PAIRED ¹⁴C AND ²³⁰Th/U DATING OF SURFACE CORALS FROM THE MARQUESAS AND VANUATU (SUB-EQUATORIAL PACIFIC) IN THE 3000 TO 15,000 CAL YR INTERVAL

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ABSTRACT. Paired radiocarbon and ²³⁰Th/U dating was performed on 13 surface corals from submerged reefs in the Marquesas and from raised terraces in Vanuatu. The absolute ages of the corals analyzed ranged from 3000 to 15,000 cal yr. Estimates of the difference between the absolute and ¹⁴C ages of these corals are in agreement with previous determinations up until 11,500 cal yr. The resulting mean sea surface reservoir age *R* is determined at 390 ± 60 yr for the Marquesas region (9°S), which is slightly higher than the *R* value at 280 ± 50 yr for the Tahiti Islands (18°S). Multiple ¹⁴C analyses of 2 corals from the Marquesas present scattered ¹⁴C ages at ~12,000 and ~15,100 cal yr. This could be attributed to rapid changes of the ¹⁴C content of surface waters around the Marquesas Islands or to a subtle submarine diagenesis.

INTRODUCTION

Radiocarbon and/or ²³⁰Th/U dating of surface corals have mainly been used to quantify the timing and amplitude of sea-level fluctuations, and also to estimate the deviation between the ¹⁴C and calendar (²³⁰Th/U) ages. Superimposed on the long-term ¹⁴C variation over the past 40,000 cal BP (Stuiver et al. 1998; Edwards et al. 1993; Bard et al. 1990a, 1993, 1998; Voelker et al. 1993; Burr et al. 1998; Hughen et al. 1998, 2000; Kitagawa and van der Plicht 1998, 2000; Beck et al. 2000), short-term fluctuations have been recorded in both terrestrial and marine materials and are considered to be linked to changes of solar activity and of the global carbon cycle (Stuiver et al. 1998). In the marine environment, the modern oceanic circulation controls mixing of "old" waters from the deep ocean with surface ocean waters. This leads to latitudinal variations of the sea surface reservoir ages, *R* (Bard 1988). As the oceanic circulation has changed in the past, so have the *R* values at midto high latitudes (Bard et al. 1994; Sikes et al. 2000; Siani et al. 2001). Modern surface corals record seasonal to centennial ¹⁴C changes of some 20–40‰, related to both the atmospheric ¹⁴C and oceanic circulation changes (Druffel 1997; Burr et al. 1998; Guilderson and Schrag 1998). Substantial local intra-ocean variations in *R* were also recently suggested at similar latitudes for the Pacific Ocean (in Stuiver et al. 1998; Goslar et al. 2002).

We present here the results of coupled ¹⁴C and ²³⁰Th/U dating of surface corals, collected from raised terraces from Vanuatu (Cabioch and Ayliffe 2000) and from submerged reefs around the Marquesas Islands (Figure 1). The absolute ages obtained for these corals by ²³⁰Th/U dating ranged from 3000 to ~15,000 cal yr. The difference between the absolute and ¹⁴C ages of these corals are compared to those found in previous studies (Figure 2) (Stuiver et al. 1998; Bard et al. 1990a, 1993, 1998; Edwards et al. 1993; Burr et al. 1998; Hughen et al. 1998, 2000). *R* estimates from the Marquesas and Tahiti are directly compared in the time interval of 9000–11,000 cal yr.

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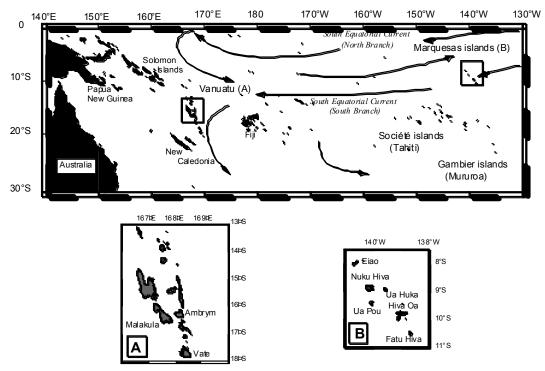


Figure 1 Sample locations; surface currents are also represented

MATERIAL AND METHODS

The surface corals (*Porites sp.*) were collected in the Marquesas Islands (6°S; 140°W) by dredging of submerged coral reefs in water depths greater than 86 m during the cruise conducted by the IRD in 1997 (Musorstom and Paleomarq), and in Vanuatu (20°S; 170°E) by direct sampling of emerged marine terraces raised by tectonic uplift (Figure 1). The corals from the Marquesas have been submerged in seawater since the time of their formation and have as a result most likely been preserved from major recrystallization/dissolution processes.

