Nowlin, 1982].

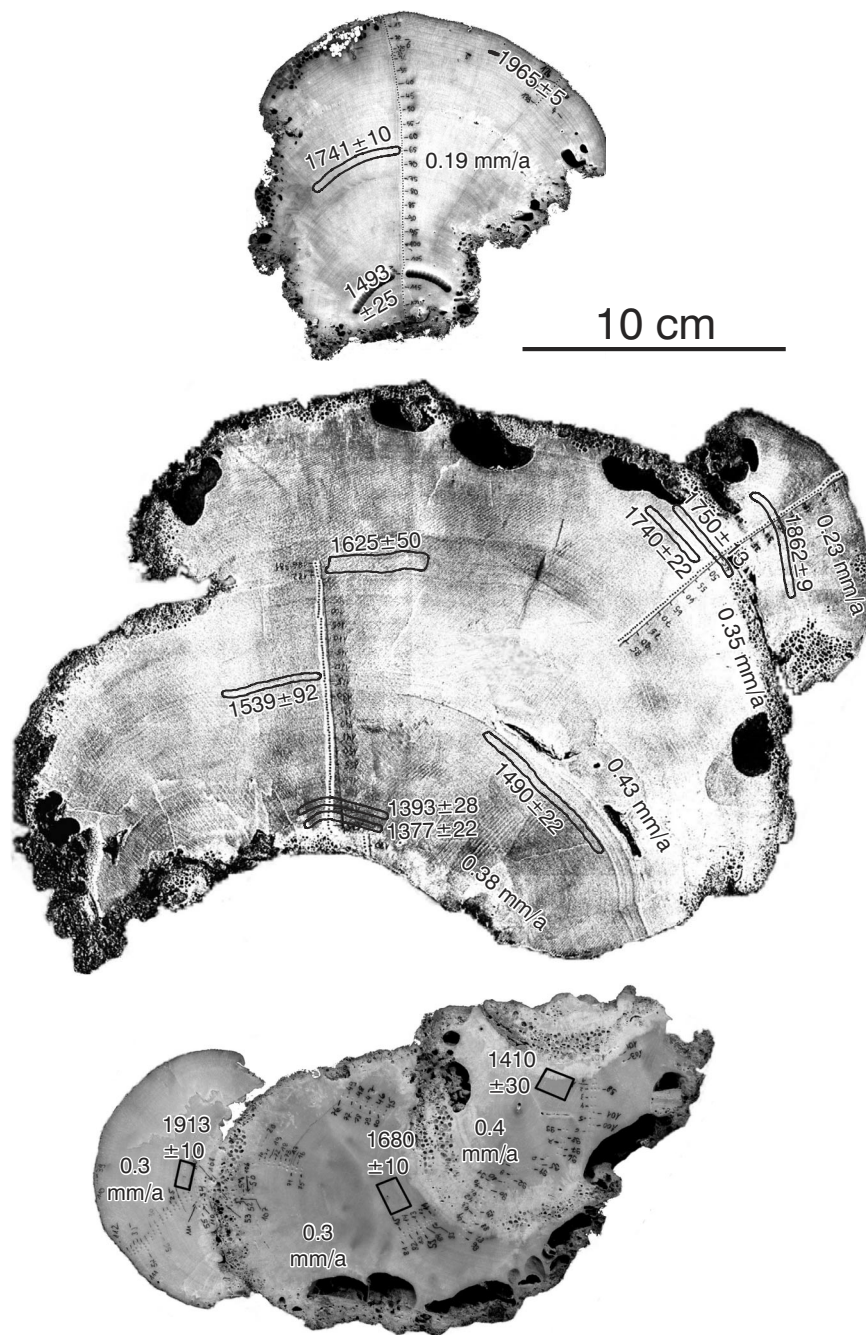
Whereas the Caribbean Surface Water is mainly derived from the tropical South Atlantic, the Subtropical Underwater forms in the tropical and subtropical North Atlantic from where it is transported southwest into the Caribbean [Schmitz and Richardson, 1991]. Thus the shallow water sponges formed their skeletons in waters that predominantly equilibrated with atmospheric CO<sub>2</sub> southeast of the Caribbean, while the deeper water sponges lived in waters that mainly reflect the atmosphere-surface water equilibration in the southwest flowing part of the North Atlantic subtropical gyre [Schmitz and McCartney, 1993].

[5] As the deeper water site is positioned near the transition from Subtropical Underwater to Caribbean Surface Water, a varying degree of mixing between the two water masses can be expected at this water depth. Upwelling of Subtropical Underwater in the southern parts of the Caribbean [Gordon, 1967] leads to further mixing of the two water masses. It should also be noted that the surface water stratification pattern of the Caribbean may have been different during the Little Ice Age.

## 3. Methods

[6] Microsamples for isotope analysis were collected with a 0.5 mm dental drill and processed using standard procedures as described by Böhm *et al.* [1996]. Analyses were carried out at the isotope lab of the Geological Institute, University of Erlangen, with a Finnigan Mat 252. Temporal resolutions of individual samples range from 2 to 4 years/sample, depending on the growth rates varying from 0.19 to 0.43 mm/year. External precision ( $2\sigma$ ) of  $\delta^{13}\text{C}$  measurements based on multiple analyses ( $N = 87$ ) of standards NBS 19 and IAEA CO1 is  $\pm 0.04\%$ . All carbon isotope ratios are given in the common delta notation, relative to V-PDB (Vienna Pee Dee Belemnite).

