ORIGINAL ARTICLE

Diversity, abundance and expression of nitrite reductase (*nirK*)-like genes in marine thaumarchaea

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Ammonia-oxidizing archaea (AOA) are widespread and abundant in aquatic and terrestrial habitats and appear to have a significant impact on the global nitrogen cycle. Like the ammonia-oxidizing bacteria, AOA encode a gene homologous to copper-containing nitrite reductases (*nirK*), which has been studied very little to date. In this study, the diversity, abundance and expression of thaumarchaeal *nirK* genes from coastal and marine environments were investigated using two mutually excluding primer pairs, which amplify the *nirK* variants designated as AnirKa and AnirKb. Only the AnirKa variant could be detected in sediment samples from San Francisco Bay and these sequences grouped with the *nirK* from *Candidatus* Nitrosopumilus maritimus and *Candidatus* Nitrosoarchaeum limnia. The two *nirK* variants had contrasting distributions in the water column in Monterey Bay and the California Current. AnirKa was more abundant in the epi- to mesopelagic Monterey Bay water column, whereas AnirKb was more abundant in the meso- to bathypelagic California Current water. The abundance and community composition of AnirKb, but not AnirKa, followed that of thaumarchaeal *amoA*, suggesting that either AnirKa is not exclusively associated with AOA or that commonly used *amoA* primers may be missing a significant fraction of AOA diversity in the epipelagic. Interestingly, thaumarchaeal *nirK* was expressed 10–100-fold more than *amoA* in Monterey Bay. Overall, this study provides valuable new insights into the distribution, diversity, abundance and expression of this alternative molecular marker for AOA in the ocean.
The ISME Journal (2012) **6,** 1966–1977; doi:10.1038/ismej.2012.40; published online 17 May 2012
Subject Category: microbial ecology and functional diversity of natural habitats
Keywords: ammonia-oxidizing archaea; copper-containing nitrite reductase; Monterey Bay time series; *nirK*; thaumarchaea

Introduction

Ammonia-oxidizing bacteria (AOB) and ammonia-oxidizing archaea (AOA) have a central role in the global nitrogen cycle by oxidizing ammonia (NH_3) to nitrite (NO_2^-). In addition to the ammonia oxidation machinery (for example, ammonia monooxygenase), most ammonia oxidizers, interestingly, also encode a copper-containing nitrite reductase (*nirK*), which catalyzes the reduction of NO_2^- to nitric oxide (NO; Casciotti and Ward, 2001; Treusch *et al.*, 2005; Garbeva *et al.*, 2007; Cantera and Stein, 2007b; Bartossek *et al.*, 2010). In AOB, NirK is involved in production of the potent greenhouse gas nitrous oxide (N_2O) through 'nitrifier denitrification' (Dundee and Hopkins, 2001; Shaw *et al.*, 2006). AOA also produce N_2O, part of which appears to be formed through a reductive pathway (for example, nitrite reduction; Santoro *et al.*, 2011). Nitrite reduction mediated by nitrifiers contributes significantly

to N_2O release to the atmosphere from both terrestrial (Webster and Hopkins, 1996; Kool *et al.*, 2011) and marine (Dore *et al.*, 1998; Santoro *et al.*, 2011) ecosystems.

The physiological importance of *nirK* in ammonia oxidizers is still unclear. Classically, ammonia oxidizers are known to use oxygen (i) for the oxidation of NH_3 to hydroxylamine (NH_2OH) and (ii) as terminal electron acceptor in their electron transport chain. For oxidizing NH_3, dinitrogen tetroxide (N_2O_4), rather than O_2, has been proposed to be the primary oxidant, as described in the NO_x cycle hypothesis (Schmidt *et al.*, 2001; Kampschreur *et al.*, 2006; Schmidt, 2008) in which NO is an important intermediate: NO produced by NO_2^- reduction can feed into the NO_x cycle and the nitrite reductase is thereby indirectly involved in ammonia oxidation. This is supported by lower growth yield and growth defects (Schmidt *et al.*, 2004) and slower ammonia oxidation rate (Cantera and Stein, 2007a) in *Nitrosomonas europaea nirK* mutants. Furthermore, when NO is scavenged from the growth media, *N. eutropha* is unable to oxidize ammonia (Zart *et al.*, 2000), whereas ammonia oxidation activity increases when NO is added to the media (Zart and Bock, 1998).

AOB have long been known to substitute O_2 with NO_2^- as the electron acceptor in 'nitrifier denitrification' under O_2-limiting conditions (Abeliovich and Vonshak, 1992; Bock *et al.*, 1995; Schmidt and Bock, 1997). This has also been suggested for AOA (Walker *et al.*, 2010) but, unlike AOB, they only encode *nirK* and none of the downstream enzymes (for example, NO reductase) involved in classical denitrification (Hallam *et al.*, 2006; Walker *et al.*, 2010; Blainey *et al.*, 2011). It is unknown whether nitrite reduction in AOA is linked to energy conservation. Finally, nitrite reductase has been proposed to be involved in detoxification of nitrite in AOB cultures (Beaumont *et al.*, 2002; Cantera and Stein, 2007a). Nitrite detoxification has yet to be examined in AOA cultures, but AOA are generally more competitive in low-ammonia environments (Martens-Habbena *et al.*, 2009; Di *et al.*, 2010; Herrmann *et al.*, 2011) where nitrite concentrations are unlikely to reach toxic levels.

Copper-containing nitrite reductase genes homologous to *nirK* genes from AOB and denitrifying bacteria have been found in thaumarchaeal soil fosmids (Treusch *et al.*, 2005; Bartossek *et al.*, 2010), environmental metagenomes (for example, Venter *et al.*, 2004; Hollibaugh *et al.*, 2011) and in sequenced genomes of two AOA cultures, *Candidatus* Nitrosopumilus maritimus (two gene copies; Walker *et al.*, 2010) and *Candidatus* Nitrosoarchaeum limnia (Blainey *et al.*, 2011). Curiously, the archaeal sponge symbiont 'Cenarchaeum symbiosum' lacks the *nirK* homolog, but does harbor a phylogenetically related multicopper oxidase (Hallam *et al.*, 2006; Bartossek *et al.*, 2010). Thaumarchaeal *nirK* sequences derived from soil and marine environments form two highly divergent clusters that are just as distantly related to each other as to AOB *nirK* sequences (Bartossek *et al.*, 2010).

Although *nirK* is implicated in the production of the greenhouse gas, N_2O, our knowledge of the diversity of thaumarchaeal *nirK* in marine systems is limited to a few meta-genomic and -transcriptomic studies (for example, Venter *et al.*, 2004; Hollibaugh *et al.*, 2011) and a single clone library from 3480 m in the Tyrrhenian Sea (Yakimov *et al.*, 2011). To obtain a broader picture of the occurrence of thaumarchaeal *nirK* in marine systems, we designed specific primers and examined the diversity, abundance and expression of the gene in samples from Monterey Bay, the California Current and San Francisco Bay. The distribution of AOA *nirK* was also compared with the diversity and abundance of thaumarchaeal ammonia monooxygenase subunit A (*amoA*) genes, and with the abundance of the Marine Group I (MGI) 16S rRNA genes.