Coral specimens were cut into small slabs with cross-sectional areas between 4–6 cm² and thickness of ~0.1 to 0.2 cm. The shortest axis of these slabs were oriented parallel to the direction of growth. Growth bands were visible and continuous in the corals sampled. The slabs cut from each specimen probably represent some 1 to 15 yr in time, given the range of possible growth rates documented by Priess (1997) of ~2 to 40 mm per yr for the *Porites* species. Each coral slab was then subsampled for X-ray diffractometry (XRD), ¹⁴C, and ²³⁰Th/U analyses. Coral samples having a magnesian calcite or pure calcite content lower than 2% were selected for dating with 1 exception (Table 1).

²³⁰Th/U Dating

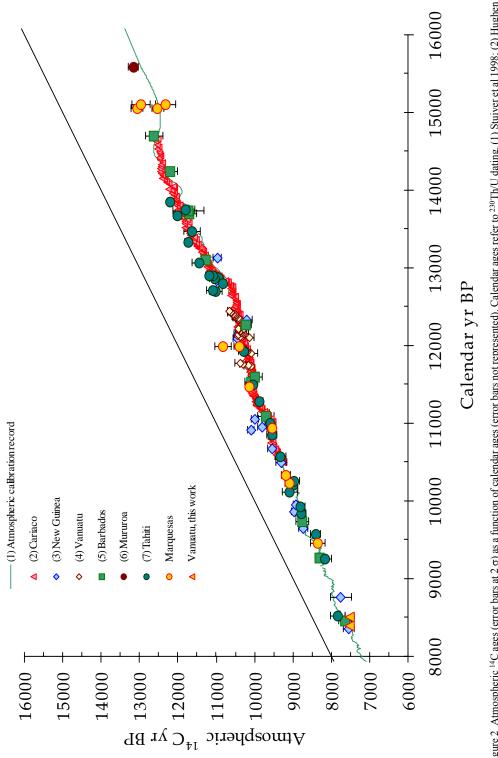
The U/Th age determination follows the procedure by Cabioch and Ayliffe (2000). Exterior surfaces of corals samples were initially cleaned by physical abrasion with a dental drill bit. The coral samples were then further cleaned by ultrasound treatment and several rinses with quartz distilled (QD) water. U and Th fractions were extracted and purified from coral samples using standard ion exchange chemistry as described by Stirling et al. (1995). The purified U and Th separates were loaded onto zone-refined Re filaments between 2 layers of colloidal graphite. Isotopes of U and Th