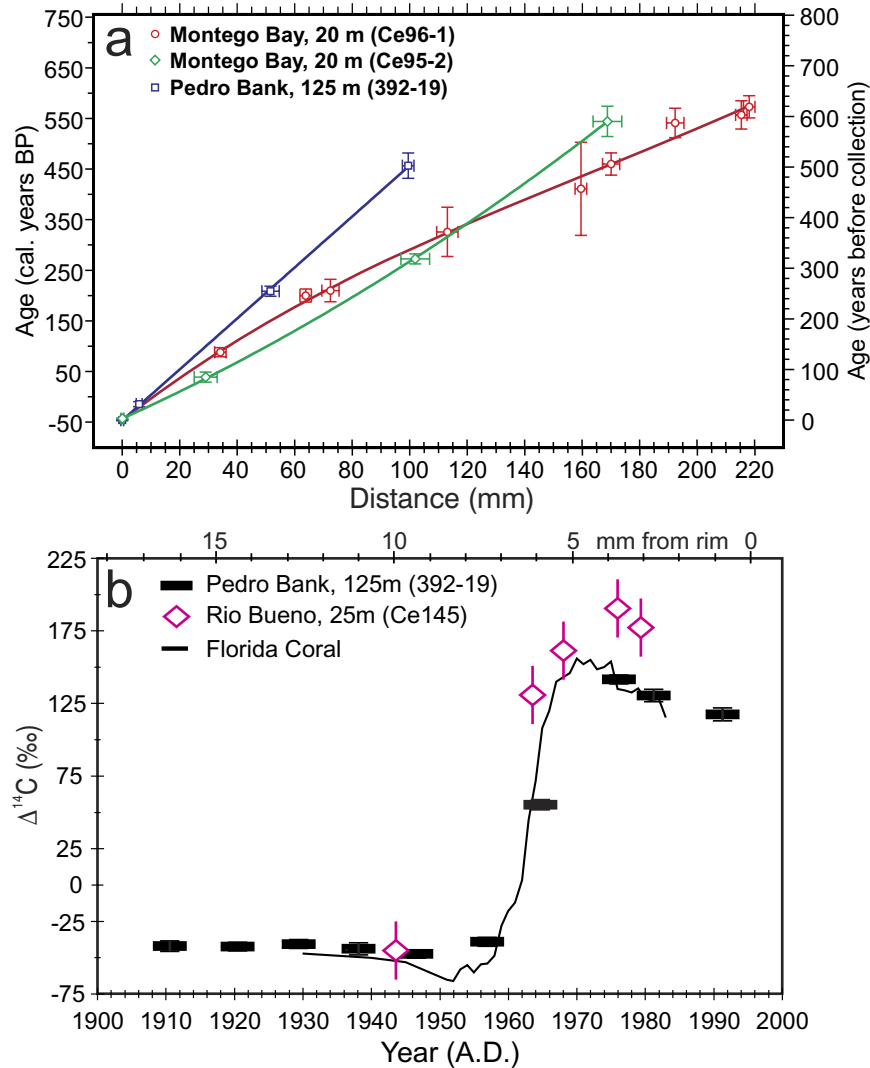
[7] Growth rates derive from U/Th dates in three specimens (measured at GEOMAR and at Göttingen University) and fits of  $^{14}\text{C}$  AMS data (Figure 2) to the radiocarbon “bomb spike” [Benavides and



**Figure 1.** Polished slabs of the investigated *Ceratoporella nicholsoni* specimens (from top 392-19, Ce96-1 and Ce95-2) with sample transects (dotted lines). Uranium/Thorium ages are given as years of formation (A.D.) of the respective layers together with standard errors ( $2\sigma$ ). The 1965 datum of the top specimen is the radiocarbon bomb spike. The sampling spots of the U/Th datings and the radiocarbon sample that was assigned to the year 1965 A.D. are marked with bold black lines. Average growth rates are calculated from U/Th ages and the age of the surface, i.e., the years of collection (1996 A.D. for Ce96-1 and 392-19 and 1995 A.D. for Ce95-2).

Druffel, 1986; Druffel, 1989] in specimen Ce145 [Böhm et al., 1996] and specimen 392-19 (measured at the Leibniz-Labor, Kiel University). For the latter we assume that the timing of the radiocarbon

