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The radiocarbon signature of microorganisms in the mesopelagic ocean

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Several lines of evidence indicate that microorganisms in the meso- and bathypelagic ocean are metabolically active and respiring carbon. In addition, growing evidence suggests that archaea are fixing inorganic carbon in this environment. However, direct quantification of the contribution from deep ocean carbon sources to community production in the dark ocean remains a challenge. In this study, carbon flow through the microbial community at 2 depths in the mesopelagic zone of the North Pacific Subtropical Gyre was examined by exploiting the unique radiocarbon signatures ($\Delta^{14}\text{C}$) of the 3 major carbon sources in this environment. The radiocarbon content of nucleic acids, a biomarker for viable cells, isolated from size-fractionated particles (0.2–0.5 μm and >0.5 μm) showed the direct incorporation of carbon delivered by rapidly sinking particles. Most significantly, at the 2 mesopelagic depths examined (670 m and 915 m), carbon derived from *in situ* autotrophic fixation supported a significant fraction of the free-living microbial community (0.2–0.5 μm size fraction), but the contribution of chemoautotrophy varied markedly between the 2 depths. Results further showed that utilization of the ocean's largest reduced carbon reservoir, ^{14}C -depleted, dissolved organic carbon, was negligible in this environment. This isotopic portrait of carbon assimilation by the *in situ*, free-living microbial community, integrated over >50,000 L of seawater, implies that recent, photosynthetic carbon is not always the major carbon source supporting microbial community production in the mesopelagic realm.

microbial metabolism | particle flux | particulate organic carbon (POC) | chemoautotrophy

Marine microbes play a critical role in transforming inorganic and organic carbon in the ocean (1), and their interactions with the carbon reservoir, in its variety of phases, have important implications for the sequestration of CO_2 in either the deep ocean or sediments. However, the role that microorganisms play in the carbon cycle of the deep ocean, the mechanisms they employ, and the extent to which they employ them, are poorly understood. Most of the ocean's reduced carbon accumulates in the meso- and bathypelagic realm as dissolved organic carbon [(DOC); ≈ 700 Pg] (2), and here is where the fate of a significant fraction of organic carbon leaving the euphotic zone is decided (3). Therefore, microbial transformations of carbon in the “twilight zone” (defined here as depths between the base of the euphotic zone and 1,000 m) (3) play an essential role in determining the ocean's ability to sequester atmospheric CO_2 .

Below the euphotic zone, first light, and then labile organic matter, become increasingly scarce. Although DOC is present in this environment, it has long been assumed that this reservoir is unavailable to support heterotrophic production. For example, depth profiles of DOC (4), dissolved combined amino acids (5, 6), and dissolved carbohydrates (6, 7) appear to be static below 600 m, consistent with findings that DOC from the deep ocean resists microbial degradation (8). The most convincing evidence of the refractory nature of deep ocean DOC is its average

radiocarbon age, which lies between 4,000 and 6,500 radiocarbon years (4, 9, 10).

Despite the apparently static nature of the largest carbon reservoir available to support heterotrophic production in the meso- and bathypelagic zones, the abundance and activity of microorganisms are still measurable (11–14). One study suggests that 20–33 Pg C yr^{-1} are respired to CO_2 in the dark ocean (15), and the majority of this carbon is most likely derived from particulate organic carbon (POC) that sinks out of the euphotic zone. Measurements using particle traps have shown that this flux of POC is strongly attenuated in the mesopelagic ocean (3, 16); and a parameterization of this attenuation—the Martin Curve (16)—demonstrated that 90% of the organic carbon leaving the euphotic zone is lost from the particle phase within the upper 1,500 m.

Bacteria attached to sinking particles and marine snow not only respire carbon but also produce hydrolytic enzymes that transform POC into the dissolved phase at a rate faster than it is assimilated (17, 18). It is hypothesized that this decoupling of hydrolysis and uptake rates produces a “plume” of dissolved organic matter and other nutrients that trail these particles as they sink through the water column (19, 20). Therefore, in addition to respiration, carbon loss from POC may occur through DOC release as well. Strong correlations have been observed between deep ocean free-bacterial abundance and both episodic and long-term particle flux out of the overlying water column (21–23), further indicating that carbon is transferred from sinking POC to DOC to free-living microorganisms.