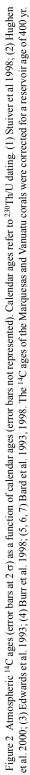
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the c rral co s 17b 5.99 5.99 vulate	2 σ (yr)	110	110	190	260	170 200	150	210	150	210	570 150	440 150		
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ory coc 2000) 2 ages 1ues (1 arenthe	Δ ¹⁴ C (‰)	-35 (15)	-38 (11)	56 (110)	57	(111)	57		88 (112)	((+1)	78 (134)		55 (100)	(168) (168)
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as and Vanuatu. Colu n m relatively to sea sed in conventional error similarly noted to correspond to	²³⁰ Th/U 2 σ ¹⁴ C age 2 σ ¹⁴ C age [*] 2 σ Δ ¹⁴ C 2 σ R 2 σ Aragonite 232] 2 σ cal yr (yr) (BP) (yr) (BP) (yr) χ^2 (%0) (%0) (yr) (yr) %0	10 20 15,050 120 12,830* 180 <u>29.166</u> 13,410 ^{\$} 200 13,560 ^{\$} 260 12,920* 150 3.364 236 25 (298)	$13,120^{*}$ 260 $13,470^{\$}$ 160 0.836 1.55 24 (214)	20 15,100 120 13,360 240 <u>13.086</u>	12,720 260 276 42 (341)		60 10 6400 60 6000 160 28 22 385 165 100 (93)	$50 ext{ 90 } 8390 ext{ 70 } 8060 ext{ 140 } 8020 ext{ 91 } 0.5765 ext{ 17 } 15 ext{ 415 } 95 ext{ 100 } 100 ext{ 781}$	7990 120	20 10 8500 60 8010 120 32 17 270 130 100
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l Vanuatu slatively i convent imilarly corresp	²³⁰ Th/U cal yr	15,050		15,100			6400	8390		8500
as and n m re sed in error s : 1000	2 a	20		20			10	06		10
Marques: altitude i ure expres hean and (nean and (ontinued)	[230/232]	2570		2980			1750	18,730		920
m the nd the ages a hted m ^k calendai	5σ	2.1		2.6			2.2	2.8		2.4
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nd ¹⁴ C a pth in m measure were us 315 n = <u>e reserve</u>	(3)	17b 2 3.429 17b 2 17b 1	17b 1	80a 1	80a 1		15a 1 2.360	74a 2 2.938	74a 2	73a 2 2.945
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were measured with a Finnigan MAT 262 solid-source mass spectrometer using a peak jumping routine with a secondary electron multiplier. Repeated analyses of the uraninite standard HU-1 containing ²³⁸U, ²³⁴U, and ²³⁰Th in radioactive equilibrium yielded an average [²³⁴U/²³⁸U] activity ratio of 1.0015 \pm 0.0032 (N = 9) (= ²³⁴U/²³⁸U atomic ratio of 54.98 \pm 0.18 \times 10⁻⁶) and a [²³⁰Th/²³⁸U] activity ratio of 1.0004 \pm 0.0044 (N = 9). Decay constants used in the age calculations were λ^{230} Th = 9.1954 \times 10⁻⁶ \pm 7.19 \times 10⁻⁸ yr⁻¹ and λ^{234} U = .8349 \times 10⁻⁶ \pm 5.7 \times 10⁻⁹ yr⁻¹. The reported ²³⁴U/²³⁸U and ²³⁰Th/²³²Th activity ratios of all the dated corals were corrected for machine biases by normalization to the corresponding ratios determined for HU-1.

14C Dating

Secondary crystallization in skeleton pores of corals or surface contaminants are usually eliminated by strong acid or stepwise leaching procedures (Bard et al. 1990b; Burr et al. 1992; Yokoyama et al. 2000). Such procedures are time consuming due to the partial dissolution of the sample in several aliquots and then accordingly to the increase of the number of ¹⁴C measurements to check cleaning efficacy (Burr et al. 1992; Yokoyama et al. 2000). In this study, coral samples (~20 mg in size) were first pre-cleaned by sand blasting until the elimination of micrite forms around skeleton pores (Figure 3), which can be carefully controlled under a microscope. Under this procedure, coral samples lose between 20–60% of their initial weight. Next, the abraded coral sample is rinsed and ultrasonically cleaned and then crushed in an agate mortar. About 10 mg of the fine powder is then immediately introduced into 1 side of a reaction vessel with 2 side arms. The coral powder is then leached in a 2-cm³ solution of HNO₃ (0.01N) for 15 min and then rinsed to neutral pH. The second arm of the reactor is then filled with 1 cm³ of H₃PO₄ and the vessel containing the wet powder is immediately attached to the vacuum line and evacuated as suggested by Schleicher et al. 1998. After hydrolysis of the coral powder by reaction with the H₃PO₄ *in vacuuo*, the evolved CO₂ is trapped into an ampoule. The CO₂ is then reduced into graphite according to the procedure of Arnold et al. (1989).

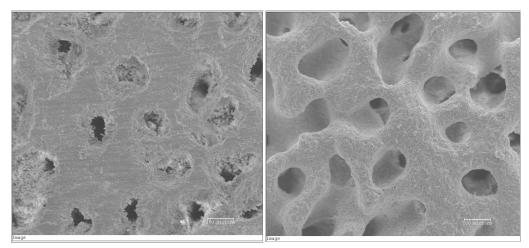


Figure 3 Scanning electron microprobe photographs of coral DR 1183 (17b) before (left) and after (right) cleaning. The sample on the right corresponds to an adjacent piece of coral treated for ¹⁴C dating.