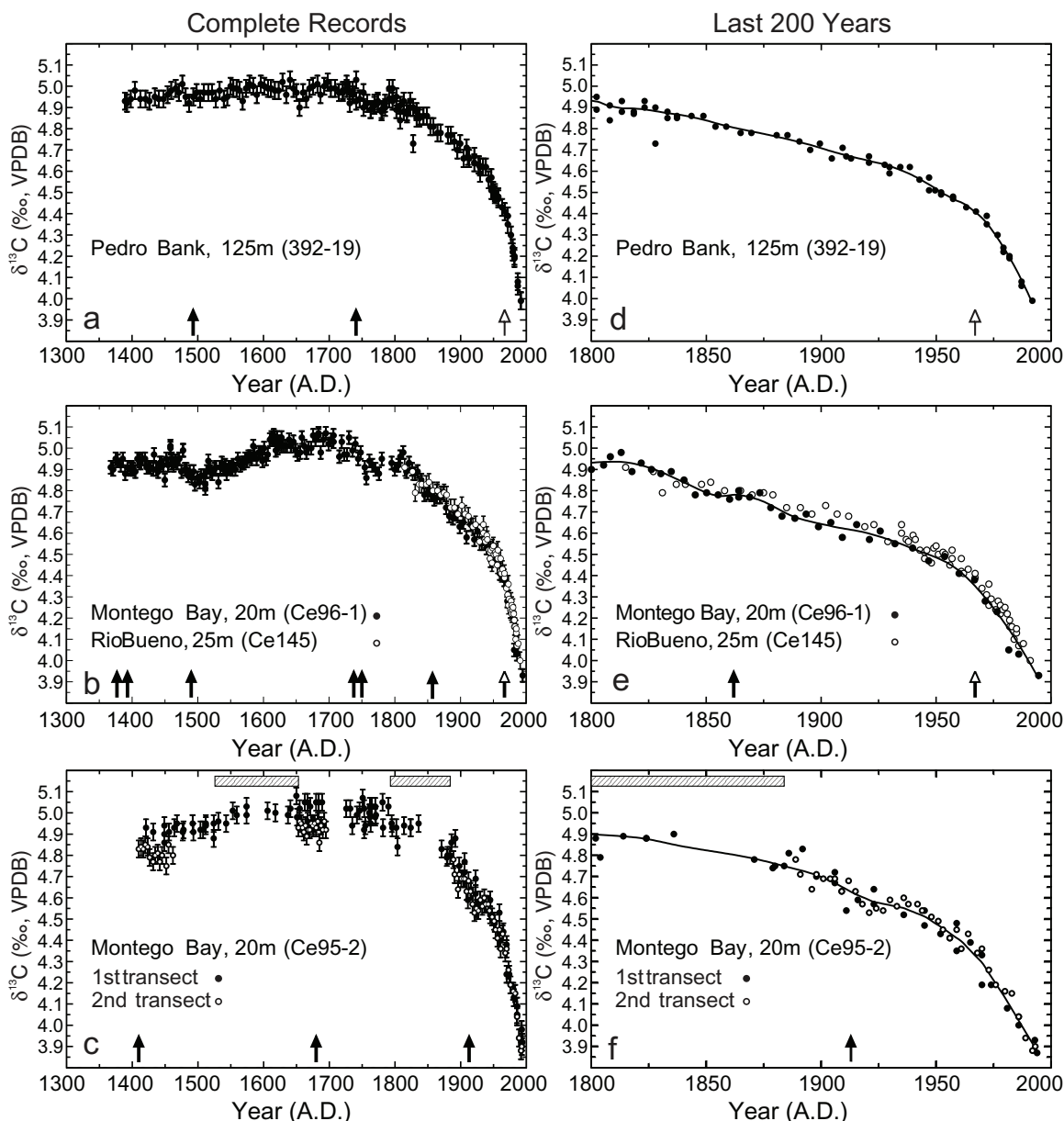
increase at 125 m depth was similar to the timing in surface waters of that region because the amplitude of change in  $\Delta^{14}\text{C}$  is similar to the surface water change (Figure 2).



**Figure 2.** (a) Age depth curves of the three investigated sponge specimens. Depth is measured along the isotope sample transects. The data points at 0 mm represent the top surfaces of the sponges. Horizontal error bars represent the sample width (measured parallel to growth axis). BP, before present (i.e. before 1950 A.D.). The data points are fitted by polynomials. (b) Fit of the measured radiocarbon values of specimens 392-19 and Ce145 to the coral  $\Delta^{14}\text{C}$  curve for Atlantic Ocean surface waters east of southern Florida [Druffel, 1989]. Error bars are smaller than symbols except where shown. The width of the symbols is equivalent to the width of the samples. The lower x axis corresponds to the Florida coral  $\Delta^{14}\text{C}$  curve. The upper x axis shows the sample distance from the top of the sponge skeleton along the growth axis. We take the value intermediate between the prebomb and postbomb levels to represent the year  $1964 \pm 5$  A.D. in both sponge skeletons. Spacing of the sponge radiocarbon samples is scaled to the lower x axis (time) by assuming a growth rate of 0.19 mm/a. Specimen Ce145 probably grew slightly faster (0.22 mm/a [Böhm *et al.*, 1996]) than specimen 392-19.

[8] On the basis of these data, we rely on steady growth rates to estimate the ages of individual sampled skeleton layers. Constant growth seems to be the rule, as growth rates calculated from U/Th dated sections of a single specimen show little variation, as long as homogeneous parts of a skeleton are considered (Figure 1). We use linear

regression to fit growth curves to the radiometrically dated tie points (Figure 2). A linear growth model provides an optimal fit to the age data of specimen 392-19. We used a second-order polynomial for specimen Ce95-2 and a third order polynomial for specimen Ce96-1. Details of the dating methods will be given elsewhere (A.