High-resolution depth profiles show that suspended POC contains bomb ^{14}C at abyssal depth (24) and support the hypothesis that rapidly sinking particles disaggregate to smaller size fractions. Additionally, several geochemical observations suggest that there is a “fresh” or new component within the mesopelagic DOC pool that is masked by the much larger refractory fraction identified above (25, 26), and sinking POC is the likely conduit for the rapid delivery of this new DOC to the deep ocean. Together these independent lines of evidence imply that some fraction of free-living heterotrophic microbes in the deep ocean is supported by inputs of recently-fixed carbon ultimately delivered by sinking particles.

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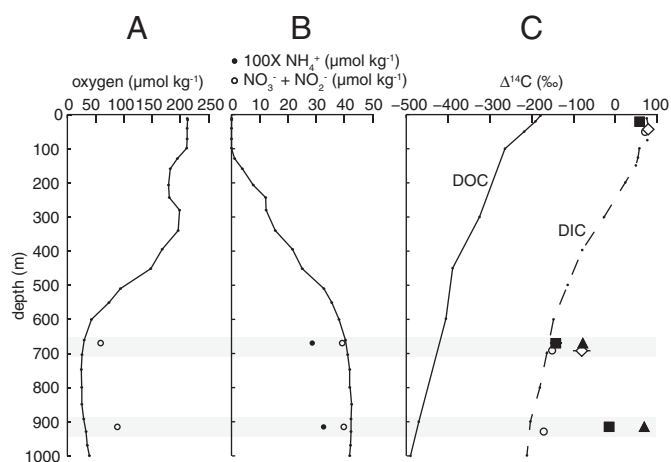


Fig. 1. Vertical profiles of oxygen (A) and nitrate + nitrite (B) data are for nearby station ALOHA (<http://hahana.soest.hawaii.edu>). Data (○) are specific to the NELHA sampling site (at 19.74°N, 156.05°W) (32, 41). (C) Corrected $\Delta^{14}\text{C}$ values for 0.2–0.5 μm (■) and >0.5- μm (▲) size fractions of DNA extracted from 21 m, 670 m, and 915 m (see *SI Text* for correction calculations), along with vertical profiles of $\Delta^{14}\text{C}$ of DOC (at 31°N, 159°W) (9) and DIC (at 35°N, 155°E) (40). $\Delta^{14}\text{C}$ -DNA error bars are propagated AMS errors and blank corrections, and if not visible, are less than the width of the marker. Also included are abundance-weighted averages and error of archaeal lipid $\Delta^{14}\text{C}$ from 21 m and 670 m (◇) (32).

indicative of aromatic groups not typically abundant in marine detritus, but found in DNA. However, NMR data could not conclusively rule out the presence of either intracellular biochemicals other than DNA or small amounts of ambient DOC in our precipitated samples. For these reasons we relied primarily on the measured C/N ratio of each DNA sample to estimate possible carbon contamination and apply the $\Delta^{14}\text{C}$ correction reported in Table 1.

A set of ^{14}C process blanks was also developed to demonstrate that $\Delta^{14}\text{C}$ -DNA was indeed reflective of available carbon sources. GF/F-filtered seawater collected off the Scripps Institution of Oceanography (SIO) pier was amended with 883 μM nitrate, 36.3 μM phosphate, and a carbon source with a known $\Delta^{14}\text{C}$ value (glucose, $\Delta^{14}\text{C} = +87 \pm 2\text{‰}$; or acetate, $\Delta^{14}\text{C} = -987 \pm 4\text{‰}$). These carbon sources were chosen to provide a range of $\Delta^{14}\text{C}$ values from modern ($\Delta^{14}\text{C} > 0\text{‰}$) to essentially radiocarbon dead ($\Delta^{14}\text{C} = -1,000\text{‰}$). The natural, marine microbial community was then incubated in the dark for 3 days until biomass was $\approx 10^9$ cells mL^{-1} and then subjected to the same filtration, lysis, and extraction method as all environmental samples. Isolated DNA was processed for $\Delta^{14}\text{C}$ and $\delta^{13}\text{C}$ analyses with the results shown in Table 2. ^{14}C sample sizes were comparable to the amount of carbon analyzed for environmental $\Delta^{14}\text{C}$ -DNA (Tables 1 and 2).

