

## ORIGINAL ARTICLE

# Unique archaeal assemblages in the Arctic Ocean unveiled by massively parallel tag sequencing

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**The Arctic Ocean plays a critical role in controlling nutrient budgets between the Pacific and Atlantic Ocean. Archaea are key players in the nitrogen cycle and in cycling nutrients, but their community composition has been little studied in the Arctic Ocean. Here, we characterize archaeal assemblages from surface and deep Arctic water masses using massively parallel tag sequencing of the V6 region of the 16S rRNA gene. This approach gave a very high coverage of the natural communities, allowing a precise description of archaeal assemblages. This first taxonomic description of archaeal communities by tag sequencing reported so far shows that it is possible to assign an identity below phylum level to most (95%) of the archaeal V6 tags, and shows that tag sequencing is a powerful tool for resolving the diversity and distribution of specific microbes in the environment. Marine group I Crenarchaeota was overall the most abundant group in the Arctic Ocean and comprised between 27% and 63% of all tags. Group III Euryarchaeota were more abundant in deep-water masses and represented the largest archaeal group in the deep Atlantic layer of the central Arctic Ocean. Coastal surface waters, in turn, harbored more group II Euryarchaeota. Moreover, group II sequences that dominated surface waters were different from the group II sequences detected in deep waters, suggesting functional differences in closely related groups. Our results unveiled for the first time an archaeal community dominated by group III Euryarchaeota and show biogeographical traits for marine Arctic Archaea.**

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**Keywords:** archaea; Arctic Ocean; pyrosequencing; microbial diversity; group III Euryarchaeota; Crenarchaeota

## Introduction

The Arctic Ocean plays a crucial role in controlling the productivity of the world oceans through regulating nutrient budgets and cycling nitrogen between regions of high denitrification activity and net nitrogen fixation. Water flowing into the Arctic from the Pacific Ocean through the Bering Strait is depleted in nitrate with respect to phosphate (Jones *et al.*, 2003). This water flows out through the Canadian archipelago and Fram Strait to the Atlantic Ocean where microorganisms use phosphate to fix nitrogen. Through its Arctic journey the water's

nutrient properties change (Yamamoto-Kawai *et al.*, 2006), and this change probably influences primary production at a global scale. Although marine microorganisms are a major component of nutrient cycles (Arrigo 2005), the diversity and distribution of microbial communities in general remain poorly understood, and because of logistic constraints the Arctic is especially underexplored.

Archaea were first described as extremophiles, but are now recognized as ubiquitous and abundant in both marine (Karner *et al.*, 2001) and freshwaters environments (Auguet and Casamayor 2008). Marine Archaea are phylogenetically distributed through four main taxonomical clusters: one cluster of Crenarchaeota, the Marine group I (MGI), and three clusters of Euryarchaeota, group II, III and IV. The only cultivated representative of planktonic Archaea is known to oxidize ammonia and belongs to Crenarchaeota MGI (Konneke *et al.*, 2005). Recent

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evidence indicates that MGI Crenarchaeota are key players in chemoautotrophic ammonia oxidation (Hallam *et al.*, 2006b; Francis *et al.*, 2007), although other new data suggest that there are vertical and latitudinal gradients in the abundance of putative Crenarchaeota nitrifiers in the Atlantic Ocean and that some Crenarchaeota may live heterotrophically (Agogue *et al.*, 2008). The abundance of Crenarchaeota and their key role on the global nitrogen cycle makes them a fundamental component of the marine ecosystem. There are no cultivated representatives of marine planktonic Euryarchaeota to date and the functional role of that archaeal phylum remains little understood.

The MGI Crenarchaeota and group II Euryarchaeota are the most abundant Archaea in the oceans (Massana *et al.*, 2000). In temperate and tropical oceans Crenarchaeota are believed to be more frequent in deep waters, whereas group II Euryarchaeota are often more abundant in surface waters (Karner *et al.*, 2001; Herndl *et al.*, 2005; DeLong *et al.*, 2006). In the Arctic, the distribution of the different archaeal groups remains poorly resolved. One study found low abundance of group II Euryarchaeota down to 500 m (Kirchman *et al.*, 2007), other studies showed communities dominated by group II Euryarchaeota in surface waters (Galand *et al.*, 2006, 2008b), whereas a basin-wide survey described Crenarchaeota as more abundant than Euryarchaeota at 55 and 131 m (Bano *et al.*, 2004). The differing conclusions of these studies may be because of the use of different methods (Fluorescence *In Situ* Hybridization (FISH) versus