Materials and methods

Sampling
Samples were collected from Monterey Bay aboard R/V Point Lobos along an onshore–offshore transect with three oceanographic stations; C1, M1 and M2. Monterey Bay is a >1000 m deep, non-estuarine embayment in central California that opens broadly to the coastal ocean. The mid-bay station, M1, is located in the center of the upwelling plume (Service *et al.*, 1998), M2 is the outer-bay station and C1 is the inner-bay 'coastal' station (see Supplementary Figure S1). Upwelling season starts in March and continues through summer. Samples were collected at discrete depths down to 200 m on May 18, June 10 and December 2 of 2010, using a 12-bottle rosette sampler with a conductivity-temperature-depth device. Samples were collected within the central California Current on cruise S310 aboard McArthur II in July 2010 down to 3000 m water depth at station 60.90 (located about 300 km offshore, position 36.608/ − 125.778). San Francisco Bay sediments samples were collected, and DNA extracted, as described in Mosier and Francis (2010).

Water samples (1 l for DNA and 1–2 l for RNA) were pressure filtered through a 0.2-μM, 25-mm diameter Supor-200 membrane filter (Pall, Ann Arbor, MI, USA) with a 10-μM polyester pre-filter (GE Water & Process Technologies, Trevose, PA, USA), using a peristaltic pump. Filters for DNA and RNA extraction were transferred to 2-ml bead-beading tubes containing about 100 μl of 0.1-mm glass beads, and immediately frozen in liquid nitrogen. The Biological Oceanography Group at Monterey Bay Aquarium Research Institute kindly provided water chemistry data.

Nucleic acid extraction and cDNA synthesis
DNA was extracted from frozen filters following the protocol in Santoro *et al.* (2010), with a few modifications: 700 μl Sucrose-EDTA Lysis buffer (0.75 M sucrose, 20 mM EDTA, 400 mM NaCl and 50 mM Tris) were added and the filters were agitated for 45 s at speed 5.5 in a FastPrep bead-beating machine (MP Biomedicals, Solon, OH, USA). A volume of 100 μl 10% sodium dodecyl sulfate and proteinase K (50 mg ml^{-1} final concentration) were added and filters were incubated at 55 °C overnight. The lysate was purified using the Qiagen Blood & Tissue DNeasy kit (Valencia, CA, USA) following the manufacturer's protocol with an additional wash step with buffer AW2. Purified DNA was quantified using a Qubit fluorometer (Invitrogen, Grand Island, NY, USA).

RNA was extracted using the *mir*Vana miRNA isolation kit (Ambion, Grand Island, NY, USA). The initial lysis was carried out by adding 750 μl lysis/binding buffer to the filters with beads followed by vortexing at full speed for 5 min. The organic extraction and on-column RNA purification were performed according to the manufacturer's protocol. The RNA was finally eluted in 75 μl warm (95 °C) RNase-free water. Contaminating DNA was removed using the TURBO DNA-free kit (Ambion) following the manufacturer's protocol. cDNA was synthesized using the SuperScript III First-Strand Synthesis System for reverse transcription-PCR (Invitrogen) according to the manufacturer's protocol except for increasing the reverse transcription incubation step

to 4 h at 50 °C. Negative reverse transcription control reactions were performed for each sample replacing the reverse transcriptase enzyme with water.

Primer design

A total of 106 thaumarchaeal *nirK*-like sequences were retrieved from available marine metagenomes from CAMERA (Sun *et al.*, 2011) and NCBI using BLASTn and tBLASTx with *Ca.* Np. maritimus ORF 1259 (NC_010085) as query sequence. Sequences were aligned in Geneious v 5.4 (Drummond *et al.*, 2011) by translation alignment and primers were chosen by visual inspection of the alignment. Two mutually excluding primer pairs amplifying the same ~500 bp of the gene were necessary to target all of the currently known sequence diversity. The primer pair for AnirKa is AnirKa-58F 5′-ACB YTA TTC GGA AGY ACA TAC ACA-3′ and AnirKa-579R 5′-GYM ATT CCG TAC ATK CCG GA-3′. The primer pair for AnirKb is AnirKb-61F 5′-CTA TTC GGA RGT WCT TTY ACT GC-3′ and AnirKb-555R 5′-ACG TGT TGG TCC ATT GCT GC-3′. To determine whether a longer fragment of the gene would give higher phylogenetic resolution, an additional reverse primer (used together with AnirKa-58F) was also designed to yield a fragment of 1050 bp (AnirKa-1108R 5′-TRM TGC RTA TRC ACC TGG KTC-3′); all sequences generated with this primer are also targeted by the primer AnirKa-579R. The longer fragment did not result in higher phylogenetic resolution. The numbering of primers refers to the corresponding position in *Ca.* Np. maritimus ORF 1259 (NC_010085). The two primer pairs do not target any known soil-derived archaeal *nirK* sequences.

PCR, cloning and community analysis

The diversity of thaumarchaeal *nirK* was analyzed in eight water samples from Monterey Bay (stations M1 and M2 at 10, 40, 100 and 200 m water depth, May 2010), two water samples from California Current station 60.90 (30 and 3000 m) and four sediment samples from San Francisco Bay (BG20, BG30, BC11 and BA10). The diversity of thaumarchaeal *amoA* was analyzed in the same samples, except for Monterey Bay M2-10 m, which had too low abundance to be amplified for cloning, and M1-100 m. Diversity of thaumarchaeal *nirK* and *amoA* mRNA transcripts was analyzed in three water samples from Monterey Bay mid-bay station M1 at 10, 40 and 200 m from May 2010.

PCR amplification of AnirKa sequences was performed with the primers AnirKa-61F and AnirKa-579R, except for the DNA samples from Monterey Bay station M1 and San Francisco Bay, these were amplified with the reverse primer AnirKa-1108R instead. PCR amplification of AnirKb sequences was performed with the primers AnirKb-58F and AnirKb-555R. Both AnirKa and AnirKb sequences were PCR amplified in triplicate 25-µl reaction mixtures (12.5 µl Premix F (epicenter), 0.1 µM of each primer, 0.63 unit AmpliTaq DNA polymerase (Applied Biosystems, Carlsbad, CA, USA)) under the following thermal cycling conditions: 95 °C for 5 min, followed by 35 cycles of 94 °C for 30 s, 50 °C for 45 s, 72 °C for 1 min and finally 72 °C for 5 min. PCR amplification of AOA *amoA* was carried out with the primers Arch-amoAF and Arch-amoAR as described in Francis *et al.* (2005). Triplicate reactions were pooled, PCR purified (the QIAquick PCR purification Kit, Qiagen), cloned using the pGEM-T Easy Vector System (Promega, Madison, WI, USA) and sequenced (ABI 3100 Capillary Sequencer, ELIM Biopharm, Hayward, CA, USA).