$\Delta^{14}\text{C}$ -DNA values, $+63 \pm 2\text{‰}$ (average \pm SD; $n = 3$) for the glucose culture and -821‰ for the acetate culture, showed rapid incorporation of the $\Delta^{14}\text{C}$ signature of carbon sources by

the microbial community. Measured $\Delta^{14}\text{C}$ -DNA values do deviate to different extents from the $\Delta^{14}\text{C}$ signature of each respective carbon source. This observed deviation could be caused by a combination of several factors: utilization of ambient DOC present in the collected seawater by cometabolism with the added glucose or acetate (42), selective utilization if the added carbon source was isotopically inhomogeneous, and contamination from either process chemicals or passive adsorption of ambient bulk DOC to filters during cell isolation as discussed previously. It is clear that no consistent carbon contamination from either a modern or ^{14}C -dead source is introduced through our extraction procedure. This finding was expected based on previous results showing negligible carbon contamination from reagents and filter materials. Nevertheless, contamination from an exogenous component, such as ambient DOC, with an intermediate $\Delta^{14}\text{C}$ value (between modern and ^{14}C -dead) could cause the observed deviation. However, source carbon and bacterial DNA had very similar $\delta^{13}\text{C}$ values after the incubation ($\Delta\delta^{13}\text{C} < \pm 1\text{‰}$; Table 2) and suggest that contamination from an exogenous source, if present, was relatively small (see *SI Text*).

Further evidence that the Natural Energy Laboratory of Hawaii Authority (NELHA; Kona, HI) samples primarily contain native DNA is their ability to be amplified by PCR. Because of the nature of the seawater supply system, we were motivated to analyze the microbial community to confirm that we were not isolating DNA from organisms that were unique to the seawater intake pipe. 16S rRNA bacterial and archaeal clone libraries establish that the microbes sampled are representative of those found in other mesopelagic marine environments (including nearby Station ALOHA) (Table S1 and Fig. S1), further confirming the presence of native DNA in our samples. No absolute trends in community composition based on depth or size fraction are apparent. The majority of archaeal clones from all depths and size fractions are Group I *Crenarchaeota* (>85%), with the remainder belonging to the Group II *Euryarchaeota*. Of note are a number of clones with 98–99% identity to the ammonia-oxidizing *N. maritimus* (33).

Significant differences were not observed for inorganic nitrogen between the 2 depths as $\text{NO}_3^- + \text{NO}_2^-$ concentrations were 39.6 $\mu\text{mol kg}^{-1}$ at 670 m and 40.1 $\mu\text{mol kg}^{-1}$ at 915 m (41). NH_4^+ concentrations were also comparable with values of 0.29 $\mu\text{mol kg}^{-1}$ and 0.33 $\mu\text{mol kg}^{-1}$ at 670 m and 915 m, respectively. There were measurable O_2 variations as the oxygen concentration at 670 m of 60.9 $\mu\text{mol kg}^{-1}$ was depleted relative to the 90.1 $\mu\text{mol kg}^{-1}$ at 915 m (Fig. 1) (41).

Discussion

Measurements of the ^{14}C content of microbial DNA directly demonstrate significant metabolic stratification in the mesopelagic water column. As expected, microorganisms in the surface ocean derived their carbon primarily from DIC and fresh DOC. In the mesopelagic realm, the free-living (0.2–0.5 μm) microbial population at 915 m incorporated carbon delivered primarily by sinking particles, but the same microbial size-fraction at 670 m relied more extensively on in situ chemoautotrophy to satisfy community carbon demand. Existing measurements do not

Table 2. Isotopic composition and sample sizes of carbon sources and microbial DNA from ^{14}C process blanks

Sample	Date, month/yr	Carbon source	Culture volume, L	$\Delta^{14}\text{C}$, ‰	Size, $\mu\text{g C}$	UCIG#	$\delta^{13}\text{C}$, ‰
glucose				$+87 \pm 2$	819	3411	-10.7
microbial DNA	10/03	glucose	20	$+63 \pm 5$	1015	1565	-11.7
microbial DNA	6/07	glucose	8	$+64 \pm 4, +62 \pm 4$	120, 200	38655, 38656	-11.7
acetate				$-988 \pm 4, -986 \pm 4$	800, 250	38653, 38654	-35.3
microbial DNA	6/07	acetate	8	-821 ± 4	110	39810	-34.5

predict this dynamic view of the deep “twilight zone” (between 670 m and 915 m). For example, the DOC reservoir is static between these depths (4–7), the geochemical environment at both depths is similar (41), the POC flux at each depth predicted by the parameterization of Buesseler et al. (3), is not consistent with the observed variability, and similar bacterial numbers are expected for both depths (43). Also significant is the demonstration that a fraction of the free-living microbial community had adapted to a metabolic lifestyle that was distinct from microorganisms found in larger size fractions, such as those associated with particles.