Two sub-samples were randomly picked from different parts of each coral slab and prepared independently. Given the range of growth rates observed for *Porites sp.* corals (Priest 1997), it is quite likely that the 2 sub-samples taken from each coral specimen grew during different seasons or years. Two targets of Fe-C powder were made from each graphite preparation in order to increase the precision of the ¹⁴C ages by accelerator mass spectrometry (AMS) at the Gif-sur-Yvette Tandétron facility (Arnold et al. 1987).

RESULTS AND DISCUSSION

Evaluation of Cleaning Procedure of Corals

The efficacy of our cleaning procedure was initially assessed through ¹⁴C analyses of old corals (Table 2). In addition, we also proceeded to new ¹⁴C dating of corals from Tahiti and Mururoa, previously dated by Bard et al. (1993, 1996, 1998) (Table 3).

Blank values were assessed from 3 corals from Mururoa (Bard et al. 1993), Vanuatu (Cabioch and Ayliffe 2000), and the Marquesas, determined by 230 Th/U dating to be beyond the limit of the 14 C dating method (Table 2). Although the mean and standard deviation obtained for the corals from the Marquesas and Vanuatu (0.21 ± 0.05 pMC: apparent age: 49,350 ± 1900 BP) are slightly higher than those obtained for 1 coral (Irene 30) from Mururoa (0.16 ± 0.02 pMC; apparent age: 51,800 ± 1050 BP), they are nevertheless statistically indistinguishable from one another. The mean and standard deviation for the old ¹⁴C-depleted corals using this cleaning procedure are 0.20 ± 0.05 pMC (apparent age 50,100 ± 2070 BP) and are similar to slightly lower than those previously measured (Bard et al. 1990b; Burr et al. 1992; Yokoyama et al. 2000). The AIEA-C1 marble, currently used to assess the full procedural blank of the AMS ¹⁴C dating method of our laboratory, has a value of 0.08 ± 0.02 pMC (n = 7; apparent age 57,280 ± 2000 BP). As for the old ¹⁴C-depleted foraminifera reported by Schleicher et al. (1998) and Nadeau et al. (2001), the mean blank value of the old corals is larger than that of the geological C1 marble, which is thus inappropriate for ¹⁴C age calculation of young biological materials.

Corals from Tahiti and Mururoa, previously dated between 11,000 and 17,000 cal yr (Bard et al. 1990, 1993, 1998), were subjected to our new cleaning procedure and also to that previously introduced by Bard et al. (1990b), which involved a strong HCl leach (Table 3). The χ^2 tests for the ¹⁴C ages obtained using the previous and new cleaning procedures suggest no significant differences (at the 95% confidence level) between any of them. Therefore, it appears that our new cleaning treatment is as efficient as the previous one for removing surface contaminants from such corals.

Comparison of the Marine Reservoir R Ages from the Tahiti and Marquesas Corals in the Time Interval 9000–11,000 cal yr

The ¹⁴C and absolute ages of the coral sub-samples are reported in Table 1. By subtraction of the ¹⁴C ages of the corals from Vanuatu and Marquesas from the corresponding atmospheric ¹⁴C ages documented in the INTCAL98 tree-ring record, we estimate a reservoir age of ~400 yr for the surface of the central Pacific Ocean prior to 9000 cal yr (Table 1). Using a similar approach between 9000 to 11,000 cal yr, we determine a weighted mean *R* value of 390 ± 60 yr (weighted 2 σ error; n = 8) for the Marquesas corals, which is close to the mean global ocean value (Stuiver et al. 1998) (Figure 4). For the same time period, a slightly lower *R* of 280 ± 50 yr (weighted mean and 2 σ error; n = 8) is determined from the Tahiti corals. This small difference, as previously observed (Stuiver et al. 1998; Bard et al. 1998; Goslar et al. 2000), may be attributed to the location of these islands with respect to the Pacific surface water currents. While the Tahiti Islands are located in the well-ventilated South Pacific gyre, the Marquesas Islands are situated within the South Equatorial westward drift (Figure 1). The likely more extensive vertical mixing of surface waters with underlying ¹⁴C-depleted waters due to the Eastern Tradewinds in this region may account for the slightly older Marquesas *R* value.