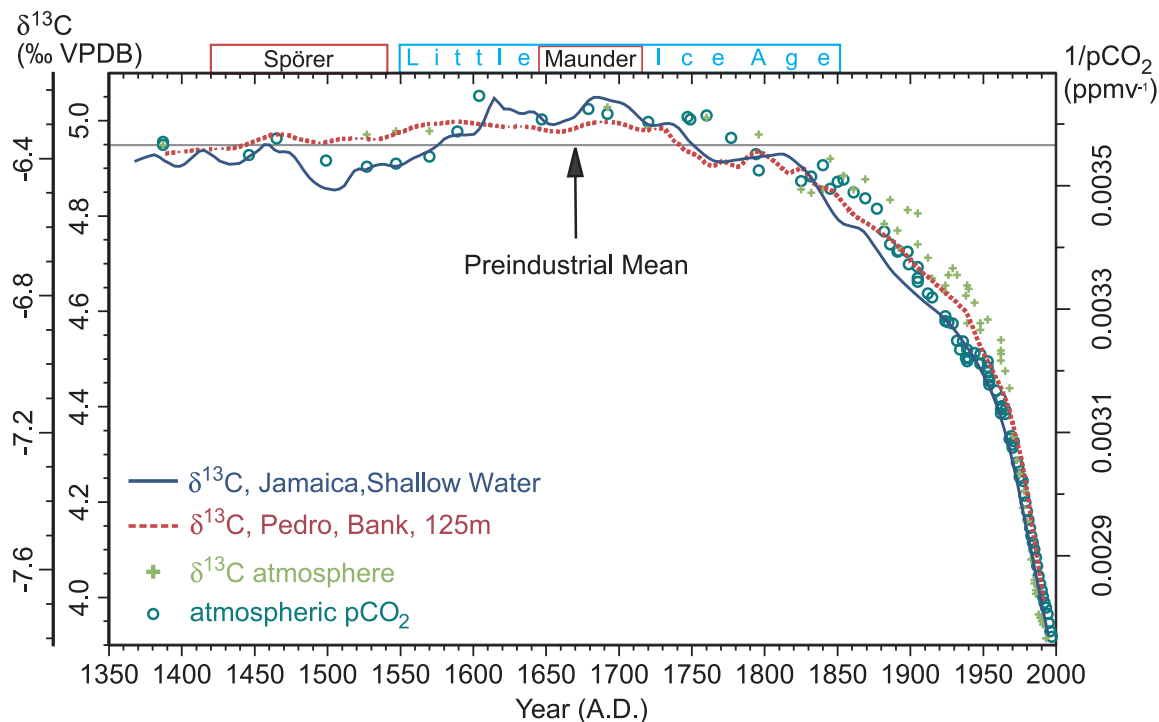


**Figure 3.** Stable carbon isotope records of the *Ceratoporella* skeletons. Error bars are  $\pm 2\sigma$  (standard deviation). The lines are locally weighted linear regressions (LOWESS, 10% window). Black arrows mark U/Th dates as shown in Figure 1. Open arrows mark positions of the bomb spike in samples Ce145 [Böhm *et al.*, 1996] and 392-19. Hatched bars in Figures 3c and 3f mark the positions of the bleached and bored zones in sample Ce95-2. Values from these bands may be slightly altered. Sample Ce145 is plotted with sample Ce96-1 for comparison.

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## 4. Results

[9] All  $\delta^{13}\text{C}$  records (appendix A) show the full extent of the industrial decline (Figure 3) caused by the anthropogenic addition of  $^{12}\text{C}$ -enriched  $\text{CO}_2$  to the atmosphere [Druffel and Benavides, 1986;



**Figure 4.** Comparison between Caribbean shallow (blue solid line) and deeper water (red dotted line) sponge  $\delta^{13}\text{C}$  records (inner left y axis),  $\delta^{13}\text{C}$  of atmospheric  $\text{CO}_2$  (green crosses, outer left y axis) and atmospheric  $p\text{CO}_2$  (circles, right y axis, reciprocal scale). Atmosphere data are derived from Antarctic ice core and firm air inclusions [Etheridge *et al.*, 1996; Francey *et al.*, 1999] and air measurements [Keeling and Whorf, 2001]. The atmospheric  $\delta^{13}\text{C}$  record is scaled for its preindustrial mean and minimum values to fit the shallow water sponge record. The  $\delta^{13}\text{C}$  curves are smoothed by locally weighted linear regressions (9% window). The horizontal line marks the preindustrial (1400–1850 A.D.) means (280.5 ppm,  $-6.37\text{‰}$ ,  $4.95\text{‰}$  VPDB, respectively). Slight offsets in the timing between atmosphere and sponge records can be artefacts of the dating methods. Blue bar at the top shows the approximate duration of the Little Ice Age [Grove, 1988]. Red bars mark the Spörer (ca. 1420–1540 A.D.) and Maunder (1645–1715 A.D.) sunspot minima.

Keeling *et al.*, 1979; Friedli *et al.*, 1986; Francey *et al.*, 1999; Quay *et al.*, 1992; Laube-Lenfant and Pierre, 1994; Bacastow *et al.*, 1996]. In all records the industrial period shows an older segment of slowly declining  $\delta^{13}\text{C}$  values. In contrast, the younger segment, starting about 1960 A.D., exhibits rapidly declining values (Figures 3d–3f). This parallels the records of atmospheric  $p\text{CO}_2$  and its  $^{12}\text{C}$  content, measured in ice core inclusions and air, which started to increase at higher rates between 1950 and 1960 A.D. [Friedli *et al.*, 1986; Etheridge *et al.*, 1996; Francey *et al.*, 1999] (Figure 4).

[10] The industrial decline in  $\delta^{13}\text{C}$  started in the first half of the 19th century after a short period of stable values around 1800 A.D. This late preindustrial plateau shows nearly constant values in all

measured sponges (Figure 3). The stable period was preceded by a slight decline in  $\delta^{13}\text{C}$  in the middle of the 18th century. The highest measured  $\delta^{13}\text{C}$  values of both shallow water records occur during the 17th and early 18th century (Figure 3b and 3c). Before that period, i.e., before about 1600 A.D., values were in the same range as in the early 19th century. In the shallow water sponge records the mean  $\delta^{13}\text{C}$  of the 1400–1500 A.D. interval is significantly lower ( $p < 0.01$ ) than the 1620–1720 A.D. mean (Table 1).

[11] Specimen Ce95-2 has two bio-eroded zones interrupting the isotope record (Figure 3c). It shows a relatively high scatter in the preindustrial part of the record. Two parallel isotope transects were sampled at a distance of  $\sim 1\text{cm}$ . We find an offset of  $\sim 0.1\text{‰}$  between corresponding values of the