A primarily autotrophic microbial community in the surface ocean would have a $\Delta^{14}\text{C}$ signature that is identical to surface DIC ($+71 \pm 3\text{‰}$ at the time of sampling) (32). Archaeal lipids conformed to this expected trend—surface glycerol dialkyl glycerol tetraether (GDGT) lipids of the *Crenarchaeota* had an average $\Delta^{14}\text{C}$ value of $+82 \pm 12\text{‰}$ (32). Sterols from autotrophs and/or heterotrophs isolated in the $>0.5\text{-}\mu\text{m}$ fraction at this site had values of $+56 \pm 9\text{‰}$ and $+69 \pm 9\text{‰}$ (32). Additionally, sugars isolated from the DOC reservoir, a possible substrate for heterotrophic microorganisms, had modern $\Delta^{14}\text{C}$ values between 40 and 67‰ at this site in 2001 (25). Therefore, heterotrophs consuming fresh DOC, such as dissolved carbohydrates, and autotrophs should have similar $\Delta^{14}\text{C}$ -DNA values in the surface ocean. The surface DNA sample (0.2–0.5 μm) isolated in the present study included both archaeal and bacterial DNA (Table S1) and had a $\Delta^{14}\text{C}$ signature of $+60 \pm 3\text{‰}$. This value fell within the reported $\Delta^{14}\text{C}$ range for modern carbon sources confirming that some combination of DIC and recently produced organic carbon was fueling the surface free-living microbial community at the time of sampling. The only previous ^{14}C study of marine microbial DNA (0.2–0.7 μm) reported depleted $\Delta^{14}\text{C}$ values ($-34 \pm 24\text{‰}$) in surface waters of the eastern North Pacific Ocean (34°50' N, 123°00' W; Station M) (38) and suggested microbial incorporation of an old DOC component ($\Delta^{14}\text{C} = -549\text{‰}$). In contrast, DNA isolated from the surface microbial community at NELHA did not indicate utilization of any refractory DOC.

Utilization of refractory DOC in the mesopelagic zone also appeared to be negligible, and the relative importance of ambient DIC-fixation varied markedly with depth. The depleted $\Delta^{14}\text{C}$ -DNA value ($-140 \pm 17\text{‰}$) for the 0.2–0.5 μm size-fraction at 670 m showed that microbes are incorporating carbon with a $\Delta^{14}\text{C}$ value that is similar to in situ DIC ($-151 \pm 3\text{‰}$) (Fig. 1) (32), either directly through chemoautotrophy or through secondary production. The depleted $\Delta^{14}\text{C}$ -DNA value (Table 1) could have also resulted from consumption of a fraction of the ambient DOC pool (Fig. S2). However, archaeal lipids produced in situ at 670 m had an average $\Delta^{14}\text{C}$ value of -112‰ and showed that the archaeal community derived 83% of their carbon from autotrophy at this depth (32). This latter finding favors the hypothesis that the observed ^{14}C -depletion of total DNA results from chemoautotrophic production supplying the majority of carbon (as much as 95%) (Fig. S2) incorporated by the microbial community at this site and depth. The available phylogenetic data (Fig. S1 and Table S1) identified microorganisms other than chemoautotrophs in the 0.2–0.5 μm size-fraction from 670 m, so $\Delta^{14}\text{C}$ -DNA signatures must further reflect some heterotrophy coupled to in situ chemoautotrophy (i.e., both primary and secondary production). For example, dissolved neutral sugars isolated from high-molecular-weight ($>1,000$ Da) DOC at this site had an average $\Delta^{14}\text{C}$ value of $-123 \pm 10\text{‰}$ at 670 m (25), indicating that a major fraction of these carbohydrates could have been freshly produced in situ through chemoautotrophy. These sugars and other compounds produced in situ would provide a labile, ^{14}C -depleted carbon source for heterotrophs at 670 m. If heterotrophic free-living bacteria at this depth rely primarily on ^{14}C -depleted, fresh DOC produced from in situ autotrophy, then

both autotrophic and heterotrophic microorganisms would have a similar $\Delta^{14}\text{C}$ signature.