Clone library coverage was assessed by calculating rarefaction curves and coverage in the Mothur software package (Schloss *et al.*, 2009; Table 1, Supplementary Table S1, and Supplementary

Table 1 Richness, diversity and coverage estimators calculated for clone libraries from all environments calculated at 97/95% OTU cutoff

Sample	Target	n	Shannon (H)	Simpson (D)	Chao	ACE	Coverage (%)
Monterey Bay: M1	AnirKa	78	1.9/1.1	0.22/0.56	16/11	18/15	95/96
	AnirKb	69	3.4/2.9	0.04/0.06	75/56	92/45	59/80
	amoA	35	2.1/1.6	0.16/0.30	16/9	19/13	83/94
Monterey Bay: M2	AnirKa	67	2.5/1.8	0.12/0.25	31/11	92/12	82/97
	AnirKb	66	3.2/2.5	0.04/0.10	57/19	110/21	70/94
	amoA	42	2.2/2.0	0.14/0.17	22/14	36/15	81/90
Monterey Bay: M1 cDNA	AnirKa	60	2.4/1.9	0.10/0.20	16/10	17/11	95/97
	AnirKb	49	3.3/2.9	0.03/0.05	122/36	218/38	45/76
	amoA	60	2.4/1.4	0.10/0.36	14/8	15/9	97/97
California current 30 m	AnirKa	16	1.4/1.0	0.22/0.38	6/5	6/8	89/89
	AnirKb	13	0.9	0.40	3	4	92
	amoA	7	0.0	1.00	1	1	100
California current 3000 m	AnirKa	16	1.0	0.48	4	5	94
	AnirKb	15	1.4	0.22	5	6	93
	amoA	12	1.9/1.8	0.09/0.11	23/10	21/11	50/67
San Francisco Bay	AnirKa	62	2.6/2.4	0.11/0.12	28/17	33/17	84/95
	amoA	44	2.5/2.3	0.10/0.12	36/22	70/42	73/82

Figure S2). Diversity indices (Shannon, Simpson) and richness estimators (Chao, ACE) were calculated in Mothur (Schloss *et al*., 2009; Table 1). Calculations were based on 477, 450 and 594 bp for AnirKa, AnirKb and AOA *amoA*, respectively. The archaeal *nirK* nucleotide sequences were aligned using translation alignment in Geneious

(Drummond *et al*., 2011) and trimmed to the length of the shortest sequence. A consensus tree based on 453 bp was constructed from maximum likelihood (PhyML (Guindon and Gascuel, 2003) using general time-reversible substitution model) and neighbor-joining (Jukes-Cantor distance correction), both with 1000 bootstrap replicates (Figure 1). Representative
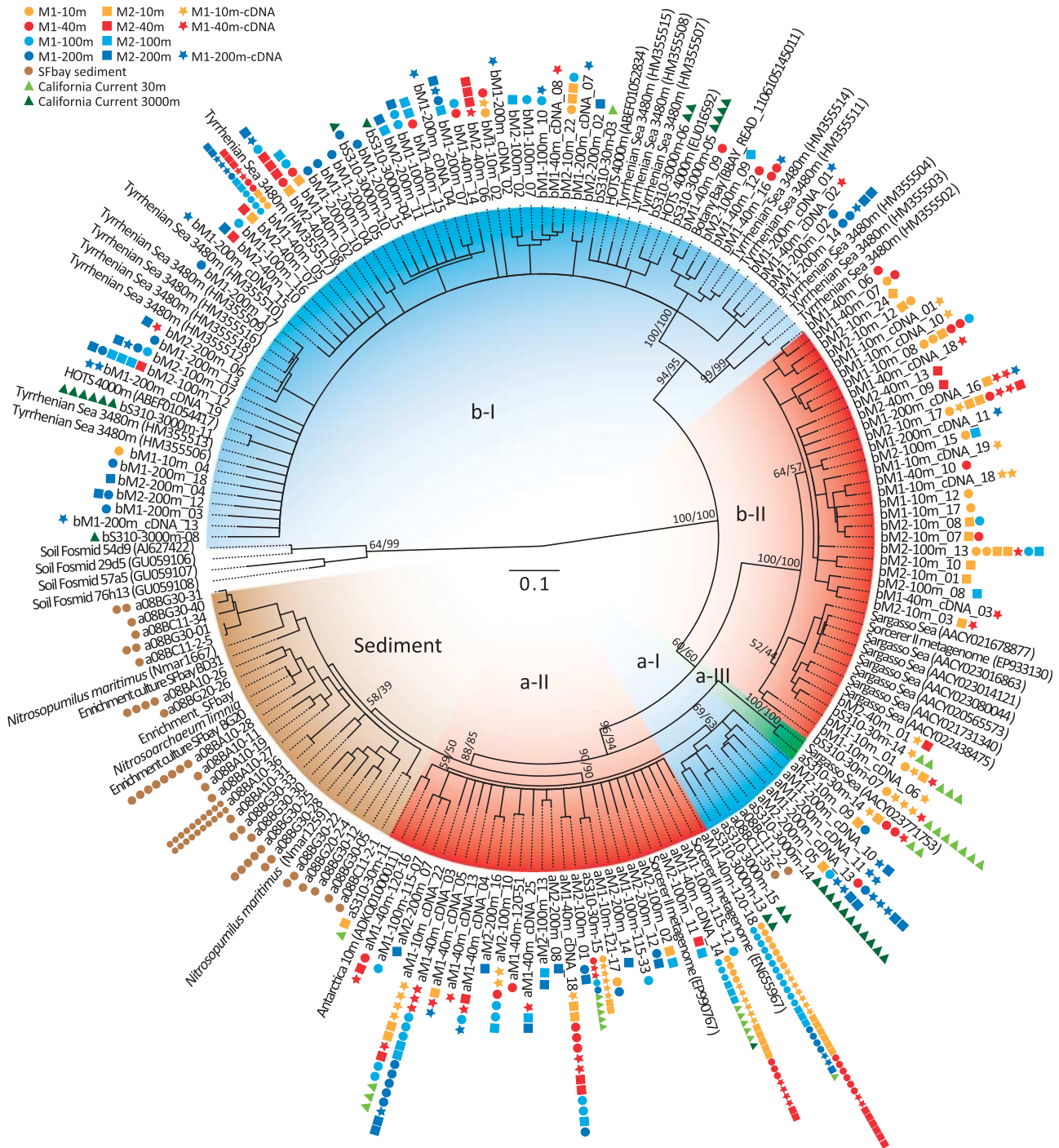
Figure 1 Phylogeny of marine thaumarchaeal *nirK* sequences. Maximum likelihood (ML; PhyML, general time-reversible (GTR)+Γ) 30% cutoff consensus tree with 1000 bootstrap replicates, based on 453 bp. Major groupings were all supported by neighbor-joining (NJ, Jukes-Cantor distance correction, 1000 bootstraps). Numbers on branches are bootstrap values from ML/NJ. Only sequence representatives from a 97% similarity OTU cutoff are shown in tree. Colored symbols show total number of clones represented by a sequence (see legend). Sequence name prefix 'a/b' refers to which primer pair the sequence was amplified with (AnirKa or AnirKb).
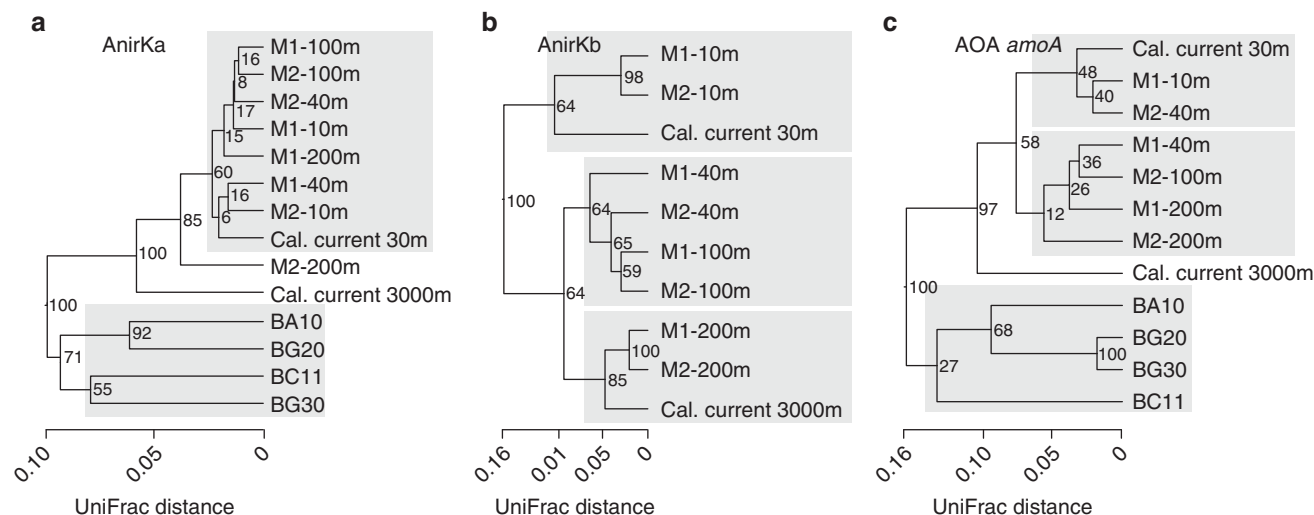
Figure 2 Dendrograms of hierarchical clustering of community similarities based on weighted UniFrac Jackknife Cluster Analysis with 100 permutations. (**a**) AnirKa sequences, (**b**) AnirKb sequences and (**c**) AOA *amoA* sequences. Numbers on branches are Jackknife values.