Lab code Cor		Coral	Date of			1 σ	Apparent	1σ	230 Th/U	1 σ	
GifA	Dredge	code	preparation	C µg	pMC	(%)	age (BP)	(yr)	cal yr	(yr)	Site
100443		Ir30	26/6	680	0.20	0.03	50,020	1100	259,000*	0009	Mururoa
100445			27/6	1350	0.17	0.03	51,400	1230			Mururoa
100489*			28/9	2000	0.15	0.02	52,470	1340			Mururoa
100499			2/10	2130	0.16	0.02	51,920	1230			Mururoa
100500			9/10	1790	0.15	0.02	52,470	1350			Mururoa
100768			24/11	1150	0.16	0.02	51,920	1230			Mururoa
100769			24/11	1240	0.14	0.02	53,070	1250			Mururoa
100469	CP1262(1)	83a	L/L	1420	0.22	0.03	49,220	1080	63,860	260	Marquesas
100469			L/L	1420	0.17	0.02	51,400	1190			Marquesas
100470	CP1262(2)	14a	10/7	1810	0.25	0.03	48,140	066	64,410	260	Marquesas
100470			10/7	1810	0.25	0.03	48,140	870			Marquesas
100756		83a	14/11	1540	0.27	0.04	47,500	1040			Marquesas
100757		14a	21/11	1700	0.23	0.03	48,840	066			Marquesas
100473	np11 (+59)	44a	3/7	1384	0.26	0.03	47,820	970	66,980	290	Vanuatu
100473			3/7	1384	0.29	0.03	46,910	870			Vanuatu
100764		44a	23/11	1450	0.22	0.03	49,220	970			Vanuatu
100764		44a	23/11	1450	0.20	0.03	50,020	1080			Vanuatu
100474	Mallicolo	13a	3/7	1494	0.10	0.02	55,180	1490	107,580	540	Vanuatu
100474	(204 m) d		3/7	1494	0.17	0.02	51,400	1130			Vanuatu
100765		13a	16/11	1930	0.20	0.03	50,020	1080			Vanuatu
100764		44a	23/11	1450	0.22	0.03	49,220	970			Vanuatu
100766	Espiegle(1)	45a	24/11	1860	0.17	0.02	51,400	1150	121,270	580	Vanuatu
			Mururoa		0.16	0.02	51,840	1040			
			Marquesas		0.23	0.04	48,780	1260			
			Vanuatu		0.20	0.06	49,810	2320			
			Total		0.00	0.05	50 1 00	0000			

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Table 3 Comparison of the ¹⁴C ages of corals, obtained from different pre-cleaning procedures; M = mechanical sand blasting. The ¹⁴C ages are given in conventional yr BP uncorrected for the reservoir age. Samples noted by an asterisk were published in Bard et al. 1998. All samples have a carbon content higher than 500 µg, except sample GifA100494* (340 µg). The χ^2 values for P_{0.95} are 3.841 for n = 2; 5.991 for n = 3; and 7.815 for n = 4.

Sample code	Lab code GifA	Nr of targets	²³⁰ Th/U (cal yr)	2 σ (yr)	¹⁴ C age (BP)	2 σ (yr)	¹⁴ C age (weighted mean)	2 σ (yr)	χ^2 test	Comment
P7-8	95647*	2	11,280	30	10,100	140	10,160	90	1.293	HC1
P7-8	100480	2			10,200	160				HCl
P7-8	100490	2			10,210	180				М
P7-9	95648*	2	11,495	30	10,280	140	10,330	100	1.542	HCl
P7-9	100481	2			10,300	320				HCl
P7-9	100491	2			10,410	160				М
P7-11	95649*	2	12,875	40	11,130	140	11,230	90	5.215	HCl
P7-11	100482	2			11,240	160				HCl
P7-11	100492	2			11,390	180				М
P7-12	95650*	2	12,800	30	11,100	160	11,130	90	5.448	HCl
P7-12	100483	2			11,270	160				HCl
P7-12	100493	2			11,010	160				М
P8-1	95654*	1	12,905	30	11,510	220	11,480	90	2.942	HCl
P8-1	96092*	2			11,540	140				HCl
P8-1	100487	2			11,350	180				HCl
P8-1	100497	2			115,00	180				М
P8-2	95653*	2	13,335	30	11,970	160	12,030	80	4.209	HCl
P8-2	96091*	2			12,000	140				HCl
P8-2	100486	2			12,190	180				HCl
P8-2	100496	2			11,980	180				М
P8-3	95652*	2	13,665	35	12,250	160	12,260	90	3.113	HCl
P8-3	96090*	2			12,350	140				HCl
P8-3	100485	2			12,170	180				HCl
P8-3	100495	1			12,180	240				М
P8-4	95651*	2	13,850	35	12,570	160	12,490	90	6.202	HCl
P8-4	96087*	2			12,560	140				HCl
P8-4	100484	2			12,350	200				HCl
P8-4	100494	1			12,280*	280				М
Mu-8-30-315	95656*	2	17,170	40	14,860	180	14,790	120	1.274	HCl
Mu-8-30-315	100488	1			14,690	280				HCl
Mu-8-30-315	100498	1			14,750	200				М