Surprisingly, DNA from the 670 m 0.2–0.5- μm sample was more depleted in ^{14}C than the same size fraction at 915 m (Fig. 1). At 915 m, DNA in the 0.2–0.5- μm size fraction contained more ^{14}C ($-15 \pm 15\text{‰}$) than DIC ($\Delta^{14}\text{C}$ -DIC = $-171 \pm 3\text{‰}$) indicating that sinking POC was more important for sustaining the free-living microbial population at this depth. Although some chemoautotrophy is occurring at 915 m, only up to 36% of DNA carbon in the 0.2–0.5- μm size fraction at 915 m ultimately could be derived from ambient DIC fixation as compared to up to 95% at 670-m depth (Fig. S2).

The $>0.5\text{-}\mu\text{m}$ size fractions at 670 m and 915 m followed a depth-trend similar to the smaller size fraction but had higher $\Delta^{14}\text{C}$ values (Fig. 1) than their free-living counterparts. Particles would be isolated into this larger size fraction and POC metabolism by particle-attached cells is the likely cause of these enriched $\Delta^{14}\text{C}$ -DNA values. As mentioned previously, particle sinking rates are high enough that little ^{14}C -decay occurs between surface waters and the mesopelagic zone, so POC and microbes that consume this POC will retain an enriched surface ocean $\Delta^{14}\text{C}$ signature relative to ambient C pools (i.e., DIC and total DOC) at mesopelagic depths. Therefore, DNA isolated from this larger size fraction should have $\Delta^{14}\text{C}$ values similar to DIC and DNA in surface waters. However, DNA in the $>0.5\text{-}\mu\text{m}$ fraction at 670 m was significantly depleted in ^{14}C relative to surface C pools, and this depletion reflects the coisolation of free-living microorganisms onto the $>0.5\text{-}\mu\text{m}$ filter. Larger filters such as GF/F filters (0.7- μm nominal pore size) have been shown to collect over 50% of marine bacteria (44, 45), and studies have observed that archaea are larger than bacteria in some marine environments (46). Therefore, $\Delta^{14}\text{C}$ -DNA values for the $>0.5\text{-}\mu\text{m}$ size fraction must integrate the ^{14}C signature of DNA from both free-living, mesopelagic microorganisms and those attached to particles sinking out of surface waters.

The greater ^{14}C -depletion of the $>0.5\text{-}\mu\text{m}$ sample at 670 m compared to 915 m indicates the coisolation of a free-living community that incorporates more in situ DIC at 670 m and is consistent with results obtained from smaller size fractions (Table 1 and Fig. 1). Together, data from the 2 different size-fractions demonstrate that some free-living microorganisms use carbon sources that are significantly different from those used by their particle-attached counterparts.

Spatial variations in the extent of autotrophy implied by the $\Delta^{14}\text{C}$ -DNA signature of both size fractions are consistent with the variability in archaeal *amoA* gene copies observed in other marine studies (35, 47–49). The latter suggest spatial gradients in the extent of archaeal ammonia oxidation and by extension, autotrophy. In an effort to explore the possible metabolisms that may have contributed to the assimilation of carbon present within the extracted DNA, quantitative PCR (qPCR) was used to determine the number of marine group I *Crenarchaeota* 16S rRNA and archaeal *amoA* gene copies present per ng total DNA in the same samples isolated for radiocarbon analyses. Although this analysis is only an indication of community composition and metabolic potential and not gene expression, a correlation has been shown between both marine *Crenarchaeota* 16S rRNA and archaeal *amoA* copy number and ammonia oxidation rates in the Gulf of California (48). qPCR assays with the 0.2–0.5 μm DNA samples showed comparable numbers of bacterial 16S rRNA genes at 670 m and 915 m, but more of both marine group I *Crenarchaeota* 16S rRNA and archaeal *amoA* gene copies per ng total DNA from 670 m than 915 m (Fig. 2). These trends are consistent with the $\Delta^{14}\text{C}$ data and indicate that DIC fixation fuels a greater proportion of free-living microbial community production at 670-m depth. In addition, a comparison of the 2 size fractions from 915 m showed greater abundance of both genes in the smaller size fraction (0.2–0.5 μm), again consistent with