sequences from an operational taxonomic unit (OTU) cutoff of 97% similarity are shown in the tree along with thaumarchaeal *nirK* sequences retrieved from NCBI and CAMERA (Sun *et al.*, 2011) covering the same region of the gene. Thaumarchaeal *nirK* genes from soil fosmids were included as outgroups (Figure 1). AOA *amoA* sequences were aligned with GenBank sequences in ARB (Ludwig *et al.*, 2004), and a nucleotide-based phylogenetic tree was constructed by neighbor-joining with Jukes-Cantor distance correction (Supplementary Figure S3).

Community composition of the different samples was compared using weighted UniFrac Jackknife Environment clustering with 100 permutations (Figure 2, Supplementary Figure S5). UniFrac is a non-OTU-based metric that compares the relatedness of different environments using a phylogenetic tree as input. The trees used in the analyses were constructed separately for AnirKa, AnirKb and AOA *amoA* by maximum likelihood (PhyML, general time-reversible) from 480, 462 and 597 nucleotide positions, respectively, using a soil fosmid *nirK* sequence (AJ627422) as the outgroup.

GenBank accession numbers are as follows: AnirKa, JN257263–JN257407, JN257543–JN257574, JN257603–JN257664 and JN620225–JN620284; AnirKb, JN257408–JN257542, JN257575–JN257602 and JN620285–JN620333; and AOA *amoA*, JN375838–JN375977 and JN624325–JN624384.

Quantitative PCR
The abundance of thaumarchaeal *nirK*, *amoA* and MGI 16S rRNA genes was estimated by quantitative PCR in depth profiles from Monterey Bay station C1, M1 and M2 from May, August and December 2010 (Figure 3). The abundance of thaumarchaeal *nirK* and *amoA* was also estimated within a single depth profile from California Current station 60.90 (Supplementary Figure S6).

Abundance of mRNA transcripts of thaumarchaeal *nirK* and *amoA* was estimated in samples from Monterey Bay station C1, M1 and M2 collected in May 2010 (Figure 6).

The two different primer sets, AnirKa and AnirKb, were used to estimate the abundance of thaumarchaeal *nirK*. Both assays used the following reaction chemistry: 12.5 µl iTaq SYBR Green Supermix with ROX (Bio-Rad, Hercules, CA, USA), 400 nM of each primer and 0.3–37.3 ng template DNA or cDNA in a final volume of 25 µl. Thermal cycling conditions were as follows: 95 °C for 3 min, 35 cycles of 95 °C for 30 s, 50 °C for 45 s, 72 °C for 45 s and a detection step at 78 °C; finally, a melting curve was run from 65 to 90 °C with 0.3 °C increments.

The abundance of AOA *amoA* was estimated using the primers Arch-amoAF and Arch-amoAR (Francis *et al.*, 2005) with the same reaction chemistry as used for nirK; the cycling conditions were as follows: 95 °C for 3 min, 35 cycles of 95 °C for 30 s, 56 °C for 45 s, 72 °C for 45 s and a detection step at 77 °C, finalized by a melting curve from 65 to 90 °C with 0.3 °C increments. The abundance of MGI 16S rRNA genes (thought to represent most marine AOA) was estimated using a TaqMan assay; the primers were GI_751F and GI_956R (Mincer *et al.*, 2007) and a probe was designed for this study based on SILVA release 104 (Pruesse *et al.*, 2007): MGI_889 (6-FAM-5′-AGT ACG TAC GCA AGT ATG AA-3′). The assay was run with the following reaction chemistry: 12.5-µl TaqMan Environmental Master Mix 2.0 (Applied Biosystems), 200 nM of each primer, 300 nM of probe and 0.5–12.4 ng of DNA template. Cycling conditions were as follows: 95 °C for 10 min, 40 cycles of 95 °C for 15 s and 56 °C for 1 min followed by detection. A clone library was generated with the primers GI_751F and GI_956R to obtain a plasmid standard.