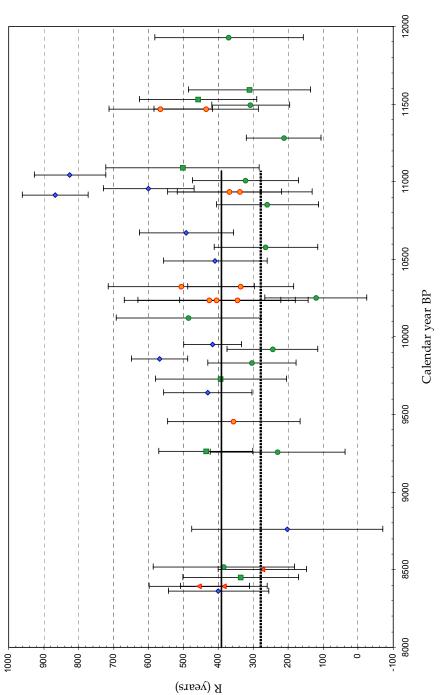


Figure 4 Sea surface reservoir ages (difference between coral and atmospheric ¹⁴C ages [Stuiver at al. 1998] of similar calendar ages). Black and dotted lines correspond to the weighted mean *R* value of the Marquesas and Tahiti corals. Same references as Figure 2.

Comparison of the ¹⁴C Ages and Δ^{14} C Values from the Marquesas and Vanuatu with Previous Marine and Atmospheric ¹⁴C Record

Between 3000 cal yr and 11,500 cal yr, the ¹⁴C ages are very reproducible within each coral (Table 1). Using an *R* value of 400 yr, the estimates of the difference between the absolute and ¹⁴C ages are in agreement with previous determinations (Stuiver et al. 1998; Burr et al. 1998; Edwards et al. 1993; Bard et al. 1993, 1998) (Figures 2, 5, 6). Beyond 11,500 cal yr, 2 corals dated at 12,000 and ~15,100 cal yr, however, present scattered ¹⁴C values.

The 2 sub-samples of coral specimen DR1183(2) dated at 12,000 cal yr gave significantly different ¹⁴C ages at the 95% confidence level (Table 1), with the marine or atmospheric Δ^{14} C values of these 2 sub-samples varying by some 50%. Two slabs from a single coral specimen (DR 1183 and DR 1183[1]) were dated each at \sim 15,100 cal yr by the ²³⁰Th/U method and were sub-sampled for the ¹⁴C dating (Figure 7). While the 2 ²³⁰Th/U dates are very similar, the ¹⁴C dates by contrast are highly scattered (Figures 2, 5, 6; Table 1). The ¹⁴C ages of the sub-samples within each slab present the same range of variations from approximately 12,800 to ~13,400 yr, corresponding to marine Δ^{14} C values of $250 \pm 35\%$ to $160 \pm 55\%$ (2 σ), respectively. They are all within error (2 σ) of the marine and atmospheric INTCAL98 values (Stuiver et al. 1998) (Figures 5, 6). Only the upper Δ^{14} C values at \sim 310‰, corresponding to the youngest ¹⁴C ages, are consistent with those of the atmospheric ¹⁴C record from Lake Suigetsu (Kitagawa and van der Plicht 2000) (Figures 2, 4, 5). Similarly, at 12,000 cal yr, only the largest Δ^{14} C value (Figures 5, 6) agrees well with those of the ¹⁴C INTCAL98 calibration record (Stuiver et al. 1998) and with those from the Lake Suigetsu ¹⁴C record (Kitagawa and van der Plicht 2000). Therefore, the addition of modern carbon either during the different steps of AMS ¹⁴C dating procedure and/or during recrystallization processes with seawater seems unlikely. This is also attested by the very low content of magnesian (Mg) calcite in the corals. The initial 234 U/ 238 U ratios of all the corals analyzed are very similar to that of sub-modern corals at ~145‰ (in Delanghe et al. 2002) and within $\pm 5\%$ of the modern seawater values (Henderson et al. 1999; Delanghe et al. 2002) (Table 1). The detrital Th contents are very low. Thus, little post-depositional alteration or recrystallization of the primary coralline aragonite has taken place. In-situ dissolution processes alone may also be ruled out, as the U concentrations of these 2 corals, DR1183(2) and DR1183(1)/DR1183, are among the largest of the measured ones (Table 1). Therefore, the ²³⁰Th/U dates may be considered as most likely valid.