**Table 1.** Mean  $\delta^{13}\text{C}$  for Certain Time Intervals Calculated From the Measured Values<sup>a</sup>

Sample Depth	Mean $\delta^{13}\text{C}$ (‰ VPDB $\pm 2\sigma/\sqrt{n}$ ) of specified intervals (A.D.)				$\Delta\delta^{13}\text{C}_s$		$\Delta\delta^{13}\text{C}_s/\Delta\delta^{13}\text{C}_a$		
	1400–1500	1620–1720	1400–1850	1960–1970	1980–1990	pi to 1960s	1960s–1980s	pi to 1960s	1960s–1980s
Ce96-1	4.92 $\pm$ 0.01	5.02 $\pm$ 0.01	4.95 $\pm$ 0.01	4.37 $\pm$ 0.03	4.06 $\pm$ 0.13	–0.58 $\pm$ 0.03	–0.31 $\pm$ 0.13	0.9 $\pm$ 0.09	0.5 $\pm$ 0.22
20 m	<i>n</i> = 39	<i>n</i> = 34	<i>n</i> = 147	<i>n</i> = 2	<i>n</i> = 3				
Ce145	–	–	(4.95)	4.41 $\pm$ 0.03	4.12 $\pm$ 0.04	–0.54 $\pm$ 0.03	–0.29 $\pm$ 0.05	0.8 $\pm$ 0.08	0.5 $\pm$ 0.10
25 m				<i>n</i> = 7	<i>n</i> = 12				
Ce95-2	4.92 $\pm$ 0.01	5.03 $\pm$ 0.02	4.98 $\pm$ 0.01	4.39 $\pm$ 0.05	4.07 $\pm$ 0.06	–0.59 $\pm$ 0.05	–0.32 $\pm$ 0.08	0.9 $\pm$ 0.11	0.5 $\pm$ 0.13
20 m	<i>n</i> = 13	<i>n</i> = 16	<i>n</i> = 71	<i>n</i> = 7	<i>n</i> = 7				
392-19	4.96 $\pm$ 0.01	4.98 $\pm$ 0.01	4.95 $\pm$ 0.01	4.42 $\pm$ 0.04	4.12 $\pm$ 0.07	–0.53 $\pm$ 0.04	–0.30 $\pm$ 0.08	0.8 $\pm$ 0.09	0.5 $\pm$ 0.14
125 m	<i>n</i> = 17	<i>n</i> = 20	<i>n</i> = 107	<i>n</i> = 4	<i>n</i> = 4				

<sup>a</sup> For sample Ce95-2, only data from the first transect are used. Here 1400–1850 means are calculated from the smoothed data sets interpolated to 1 year intervals. Decadal means are read from linear regression lines through respective decadal data. Preindustrial record of sample Ce95-2 is fragmentary; therefore the preindustrial mean may be slightly biased. Last four columns show measured marine decrease ( $\Delta\delta^{13}\text{C}_s$ ) from preindustrial mean (pi) until 1960s and from 1960s to 1980s. Preindustrial mean for Ce145 is copied from Ce96-1. Last two columns give ratios between these values and the respective atmospheric carbon isotopic decline ( $\Delta\delta^{13}\text{C}_a$ ; ice core, and firm air inclusion data [Francey *et al.*, 1999]), which is pi to 1960s:  $-0.67\text{‰}$ , 1960s to 1980s:  $-0.59\text{‰}$ .

two transects in the preindustrial part (Figure 3c). The second transect was sampled almost 3 years after the first one; therefore the observed  $\delta^{13}\text{C}$  offset may represent a similar ageing effect as previously described for sponge aragonite [Druffel and Benavides, 1986]. Nevertheless, the means of the 1400–1500 A.D. interval (transect 1: 4.92‰; transect 2: 4.81‰) and of the 1620–1720 A.D. interval (transect 1: 5.03‰; transect 2: 4.93‰) are again significantly different ( $p \ll 0.01$ ) for both transects respectively. Furthermore, the means of the respective intervals from the first transect and sample Ce96-1 are identical (Table 1).

[12] The deeper water sample from Pedro Bank shows no obvious  $\delta^{13}\text{C}$  maximum during the 17th or 18th century but rather very constant values in most of the preindustrial interval (Figure 3a). However, even in this sample we find a 1400–1500 A.D. mean that is slightly lower than the 1620–1720 A.D. mean (Table 1). The difference is significant at a 95% probability level.

## 5. Discussion

### 5.1. Preindustrial Variations

[13] The sponge records provide the first preindustrial marine  $\delta^{13}\text{C}$  data with decadal resolution. Additionally, they cover the entire industrial period with higher accuracy, better temporal resolution and better timing constraints than previous records [Druffel and Benavides, 1986; Böhm *et al.*, 1996]. The excellent reproducibility between records from different sites of similar water depth (Figure 3b and 3e) shows that sponge skeletons can reliably register regional environmental signals. The great similarity between the Jamaica and Pedro Bank records indicates that the measured carbon isotope signal is representative for open sea surface waters of the central Caribbean and of the equatorial to tropical/subtropical Atlantic Ocean.

[14] Surface water studies [Bacastow *et al.*, 1996; McNichol and Druffel, 1992; Gruber *et al.*, 1998] show a high  $\delta^{13}\text{C}_{\text{DIC}}$  variability at timescales of days to months with amplitudes significantly greater than 0.1‰. Variations with periods of 3–5 years and amplitudes of  $\sim 0.1\text{‰}$  have been

observed on interannual timescales [Bacastow *et al.*, 1996]. Also,  $\Delta^{14}\text{C}$  in subtropical surface waters may vary significantly on subdecadal to decadal timescales while lacking the expected Suess effect long-term trend [Druffel, 1997]. Such subdecadal variations are explained by advection, vertical mixing, and productivity changes. Longer surface water  $\delta^{13}\text{C}_{\text{DIC}}$  time series have not been available up to now. With the precision and resolution of our records (sample spacing  $\sim 2\text{--}6$  years), we can reliably detect  $\delta^{13}\text{C}$  variations  $> 0.05\text{‰}$  on interdecadal and centennial timescales ( $> 15\text{--}20$  years). Our records (Figure 3) demonstrate that variations in this time band are very small in the Caribbean. This is most obvious for the deeper water setting at Pedro Bank, where the preindustrial (A.D. 1400–1850)  $\delta^{13}\text{C}$  variation is only  $\pm 0.07\text{‰}$ . Most of this variability is caused by a decrease of roughly  $0.1\text{‰}$  between A.D. 1730 and 1760 (Figure 4).

[15] The most prominent feature of the preindustrial shallow water record (Figure 3b) is the 50 years long  $0.1\text{‰}$  increase in  $\delta^{13}\text{C}$  from about 1550 to 1600 A.D. and the subsequent decrease until about 1770 A.D. (Figure 4). The  $\delta^{13}\text{C}$  of atmospheric  $\text{CO}_2$  increased by  $\sim 0.05$  to  $0.10\text{‰}$  [Francey *et al.*, 1999] and atmospheric  $p\text{CO}_2$  decreased by  $\sim 6$  ppm [Etheridge *et al.*, 1996] during the same period (Figure 4). Unfortunately, the coarse resolution and limited dating accuracy of both the sponge and ice core records currently prohibit conclusions about the precise temporal relation between the atmospheric and marine records.