All quantitative PCR reactions were run in triplicate using a StepOnePlus Real-Time PCR
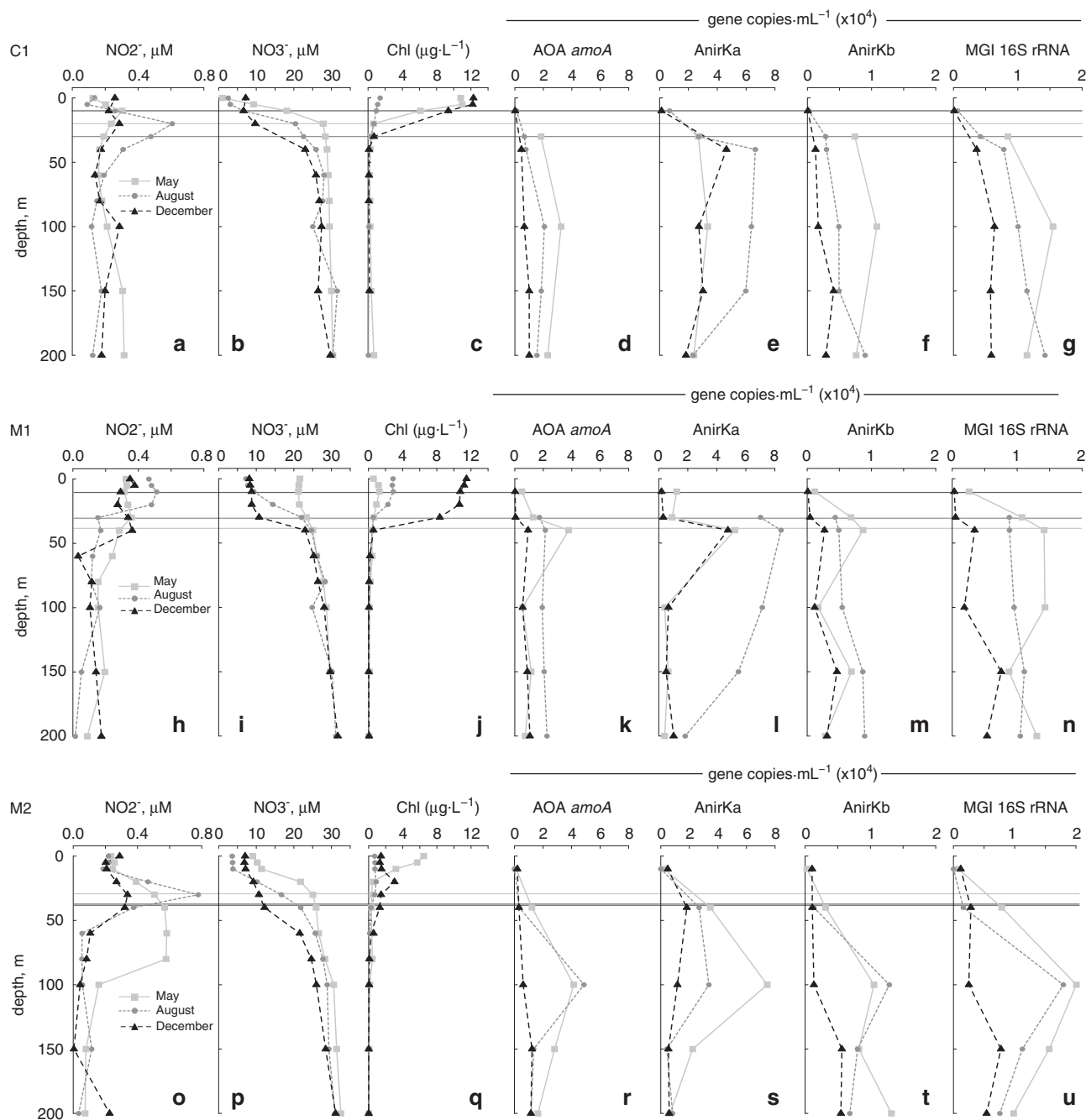
Figure 3 Water column profiles from Monterey Bay stations C1, M1 and M2 from May, August and December 2010. Profiles show NO_2^-, NO_3^-, chlorophyll and gene copy abundances of AOA *amoA*, AnirKa, AnirKb and MGI 16S rRNA. Horizontal lines show the approximate depth of light penetration (1% of surface irradiance) in the different months.

System (Applied Biosystems). Standard curves ranging from 5 to 1×10^6 copies per reaction were generated from purified, linearized plasmids obtained from the clone libraries described above. If the s.d. of a sample exceeded 10% of average value, one of the triplicates was omitted or the sample was re-run. Assay efficiencies were 73–78% for AnirKa, 85–88% for AnirKb, 79–90% for AOA *amoA* and 95–96% for MGI 16S rRNA genes.

All assays produced consistent and reproducible results.

Statistical analysis
Spearman's rank-order correlation was used to test the correlations between gene copy numbers (Figure 4); a non-parametric method was used because the gene copy abundance data did not
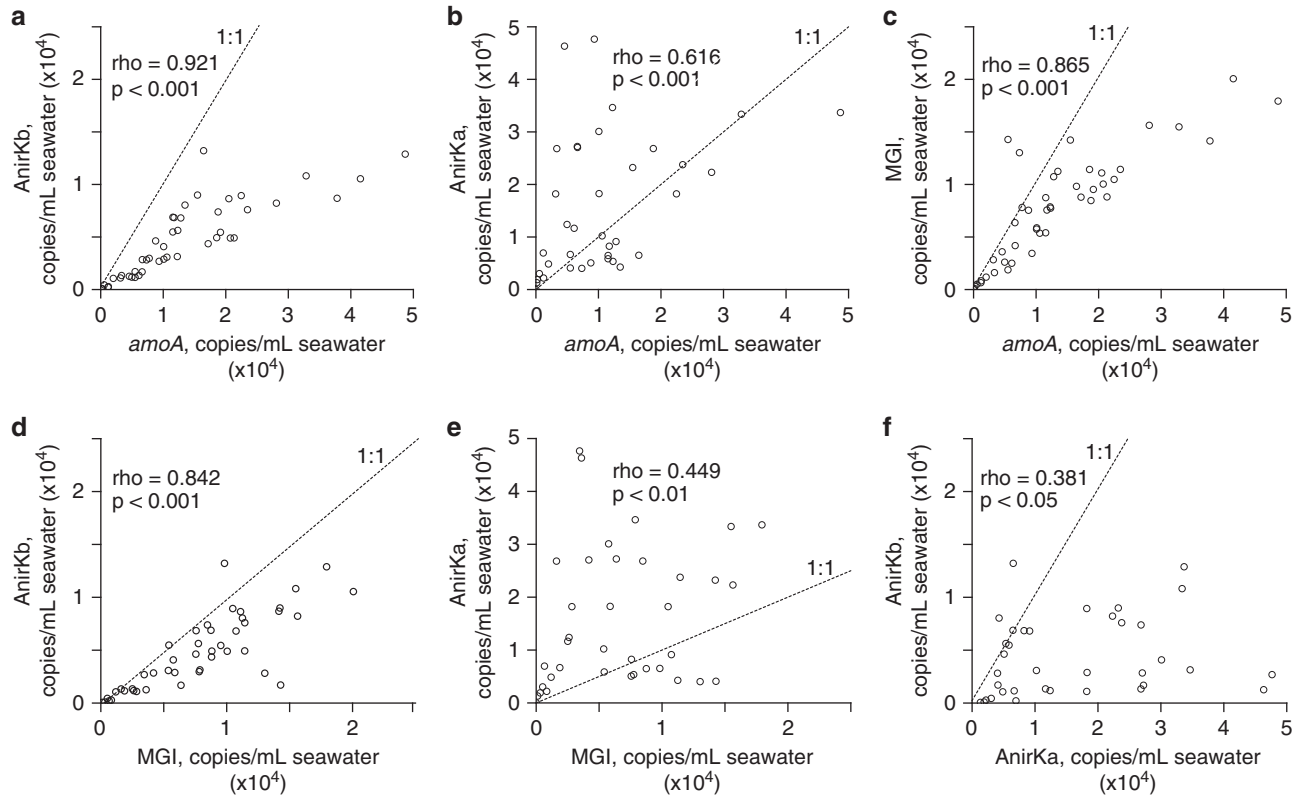
Figure 4 Correlations of gene copy abundances of AOA *amoA*, AnirKa, AnirKb and MGI 16S rRNA from Monterey Bay (**a–f**). Spearman's rank-order correlations (rho) and significance levels are shown in each plot. Dotted lines show the 1:1 relationships.
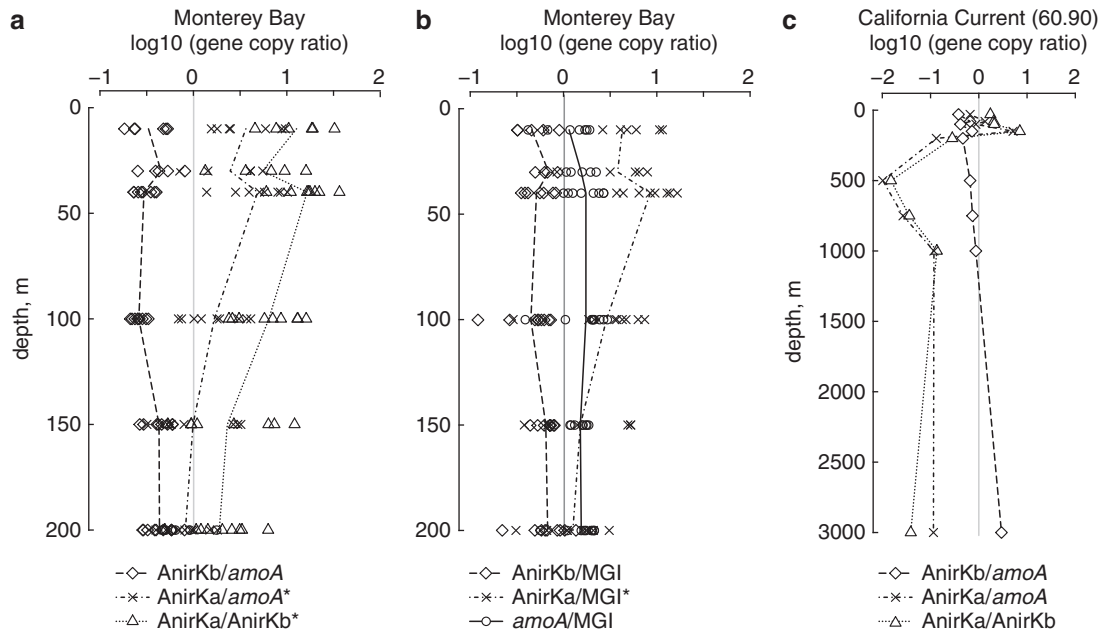
Figure 5 Water column profiles of gene copy ratios from (**a**, **b**) Monterey Bay and (**c**) California Current station 60.90. Vertical gray lines show the 1:1 relationships. For Monterey Bay, every data point is plotted and the mean values are connected with lines. *Average gene copy ratio changes with water depth (one-way ANOVA, $P < 0.001$).