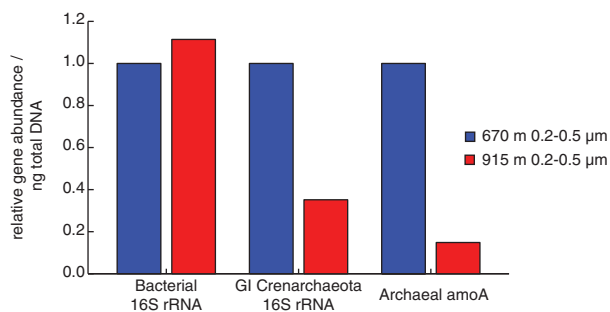


Fig. 2. qPCR data for 0.2–0.5- μ m-size fraction DNA samples showing relative abundances of bacterial 16S rRNA, group I *Crenarchaeota* 16S rRNA, and archaeal *amoA* gene copies per ng total DNA, as quantified by PicoGreen. For ease of sample comparison the qPCR data are normalized to the 670-m 0.2–0.5- μ m sample.

radiocarbon results. Although the focus here has been on archaeal autotrophy, carbon fixation by other members of the microbial community cannot be ruled out. For example, so far planktonic archaea appear to be primarily nitrosifiers, able to perform only the first step in the process of nitrification; therefore, the contribution to autotrophy by bacterial nitrifiers must also be considered (33, 47).

As discussed in previous studies (e.g., 32, 35), widespread chemoautotrophy in the deep ocean, if fueled primarily by ammonium oxidation, necessitates a starring role for regenerated, reduced N in the ocean's twilight zone. Therefore, the dominant free-living microbial metabolisms identified at both depths in this study (autotrophy, if supported primarily through nitrification, at 670 m, and heterotrophy of recently produced carbon at 915 m) require fresh organic carbon and nitrogen delivered presumably from sinking particles. The current study estimates that the majority of microbial carbon accumulating at 670 m is ultimately derived from chemoautotrophy. In the deep ocean several factors such as in situ microbial growth rates, and carbon fixation pathways and rates determine the magnitude of the reduced N requirement, and these parameters remain poorly constrained (29). Furthermore, particle flux to the deep ocean is unsatisfactorily defined (3) and may be inadequate to support carbon respiration rates (e.g., 15) or calculated archaeal nitrification rates in the mesopelagic realm. In fact, at nearby Station ALOHA bacterial carbon demand in the mesopelagic zone has been shown to exceed POC flux by 3- to 4-fold (43), and investigators have invoked vertically migrating zooplankton as one additional mechanism of carbon transport to mesopelagic depths at this site (43, 50). In situ carbon-fixation could also alleviate some of the heterotrophic carbon demand at these depths, although it remains enigmatic what process might supply the reductant. One possibility is a decoupling of dissolved nitrogen from dissolved carbon delivery to the mesopelagic zone. Nitrogen is a major waste product of vertically migrating zooplankton (e.g., 51), which could impact the nitrogen cycle at depth (52) and could mediate the depth-dependent requirement for excess reduced N, as would be necessary for intensive nitrification. The results of this study argue for a concerted focus on delineating the in situ microbial physiology of the deep ocean and constraining carbon and nitrogen fluxes into this environment.

Our approach provides a true in situ measurement with no tracer amendments or environmental manipulations and integrates production by the total microbial community in a significant volume of seawater ($\approx 50,000$ L) over approximately 1 week of sampling. This approach differs significantly from the snapshot view captured in other studies via discrete water column sampling, such as those that assess RNA gene expression. The

data show conclusively that source(s) of carbon used by the microbial community in the subsurface ocean can vary with depth (670 m vs. 915 m) and between free-living and particle-attached organisms (0.2–0.5 μ m fraction vs. >0.5 - μ m fraction). The direct demonstration of the importance of both rapid DOC delivery to the mesopelagic ocean and chemoautotrophy in maintaining total microbial production in this environment was only possible through the $\Delta^{14}\text{C}$ -DNA method developed here. This technique could provide a microbial perspective on the shape of the Martin Curve (16)—a relatively high-resolution depth profile of the $\Delta^{14}\text{C}$ signature of microbial DNA would demonstrate the importance of surface-derived carbon to microbes living in the mesopelagic zone and identify depths where significant respiration of vertical carbon flux occurs.