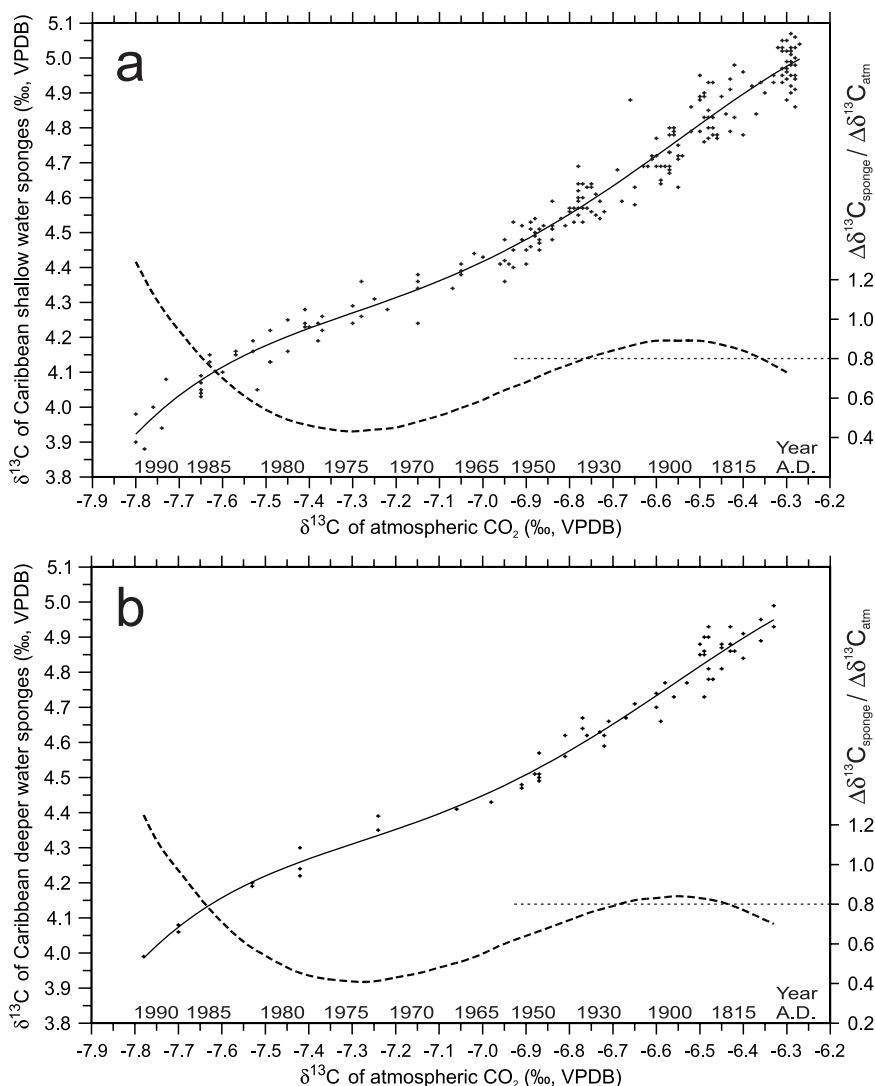
[16] The  $\delta^{13}\text{C}$  of atmospheric or marine dissolved total  $\text{CO}_2$  is expected to correlate with  $1/p\text{CO}_2$  [Keeling *et al.*, 1979] as long as  $\text{CO}_2$  is sequestered by or released from organic carbon. This is due to the strong  $^{13}\text{C}$  depletion of organic carbon. Addition of such isotopically light carbon to the atmosphere/ocean inorganic carbon pool will in parallel increase the  $\text{CO}_2$  and its  $^{12}\text{C}$  content. There are, of course, other mechanisms to change atmospheric  $p\text{CO}_2$  and the isotopic composition of  $\text{CO}_2$  or DIC independently, for example, changes in the ocean salinity or in the deepwater carbonate ion content will affect atmospheric  $p\text{CO}_2$  without isotopic effects [Broecker, 1995].

[17] The period of  $p\text{CO}_2$  reduction and  $\delta^{13}\text{C}$  increase (1550–1700 A.D.) starts after the Spörer sunspot minimum and coincides with the Maunder sunspot minimum (approximately 1420–1540 and 1645–1715 A.D.; Figure 4). These are periods of reduced solar activity [Stuiver and Quay, 1980; Lean *et al.*, 1995]. Moreover, the period between 1550 and 1850 A.D. was an interval of intermittent cool periods, called “Little Ice Age” [Grove, 1988; Jones *et al.*, 2001]. Thus global cooling may have been the cause of the parallel  $p\text{CO}_2$  reduction and atmospheric/marine  $\delta^{13}\text{C}$  increase [Trudinger *et al.*, 1999]. Increased solubility of  $\text{CO}_2$  in colder surface waters or carbon sequestration in the terrestrial biosphere was discussed previously [Etheridge *et al.*, 1996; Trudinger *et al.*, 1999; Joos *et al.*, 1999]. Only the latter can explain the parallel atmospheric and marine  $\delta^{13}\text{C}$  increase.