follow a normal distribution. Correlations between environmental parameters (NO_2^-, NO_3^-, water depth, temperature, salinity and chlorophyll) and gene copy abundances were also calculated using non-parametric Spearman's rank-order correlation (Supplementary Table S3). Change in average gene copy ratio with water depth (Figure 5a) was tested with one-way analysis of variance (ANOVA).
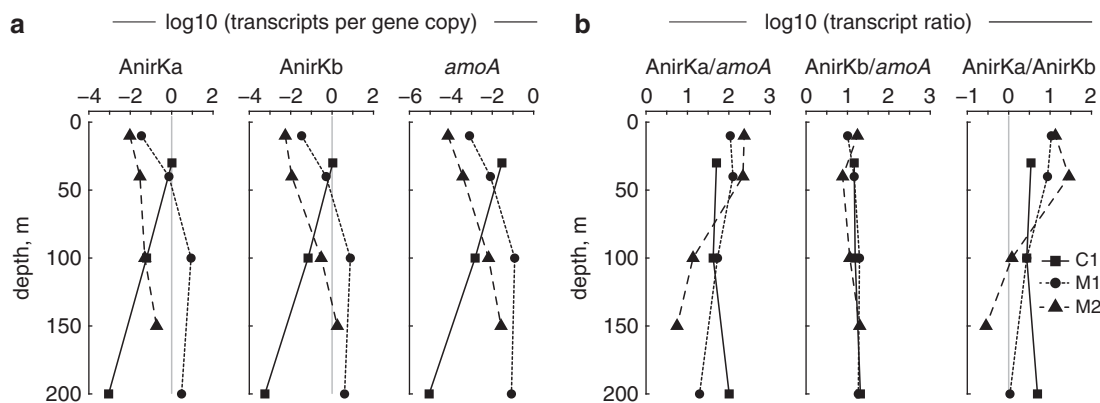
Figure 6 Transcript abundances from Monterey Bay stations C1, M1 and M2 from May 2010. (**a**) Ratios of transcript copies (cDNA) to gene copies (DNA) for AnirKa, AnirKb and AOA *amoA*. (**b**) Ratios of transcript copies (cDNA/cDNA) of AnirKa/*amoA*, AnirKb/*amoA* and AnirKa/AnirKb. Vertical gray lines show 1:1 relationships.

All calculations were performed in R Commander (Fox, 2005).

Results

Community composition of thaumarchaeal *nirK* and *amoA*

The phylogeny of thaumarchaeal *nirK* sequences is shown in Figure 1. To target the known sequence diversity in marine environments, two mutually excluding primer sets were designed and used, and the derived *nirK* sequence types will be referred to as AnirKa and AnirKb throughout. AnirKa sequences form four major clusters with an average nucleotide-based identity of 83%. Group a-I, a-II and a-III contain water column samples from Monterey Bay and the California Current; group a-I is primarily comprised of sequences from below 200 m depth. Sediment-derived *nirK* sequences from San Francisco Bay form a separate cluster along with *nirK* from the cultured AOA representatives *Ca.* Np. maritimus (two gene copies) and *Ca.* Na. limnia, the latter of which was originally isolated from San Francisco Bay sediments (Blainey *et al.*, 2011). AnirKb sequences form two major clusters with an average nucleotide-based identity of 74%. Group b-I primarily contains sequences from below 100 m depth (meso- to bathypelagic zone) and group b-II primarily contains sequences from above 100 m depth (epi- to mesopelagic zone; Figure 1, Supplementary Figure S4). Most of the metagenomic sequences from the Sargasso Sea shotgun library (Venter *et al.*, 2004) and the Sorcerer II expedition (Yooseph *et al.*, 2007) fall in group b-II, whereas metagenomic and fosmid sequences from Hawaii Ocean Time series (Konstantinidis and DeLong, 2008) and all of the sequences from 3480 m water depth in the Tyrrhenian Sea (Yakimov *et al.*, 2011) fall in group b-I; however, it should be noted that the primers used in the latter study do not target all sequences amplified by the AnirKa primers.