The anomalous ¹⁴C data at ~12,000 and ~15,000 cal yr would be the low Δ^{14} C values, and thus the old 14C ages when compared to the determined atmospheric ones from Lake Suigetsu (Kitagawa and van der Plicht 2000) (Figures 2, 6). This may be related to a subtle submarine diagenesis, which would include here dissolution of old carbonates and secondary Mg-calcite precipitation. Due to organic matter degradation, coral pore waters have a lower pH than that of open reef waters (Enmar et al. 2000) that may favor dissolution of carbonates. Secondary precipitation of aragonite and/or Mg-calcite micrites or needles of some 10 µm often fill voids in skeleton pores of sub-marine fossil corals (Enmar et al. 2000) (Figure 3). A 10% result in weight of secondary crystallization deriving from old ¹⁴C-depleted carbonates would account for the anomalously old ¹⁴C ages at 15,100 cal yr and 5% at 12,000 cal yr. Such a parallel contamination by "old" marine uranium with ²³⁴U/²³⁸U activity of 1.1, close to the value found in last interglacial corals at the time of deposition, would lead to a slight decrease of the $\delta^{234}U_{initial}$ from the value of 149 to ~144‰. This may explain the slightly lower δ^{234} U_{initial} values of these 2 corals with respect to those of the other corals, although the difference is not significant at the 95% confidence level (Table 1). However, this represents a very extreme case, and in the marine environment, the diagenetic changes in corals with recrystallization and secondary aragonite or Mg-calcite precipitation within seawater are usually related to increasing δ^{234} U_{initial} values (Bar-Matthews et al. 1993).

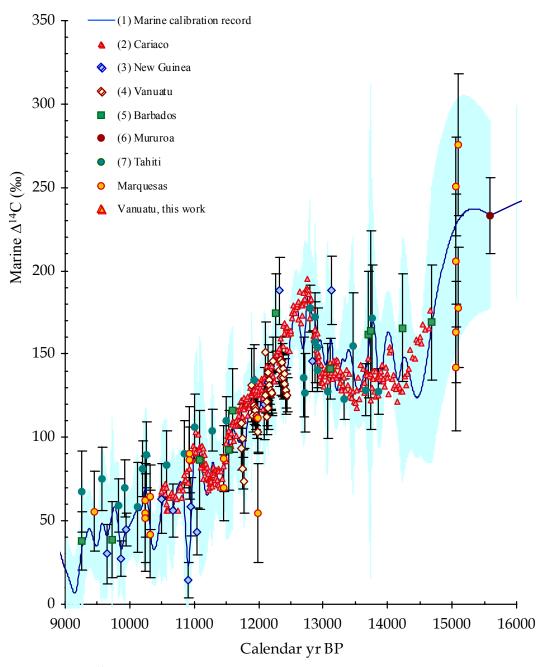


Figure 5 Marine Δ^{14} C changes at the 2- σ confidence level as a function of calendar ages. The blue envelope figured out the 2- σ error of the marine INTCAL98 Δ^{14} C values. (1) Stuiver et al 1998; (2) Hughen et al. 2000; (3) Edwards et al. 1993; (4) Burr et al. 1998; (5, 6, 7) Bard et al. 1993, 1998. Same references as Figure 2.