[18] Global cooling of sea surface temperatures would sequester atmospheric  $\text{CO}_2$  due to its higher solubility by  $\sim 10$  ppm/K. Models show that on timescales of some decades about half of this amount will be taken up by the oceans [Bacastow, 1996]. Thus global sea surface temperature cooling alone could explain the  $p\text{CO}_2$  decrease, if the globally averaged cooling during the Little Ice Age was  $\sim 1$  K [Grove, 1988; Trudinger *et al.*, 1999; Keigwin, 1996; Kitagawa and Matsumoto, 1995; Watanabe *et al.*, 2001]. However, a global 1 K cooling of sea surface temperatures would increase the isotopic equilibrium fractionation between air and water by  $\sim 0.1\text{‰}$  [Zhang *et al.*, 1995] and this would only lower the  $\delta^{13}\text{C}$  of atmospheric  $\text{CO}_2$  and leave the marine  $\delta^{13}\text{C}_{\text{DIC}}$  almost unchanged. This is because on a multi-decadal timescale the exchangeable carbon mass in the ocean is much larger than the mass of carbon in the atmosphere [Keir *et al.*, 1998]. Therefore the observed parallel increase of  $\delta^{13}\text{C}_{\text{DIC}}$  and of atmospheric  $\delta^{13}\text{C}$  during the period of lower  $p\text{CO}_2$  requires an alternative explanation.

[19] The observed close correlation of our data with atmospheric  $p\text{CO}_2$  indicates a common and therefore global cause. The most likely common cause of all observed carbon cycle changes is an increase in the global biogenic carbon mass during the Little Ice Age [Trudinger *et al.*, 1999; Joos *et al.*, 1999].





**Figure 5.** Cross plot of carbon isotope values of Caribbean shallow water sponge aragonite and atmospheric CO<sub>2</sub> [Francey *et al.*, 1999]. Age increases from left to right with increasing atmospheric values. The continuous line is a fourth-order polynomial fit to the data. The dashed line (right y axis) is the first derivative of the polynomial fit. It represents the ratio of carbon isotopic change between Caribbean surface water DIC and atmospheric CO<sub>2</sub>. This ratio varied around an average value of 0.8 before 1960, declined significantly during the 1960s and 1970s and reached its highest values near 1 in the 1990s. (a) Shallow water specimens from Jamaica. (b) Deeper water specimen from Pedro Bank.

At the same time, the combined atmospheric and marine data allow for only minor global cooling of the ocean surface.

[20] It is unknown whether sequestration of organic carbon to the deep sea by increased oceanic productivity or rather increased storage of organic matter in terrestrial environments caused the observed changes in the earth's carbon budget during the Little Ice Age. Changes in the large-

scale ocean circulation [Broecker, 2000] but also increased wind induced upwelling in the southern Caribbean and subtropical North Atlantic [Black *et al.*, 1999; deMenocal *et al.*, 2000] may additionally have contributed to the observed variation of Caribbean  $\delta^{13}\text{C}_{\text{DIC}}$  during the Little Ice Age. Only additional long-term records of  $\delta^{13}\text{C}_{\text{DIC}}$  from other oceans will help to find out whether our observations represent a purely global signal. The only published Pacific Ocean carbon isotope record



reaching back to the Little Ice Age of a coralline sponge from the Great Barrier Reef [Wörheide *et al.*, 1997] shows slightly higher values ( $\sim 0.05\%$ ) during the 17th century, which is in accord with a global nature of the observed marine variations.

## 5.2. Industrial Period

[21] The sharp contrast between the almost invariant preindustrial record and the strong decline in  $\delta^{13}\text{C}$  since 1850 A.D. underlines the exceptional quality of this recent anthropogenic disturbance of the global carbon cycle. Comparing the observed decrease of  $\delta^{13}\text{C}$  in Caribbean surface water and in the atmosphere [Francey *et al.*, 1999], we find a consistent and significant decline in the ratio of the marine to the atmospheric change ( $\Delta\delta^{13}\text{C}_{\text{DIC}}/\Delta\delta^{13}\text{C}_{\text{Atm}}$ ) from the pre-1960 period to the 1970s and 1980s (Table 1, Figure 5): The marine  $\delta^{13}\text{C}$  decrease from preindustrial times to the 1960s was  $\sim 80\%$  of the atmospheric change. The marine  $\delta^{13}\text{C}$  decrease from the 1960s to the 1980s, however, was only half the atmospheric change. This decline in the apparent isotopic equilibration between atmosphere and surface waters is evident in all four sponge records. Therefore it is unlikely that this observation represents just an artefact of insufficient dating control and unnoticed decadal growth rate variations of our sponges.

[22] In contrast to our observation of a reduced isotopic ocean-atmosphere equilibration after the 1960s, Gruber *et al.* [1999] report a  $\delta^{13}\text{C}_{\text{DIC}}$  decrease during the decade from 1983 to 1995 in the Sargasso Sea that equals or possibly exceeds the contemporary atmospheric  $\delta^{13}\text{C}$  decline. However, the youngest part of our isotope record from Specimen Ce95-2 points to a decrease of  $\sim 0.2 \pm 0.07\%$  from the 1980s until 1995 (Figure 3f). This decrease is compatible with the observations of Gruber *et al.* [1999]. Obviously, after the minimum of the 1970s (Figure 5), the isotopic ocean-atmosphere equilibration increased again to the very high values of the 1990s reported by Gruber *et al.* [1999] and indicated by the youngest portion of our sponge records.

[23] These variations of the isotopic ocean-atmosphere disequilibrium in the Caribbean Sea can be explained by the history of the atmospheric  $\delta^{13}\text{C}$ .