The phylogeny of thaumarchaeal *amoA* is shown in Supplementary Figure S3; the sequences from the San Francisco Bay sediments group with other low-salinity sediment samples and with soil/sediment samples. The AOA *amoA* sequences from Monterey Bay fall primarily in the marine 'shallow' group A, although the proportion of sequences associated with the 'deep' group B increases with water depth. All AOA *amoA* sequences from 30 m in the California Current fall within group A and all samples from 3000 m fall within group B (Supplementary Figures S3, S4).

In Monterey Bay, the diversity (estimated by Shannon and Simpson diversity indices) of AnirKb is much greater than both AnirKa and AOA *amoA* (Table 1), and the OTU richness (estimated by Chao and ACE) is two to six times higher at a 95% similarity cutoff. In the California Current, the overall diversity is lower than in Monterey Bay. In San Francisco Bay sediment, the diversity and richness of AnirKa and AOA *amoA* are similar.

Similarities in community composition between the different samples were analyzed by weighted UniFrac cluster analysis (Figure 2). The sediment samples from San Francisco Bay have a distinct community based on AnirKa and AOA *amoA* (Figure 2a and c). The water column samples from Monterey Bay and California Current show a clear clustering with water depth for AnirKb, forming three groups with samples from 10 and 30 m, 40 and 100 m and 200 and 3000 m water depths, respectively. Environment clustering with water depth is also evident for AOA *amoA*, with two distinct depth groupings: 10, 30 and 40 m, and 40, 100 and 200 m; in contrast, the 3000-m sample from the California Current is distinct from the remaining water column samples. For AnirKa, there is no clear environment clustering with water depth, except for the samples from California Current (3000 m) and M2-200 m, both of which are distinct from the remaining water column samples.

Clone libraries of thaumarchaeal *nirK* and *amoA* mRNA transcripts from Monterey Bay Station M1 (10, 40 and 200 m) showed similar diversity and OTU-based richness as the corresponding DNA-based clone libraries (Table 1), except for AnirKb transcript richness, which was higher. For AnirKa and AOA *amoA*, the gene- and transcript-based communities from all water depths are very similar, as indicated by the overall low UniFrac distance of about 0.05 (Supplementary Figure S5). For AnirKb, there is clustering of the DNA- and cDNA-based communities: the corresponding libraries from M1-200 m form one cluster and the corresponding libraries from M1-10 and 40 m form another cluster (Supplementary Figure S5).

Thaumarchaeal gene copy abundances
The abundance of thaumarchaeal *nirK*, *amoA* and MGI 16S rRNA gene sequences was estimated in depth profiles in Monterey Bay from 10 to 200 m. All four assays showed low gene abundance at 10 m and a clear increase in copy numbers below the photic zone (Figures 3d–g, k–n and r–u). At the inner- and mid-bay stations (C1 and M1), AOA abundance peaked at ~40 m, but peaked deeper in the water column at the outer-bay station, M2. Environmental parameters (NO_3^-, depth, temperature, salinity and chlorophyll, all correlating with water depth) showed clear Spearman's rank correlations with AnirKb, AOA *amoA* and MGI 16S rRNA gene copy abundances, but not with AnirKa abundance (Supplementary Table S3).

The overall abundances of both AOA *amoA* and MGI 16S rRNA genes correlate strongly with AnirKb abundance (Spearman's correlations of 0.921 and 0.842, respectively), whereas the correlation with AnirKa is weak (Spearman's correlations of 0.616 and 0.449, respectively; Figures 4a, b, d and e). There is a strong correlation between MGI 16S rRNA and AOA *amoA* gene abundances (rho = 0.865, Figure 4c), and a weak correlation between AnirKb and AnirKa abundances (rho = 0.381, Figure 4f).

Gene copy ratios reveal different distributions of AnirKb and AnirKa in the water column (Figure 5a): AnirKb is less abundant than both AOA *amoA* and MGI 16S rRNA genes with average gene copy ratios of 0.4 and 0.6, respectively, and the ratios do not change with water depth (one-way ANOVA, $P > 0.05$; Figures 5a and b). In contrast, the gene copy ratios of AnirKa to AOA *amoA* and MGI 16S rRNA genes vary significantly with water depth (one-way ANOVA, $P < 0.001$), with AnirKa being five and sevenfold more abundant than AOA *amoA* and MGI 16S rRNA genes, respectively, in the upper 40 m of the water column and approaching ratios of 0.5 and 2 toward 150–200 m depth (Figures 5a and b). In addition, AnirKa was relatively more abundant closer to the coast, with a significantly higher ratio of AnirKa to AnirKb at the inner Monterey Bay station C1 than at station M1, M2 and California Current station 60.90 (Supplementary Figure S7).

The ratio of AnirKa to AnirKb copy numbers also varies strongly with depth (one-way ANOVA, $P < 0.001$), with AnirKa outnumbering AnirKb 15-fold in the upper 40 m of the water column, dropping to threefold as abundant toward 200 m depth (Figure 5a). AOA *amoA* is on average 1.7 times more abundant than MGI 16S rRNA genes and the gene ratio shows no variation with water depth from 10 to 200 m (one-way ANOVA, $P > 0.05$). A ratio of 1 has previously been reported from station M1 in Monterey Bay (Mincer *et al.*, 2007), whereas a ratio >1 has been found in a number of oceanic regions, including the Gulf of California (Beman *et al.*, 2008), the California Current (Santoro *et al.*, 2010), and the epipelagic of the central Pacific Ocean (Church *et al.*, 2010) and South China Sea (Hu *et al.*, 2011).

The patterns in gene copy ratios with water depth in Monterey Bay are also found in a depth profile from California Current station 60.90 from 30 to 3000 m (Figure 5c): AnirKb is less abundant than AOA *amoA* throughout the water column (except at 3000 m) and the ratio varies very little with water depth. In contrast, the ratio of AnirKa to AOA *amoA* fluctuates greatly with water depth, with an ~1:1 relationship in the upper 150 m of the water column and a decrease to 10–100 times less AnirKa than AOA *amoA* below 200 m.

Thaumarchaeal nirK and amoA transcript abundances
The abundance of mRNA transcripts of thaumarchaeal *nirK* and *amoA* was examined in samples from Monterey Bay Stations C1, M1 and M2 (May 2010). The mRNA abundance, in particular for AOA *amoA*, was low compared with the corresponding gene copy abundance (Figure 6a). AnirKb transcripts were ~15 times more abundant than AOA *amoA* at all stations throughout the water column (Figure 6b). AnirKa was on average 88 times more expressed than AOA *amoA*, and the ratio changed with water depth: at the outer-bay station M2, and at the mid-bay station M1, AnirKa was more expressed in the upper 40 m of the water column (108–235 times) than at 100–200 m depth (6–53 times; Figure 6b). The relative abundance of AnirKa to AnirKb was also higher in the upper 40 m (3–29 times) than at 100–200 m depth (0.3–7 times) at station M1 and M2.