Methods

Sampling and DNA Extraction. Cellulose ester filters (0.5- μ m) (Millipore Opticap) plumbed in-line with 0.2- μ m polyethersulfone filters (Pall Supor) were used to collect particulate organic matter from approximately 50,000 L of surface (21 m), 670-m, and 915-m seawater by using the large-volume pumping capabilities at NELHA (Kona, HI) (19.74°N, 156.05°W). Filters were stored at -80°C until DNA was extracted. To isolate the nucleic acids, a method modified from Blair et al. (53) was used to minimize organic carbon contamination for ^{14}C analysis. Filters were cut open by using a combusted hacksaw, unfolded, and extracted in 1.5 M NaClO₄ at 4 $^\circ\text{C}$ for 48 h with shaking to lyse collected cells. The resulting 2-L solution was ultrafiltered (Pellicon MWCO 5,000 Da; Millipore) down to 20 mL and an organic extraction was performed with an equal volume of chloroform and isoamyl alcohol (24:1). Nucleic acids were precipitated from the resulting aqueous fraction with ethanol, pelleted by centrifugation, and re-dissolved in sterile water. All reagents used in the extraction of environmental DNA were the same and used in the same quantities as for the ^{14}C process blanks. Fractions were set aside for elemental and stable isotope analysis (University of California-Davis Stable Isotope Facility; Scripps Institution of Oceanography Analytical Facility), $^1\text{H-NMR}$ spectroscopy, and molecular biology procedures, and the majority of the sample was lyophilized in 9-mm quartz tubes, combusted to CO₂ at 850 $^\circ\text{C}$ with the addition of Ag and CuO, quantified, and converted to graphite according to the methods of Vogel et al. (54) for accelerator mass spectrometry (AMS) analysis at the University of California Irvine Keck Carbon Cycle AMS facility. $\Delta^{14}\text{C}$ values are reported as per geochemical samples according to Stuiver and Polach (55).

PCR, Cloning, and Sequencing. Portions of the 16S rRNA gene were amplified from environmental DNA samples by using Takara ExTaq Polymerase under the conditions described by the manufacturer's protocol. Universal bacterial (27F, 1492R) and archaeal (21F, 958R) primers were used at a final concentration of 0.5 μM . Products from 6 reactions were pooled, cleaned by using the Qiagen PCR Purification kit, and eluted in 30- μl water. Four microliters of this cleaned product was used in a TOPO cloning reaction with vector pCR2.1 as described by the manufacturer (Invitrogen). Plasmid DNA was extracted from white colonies by using a Qiagen Miniprep kit and was submitted to SeqXcel for sequencing. The 5' end of both strands of the bacterial 16S rRNA gene was sequenced with primer 536r and M13f/r, whereas M13f&r were used to sequence entire archaeal inserts. DNA sequences were assembled and edited by using VectorNTI (Invitrogen), checked for chimeras by using the Bellerophon server (56), aligned to the Silva database (57), grouped into operational taxonomic units by using DOTUR (58), and inserted into ARB for phylogenetic analysis (59). Sequences were deposited into GenBank under accession numbers EU817582–EU817654.

Quantitative PCR. For the qPCR assays, each reaction consisted of 12.5 μL Power SYBR[®] Green PCR Master Mix (Applied Biosystems), 0.4 μM each primer, 2 ng BSA, and 1–8 ng, 670-m and 915-m 0.2–0.5- μ m size fraction DNA as determined by PicoGreen (Molecular Probes) in a final volume of 25 μL . Archaeal *amoA* assays were run in triplicate by using primers and cycling parameters from Beman et al. (48), although run for 40 cycles with the marine group I *Crenarchaeota* assays similarly adapted from Mincer et al. (47). Bacterial 16S rRNA genes were quantified by using primers 932F and 1062R (60). PCR products generated by using the qPCR primers were used as standards in tenfold dilution series from 10⁷ to 10² copies based on PicoGreen quantification. PCR efficiencies and correlation coefficients for standard curves were as follows: 96% and $r^2 = 0.997$ for archaeal *amoA*; 82% and $r^2 = 0.994$ for GI *Crenarchaeota* 16S rRNA; and 94% and $r^2 = 0.999$ for bacterial 16S rRNA.

Results are presented as the relative abundance of gene copies per ng total DNA normalized to the 670-m 0.2–0.5- μ m DNA sample.

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