The rate of decline of atmospheric  $\delta^{13}\text{C}$  increased during the 1950s and 1960s, remained high until the late 1980s, and was low during the 1990s [Francey *et al.*, 1999; Gruber *et al.*, 1999]. During the first half of the 20th century the carbon isotopic decline of the atmosphere was slow enough for the tropical surface ocean DIC to follow closely ( $\sim 80\%$  “equilibration”, i.e.,  $\Delta\delta^{13}\text{C}_{\text{DIC}}/\Delta\delta^{13}\text{C}_{\text{Atm}}$ , in Caribbean and tropical North Atlantic surface waters, Table 1, Figure 5). However, with the finite carbon isotopic equilibration time between atmosphere and surface waters, the faster atmospheric  $\delta^{13}\text{C}$  decrease during the 1960s and 1970s led to a lower degree of equilibration ( $\sim 50\%$ ). The subsequent slow down of the 1990s allowed the oceans to start catching up on the isotopic equilibrium (100–125% equilibration [Gruber *et al.*, 1999];  $\sim 100\%$  in Figure 5). This varying degree of isotopic equilibration between atmospheric  $\text{CO}_2$  and surface water DIC clearly reflects the current state of the global carbon cycle, far from steady state due to the serious anthropogenic interference of the last 200 years.

## 6. Conclusions

[24] This study shows that the coralline sponge *Ceratoporella nicholsoni* is a very accurate recorder of  $\delta^{13}\text{C}_{\text{DIC}}$ . Coralline sponges are very useful in providing long-term, high-resolution  $\delta^{13}\text{C}$  records. These provide useful information for estimating ocean uptake of anthropogenic  $\text{CO}_2$  [Quay *et al.*, 1992; Bacastow *et al.*, 1996; Sonnerup *et al.*, 1999, 2000] and the terrestrial biota turnover times [Keir *et al.*, 1998].

[25] Our data shows a period of higher values during the Little Ice Age. These preindustrial variations are in concert with changes in the atmospheric carbon reservoir. Our data provide an independent evidence for variations in the global carbon reservoirs prior to the onset of anthropogenic perturbations.

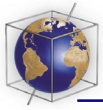
[26] Varying degrees of carbon isotopic equilibration between atmospheric  $\text{CO}_2$  and surface water DIC during the industrial increase of atmospheric  $p\text{CO}_2$  reflect the dynamic response of the carbon cycle to human interference with magnitudes and



**Appendix A.** Carbon Isotope Values From Caribbean Sponge Specimens Used in This Study<sup>a</sup> [The full Appendix A is available in ASCII tab-delimited format at <http://www.g-cubed.org>.]

Ce145, 25 m		Ce96-1, 20 m		Ce95-2, 20 m, First Transect		Ce95-2, 20 m, Second Transect		392-19, 125 m	
Year A.D.	d <sup>13</sup> C	Year A.D.	d <sup>13</sup> C	Year A.D.	d <sup>13</sup> C	Year A.D.	d <sup>13</sup> C	Year A.D.	d <sup>13</sup> C
1986	4.07	1995	3.93	1994	3.92	1993	3.90	1992	3.99
1985	4.13	1986	4.03	1993	3.98	1992	3.88	1992	3.99
1985	4.15	1982	4.05	1986	4.09	1989	3.94	1987	4.06
1984	4.10	1977	4.23	1986	4.05	1986	4.04	1987	4.08
1983	4.16	1971	4.28	1981	4.13	1983	4.15	1982	4.19
1982	4.19	1967	4.38	1974	4.24	1980	4.16	1982	4.20
1981	4.22	1960	4.41	1970	4.38	1976	4.19	1979	4.22
1980	4.25	1954	4.49	1970	4.24	1973	4.26	1979	4.24
1979	4.24	1947	4.47	1965	4.44	1970	4.36	1977	4.30
1978	4.22	1940	4.53	1959	4.53	1968	4.34	1972	4.35
1977	4.23	1932	4.55	1959	4.40	1964	4.43	1972	4.39
1977	4.28	1926	4.61	1952	4.48	1961	4.36	1967	4.41
1976	4.24	1921	4.57	1945	4.59	1959	4.45	1963	4.43
1974	4.29	1915	4.64	1945	4.59	1956	4.41	1957	4.48
1973	4.26	1909	4.58	1936	4.57	1953	4.45	1957	4.47
1972	4.31	1904	4.65	1923	4.69	1951	4.49	1952	4.50
1970	4.34	1899	4.63	1923	4.62	1948	4.51	1952	4.49
1967	4.41	1894	4.69	1916	4.64	1944	4.54	1950	4.51
1964	4.43	1889	4.67	1911	4.59	1942	4.57	1947	4.57
1962	4.41	1883	4.68	1906	4.77	1938	4.57	1947	4.51
1961	4.42	1878	4.72	1906	4.72	1936	4.60	1943	4.56
1957	4.48	1873	4.79	1898	4.75	1933	4.56	1939	4.62
1956	4.45	1869	4.77	1892	4.88	1930	4.59	1934	4.62
1954	4.50	1864	4.80	1886	4.86	1927	4.54	1930	4.62
1953	4.51	1860	4.77	1884	4.80	1924	4.55	1930	4.59
1951	4.50	1860	4.76	1880	4.80	1921	4.53	1928	4.63
1949	4.53	1855	4.78	1879	4.79	1918	4.57	1928	4.63
1948	4.46	1850	4.79	1871	4.83	1915	4.63	1921	4.67
1946	4.52	1845	4.78	1836	4.95	1912	4.68	1921	4.64
1945	4.51	1840	4.85	1824	4.93	1909	4.63	1913	4.66
1943	4.52	1835	4.89	1824	4.93	1906	4.69	1911	4.67
1941	4.57	1830	4.88	1814	4.94	1904	4.69	1909	4.71
1938	4.56	1826	4.90	1804	4.84	1901	4.69	1905	4.66
1937	4.57	1822	4.93	1802	4.93	1898	4.71	1899	4.73
1935	4.64	1818	4.89	1796	4.93	1896	4.64	1895	4.70
1991	4.00	1813	4.98	1795	4.95	1891	4.71	1891	4.74
1988	4.08	1808	4.96	1790	5.03	1889	4.78	1885	4.77

<sup>a</sup> Note that the ages given are only approximate estimates based on interpolations of U/Th dated layers! Carbon isotope values are given in ‰ VPDB.



rates of change that are unprecedented for the last 600 years and probably for the whole Holocene.

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