Discussion

Depth distribution of thaumarchaeal nirK genes in the water column
The two thaumarchaeal *nirK* variants, AnirKa and AnirKb, have contrasting distributions in the marine water column, in terms of community composition, relative gene abundance, as well as gene expression. AnirKa is most abundant in the epi- to mesopelagic, and possibly coastal waters (Figures 5a and c, Supplementary Figure S7). The community

composition of AnirKa shows little change with water depth (Figure 2a, Supplementary Figure S4): there is one dominant AnirKa phylotype in the epi- to mesopelagic (group a-II) and one phylotype (group a-I) comprised mostly of deep-water sequences from 3000 m in the California Current and 200 m at M2 in Monterey Bay (Figure 2a).

Both the abundance and community composition of AnirKb are highly similar to that of AOA *amoA* (Figures 2 and 5). The AnirKb community shifts from primarily group b-II in the upper 40 m of the water column to group b-I dominated below 40 m (Figures 1 and 2b, Supplementary Figure S4). A similar community shift is evident for AOA *amoA* where the proportion of sequences associated with the 'deep' group B increases with water depth (Figure 2c, Supplementary Figures S3, S4). This follows the general distribution of thaumarchaeal *amoA* ecotypes in which 'shallow' group A sequences are found throughout the water column, whereas group B sequences are more abundant in deeper water (Francis *et al.*, 2005; Hallam *et al.*, 2006; Beman *et al.*, 2008; Santoro *et al.*, 2010). The relative abundance of mRNA transcripts supports the overall picture that AnirKa is more abundant in epipelagic waters, whereas AnirKb is consistently tracking the distribution of AOA *amoA* genes (Figure 6b).

The divergent distributions of AnirKa and *amoA* suggest that the underlying mechanisms controlling the abundance of the AnirKa variant may be different from those governing the distribution of the traditional thaumarchaeal molecular marker gene, *amoA*. This incongruity implies that either (i) the primers targeting thaumarchaeal *amoA* are missing substantial diversity in the epipelagic or (ii) the AnirKa gene variant is not specific to thaumarchaeal ammonia oxidizers. The *nirK* gene from both *Ca*. Np. maritimus and *Ca*. Na. limnia group with the sediment-associated AnirKa sequences (Figure 1). However, none of the sequences in the three water column-associated groups are derived from cultured organisms or have genomic context, and thus cannot be conclusively linked to ammonia-oxidizing thaumarchaea at this time. Finally, differences in gene copy numbers between *amoA* and *nirK* within a given AOA genome may also contribute to the higher abundance of AnirKa in the epi- and mesopelagic: AOA are thought to only have a single *amoA* copy, whereas *Ca*. Np. maritimus has two *nirK* copies (Walker *et al.*, 2010) and *Ca*. Na. limnia has one *nirK* copy (Blainey *et al.*, 2011). However, the up to 10 times higher AnirKa abundance in surface water (Figure 5) is not likely to be due solely to differences in gene copy numbers.

Expression of thaumarchaeal *nirK* and *amoA* in Monterey Bay

Ammonia-oxidizing Thaumarchaea are thought to be a major source of N_2O in marine systems, based on its natural abundance isotopic composition (Santoro *et al.*, 2010; Santoro *et al.*, 2011). Nitrite reductase is potentially involved in N_2O production and we find thaumarchaeal *nirK* to be highly expressed in the marine water column (Figure 6b). A one to twofold higher expression of *nirK* over *amoA* has previously been reported in two meta-transcriptomes from coastal water (Hollibaugh *et al.*, 2011), suggesting that elevated *nirK* expression may be a general feature of AOA. In soil systems, where nitrifier activity can also be a significant contributor to N_2O emissions (Webster and Hopkins, 1996; Kool *et al.*, 2011), microcosm experiments did not show any correlation between thaumarchaeal *nirK* expression and N_2O production (Bartossek *et al.*, 2010). Similar experiments correlating thaumarchaeal *nirK* expression and N_2O production, or natural abundance isotope composition, have yet to be carried out in marine systems.

The mRNA abundance, in particular for AOA *amoA*, was low compared with the corresponding gene copy abundance (Figure 6a). This has also been reported in the Black Sea (Lam *et al.*, 2007), the Baltic Sea (Labrenz *et al.*, 2010) and in soil (Nicol *et al.*, 2008), whereas in the North Pacific Ocean the mRNA transcripts were 10–500-fold more abundant than gene copies in the upper 200 m of the water column (Church *et al.*, 2010).

Thaumarchaeal *nirK* diversity in marine sediments

Only the AnirKa primers amplified thaumarchaeal *nirK* sequences from sediment samples in San Francisco Bay (Figure 1). The samples span a salinity gradient from low salinity (BG20 ∼1 and BG30 ∼0.7) to high-salinity sediments (BC11 ∼30.5 and BA10 ∼25.5; Mosier and Francis, 2010), and a distinct low-salinity AOA community has previously been identified based on *amoA* gene sequences (Mosier and Francis, 2008). The low-salinity ecotype of thaumarchaeal *amoA* was also identified in this study (Supplementary Figure S3). UniFrac cluster analysis (Figure 2c) shows a high similarity in *amoA*-based AOA community composition between the two low-salinity sites, BG20 and BG30; however, the pattern is not reflected in the UniFrac cluster analysis of AnirKa sequences (Figure 2a). Thus, AnirKa also has a different distribution than AOA *amoA* in sediments.

Linking thaumarchaeal *nirK* and *amoA* sequence types

A direct link between *nirK* and *amoA* sequence types can be obtained through the available genomes of the sediment-derived AOA; *Ca*. Np. maritimus (Walker *et al.*, 2010) and *Ca*. Na. limnia (Blainey *et al.*, 2011). The *amoA* sequences from these two organisms share 89% nucleotide identity and belong to two different sequence clusters; *Ca*. Np. maritimus *amoA* groups with environmental sequences from marine/estuarine sediments and aquarium

biofilters, whereas *Ca.* Na. limnia *amoA* groups with sequences from low-salinity sediments (Supplementary Figure S3). *Ca.* Np. maritimus has two *nirK* copies and their nucleotide identities with *Ca.* Na. limnia *nirK* are 82 and 85%. All three *nirK* gene sequences cluster with the sediment group (Figure 1) where they fall into two different subgroups; however, a more extensive study of thaumarchaeal *nirK* from diverse sediment types is necessary to identify putative *nirK* ecotypes within sediments analogous to those typically found for *amoA* genes (Mosier and Francis, 2008).

In the marine water columns examined in this study, the similar distribution of the *nirK* group b-I and *amoA* group B sequence types suggests that these genes could be linked within the same organisms. AOA *amoA* group A sequence types are present throughout the water column, but predominating in the upper 10–100 m (Supplementary Figures S3 and S4). This distribution is consistent with a mixture of two *nirK* sequence types: group b-II, present in the upper water column, and a-II, present throughout the water column. Although a conclusive link between thaumarchaeal *nirK* and *amoA* sequences from the water column cannot be made with the present data alone, this could indeed be achieved via single-cell-based molecular techniques and/or metagenomic sequencing of samples (or cultures) highly enriched in marine planktonic AOA.

Acknowledgements

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