The corals dated at ~12,000 and ~15,100 cal yr were sampled in the same dredge (DR1183: $8^{\circ}45.5$ 'S; 140°03.8'W) and would reflect local and brief changes of the seawater chemical composition in the vicinity of Marquesas. Seasonal to annual variability of ¹⁴C ages, which can be

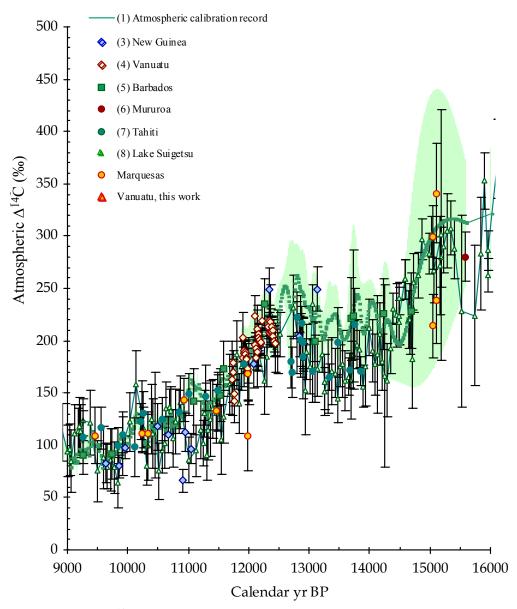


Figure 6 Atmospheric Δ^{14} C changes at the 2- σ confidence level as a function of calendar ages. The green envelope figured out the 2- σ error of the atmospheric INTCAL98 Δ^{14} C values. (8) Lake Suigetsu record (Kitagawa and van der Plicht 2000). Same references as Figure 2.

approached from our random sampling, was observed in modern banded coral samples from the Pacific Ocean (Brown et al. 1993; Druffel and Griffin 1993; Guilderson and Schrag 2001). Such variability was attributed to changes in the extent and intensity of the equatorial Pacific upwelling. However, this natural ¹⁴C variability did not exceed 10–15‰ (Druffel and Griffin 1993), which is significantly lower than the Δ^{14} C changes we observed in the fossil Marquesas corals. Thus, causes of the variation of the ¹⁴C ages in these corals remain a puzzling question.

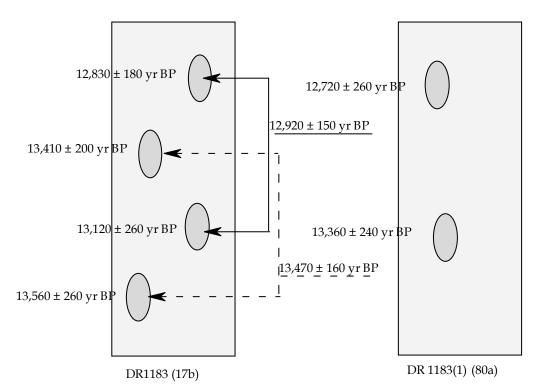


Figure 7 Sketch of 2 slabs, DR1183 and DR1183(1), of the same coral which was sub-sampled for AMS ¹⁴C dating. The weighted mean of the ¹⁴C ages and the weighted 2 σ from DR1183 (Table 1) are underlined.

CONCLUSION

Paired ¹⁴C and ²³⁰Th/U dating of corals from the Marquesas and Vanuatu provide additional estimates of the difference between the absolute and ¹⁴C ages of marine biogenic materials. These estimates are in agreement with those determined in previous studies between 3000 and 11,500 cal yr (Edwards et al. 1993; Stuiver et al. 1998; Bard et al. 1990a, 1993, 1998; Burr et al. 1998; Hughen et al. 2000; Kitagawa and van der Plicht 2000). Over this time interval, the reservoir age of the surface waters from the Marquesas differ by 100 yr from *R* values of corals from Tahiti. This is compatible with previous suggestions that variations of *R* exist in the Pacific Ocean (Stuiver et al. 1998; Goslar et al. 2002).

Among all the corals analyzed, 2 of them from the Marquesas dated at ~12,000 and ~15,000 cal yr present scattered ¹⁴C ages during the period of coral growth. Such variability may be related to rapid changes of the ¹⁴C content of the surface waters around the Marquesas or to a subtle submarine diagenesis. Additional coupled ¹⁴C and ²³⁰Th/U dating of corals from the Pacific should refine our knowledge of the extent and magnitude of these rapid changes in surface ¹⁴C in this region, and will be important in understanding the underlying mechanism responsible for such fluctuations.

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