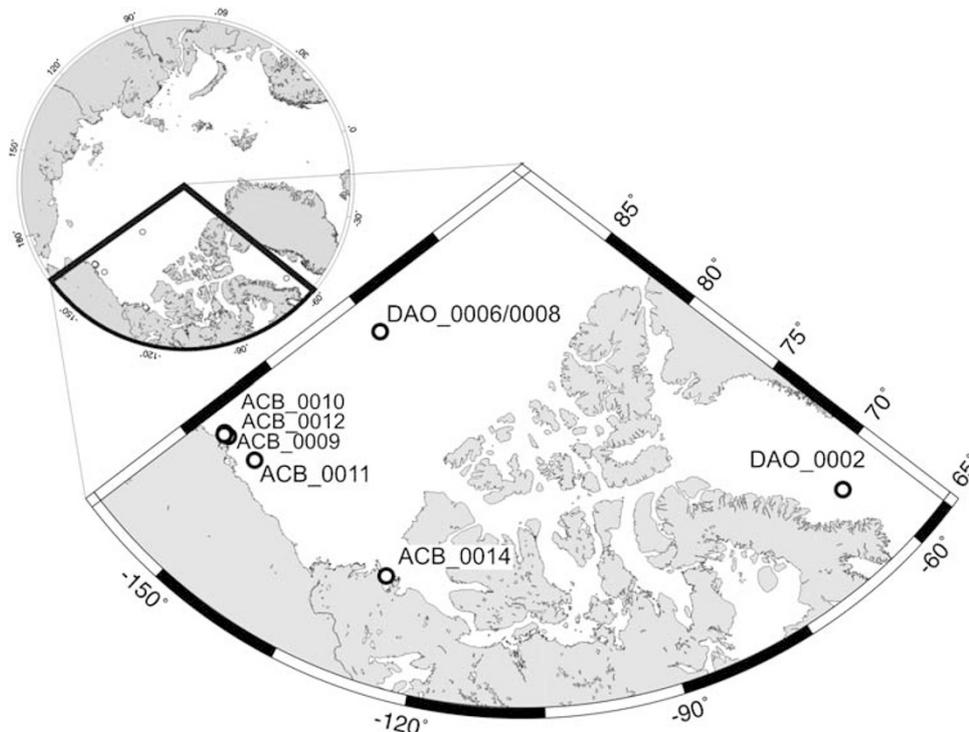
clone libraries), but could also be linked to the physical oceanography of the Arctic region, where archaeal groups are more closely linked to parent water masses than to depth alone (Galand *et al.*, 2009). The diversity and distribution of archaeal communities in the deeper Arctic Ocean remains unexplored.

The aim of this study was to describe the composition of archaeal assemblages in the Arctic Ocean, and to resolve the distribution of archaeal diversity within different water masses. We targeted five different water masses from both the coastal and central Arctic Ocean, that is, the coastal Pacific surface water and the polar mixed layer in the Beaufort Sea, the lower halocline and deep Atlantic water from the Canada Basin, and intermediate water from the Baffin Bay. To get an extended description of the diversity and a high coverage of the *in situ* populations, we characterized the archaeal assemblages by applying massively parallel tag sequencing to the hypervariable V6 region of the 16S rRNA gene (Sogin *et al.*, 2006). We assigned a taxonomical identity to the archaeal sequence tags and were able to describe unique assemblages within the archaeal community.

## Materials and methods

### Sampling sites and collection

We used eight samples collected from different geographical regions of the western Arctic Ocean (Figure 1). Sample DAO\_0002, 0006 and 0008 were



**Figure 1** Map of the Arctic Ocean showing the locations of sampling stations.

collected from the Canadian icebreaker *CCGS Louis St Laurent* as part of the International Polar Year study Canada's three Oceans project (C3O), ACB\_0014 was collected from the *CCGS Amundsen* as part of the Canadian Arctic Shelf Exchange Study. Samples ACB\_0009, 0010, 0011 and 0012 were collected as described earlier (Kirchman *et al.*, 2007; Malmstrom *et al.*, 2007). All ACB samples were collected from surface water masses, whereas the DAO samples were from deeper water masses. The water mass definition followed earlier descriptions (McLaughlin *et al.*, 2004; Rudels *et al.*, 2004; Tang *et al.*, 2004). DAO\_0002 was from the Baffin Bay intermediate water, DAO\_0006 from the deep Atlantic water and DAO\_0008 from the lower halocline of the Canada Basin (Table 1). Samples ACB\_0009, 0010, 0011 and 0012 were from the Pacific surface water taken from the Chukchi and Beaufort Sea (Table 1). Sample ACB\_0014 was from the polar mixed layer taken in Franklin Bay in the Amundsen Gulf.

The DAO and ACB\_0014 samples were collected with a rosette system equipped with a Seabird (Bellevue, WA, USA) 911 conductivity temperature and depth profiler. The rosette system was fitted with 12-L PVC bottles (Ocean Test Equipment Inc., Fort Lauderdale, FL, USA). Bottles were tripped on the upcast and seawater was collected from the rosette into bottles that had been cleaned in 10% (v/v) HCL, rinsed thrice with MilliQ water, and then thrice with sample water before filling. Within 30 min of collection, 6 l of seawater were filtered successively through a 50- $\mu$ m mesh, 3- $\mu$ m pore size 47-mm polycarbonate filter, and finally through a 0.2- $\mu$ m pore size Sterivex unit (Millipore Canada Ltd, Mississauga, ON, USA). The whole cell concentrates in the Sterivex unit were preserved in 1.8 ml of buffer (50 mM Tris-HCl, 0.75 M sucrose and 40 mM EDTA) and frozen at  $-80^{\circ}\text{C}$ . Other ACB samples were collected as described earlier (Malmstrom *et al.*, 2007).

Chlorophyll a was estimated using standard techniques (Nusch 1980). Total prokaryote counts for the DAO samples were kindly supplied by Dr WKW Li (Bedford Institute of Oceanography, Halifax, NS, Canada) from flow cytometry counts, and from microscopic analysis for ACB samples.

#### DNA extraction and pyrosequencing

DNA was extracted by digesting the cells with lysozyme (final concentration, 1 mg ml $^{-1}$ ) and proteinase K (0.21 mg ml $^{-1}$ ). Nucleic acids were then separated with phenol/chloroform (ACB samples) or a salt solution (DAO samples), and precipitated with ethanol. Earlier experiences indicated no differences in DNA recovery efficiency between the two techniques (C Lovejoy unpublished data). The archaeal hypervariable V6 region of the 16S rRNA gene was amplified using primers: 958arcF 5'-AATTGGANTCAACGCCGG-3', 1048arcRmajor

**Table 1** Physicochemical characteristics, water mass of origin and diversity estimates (calculated at the 3% difference level between 16S rRNA gene fragments) for eight archaeal assemblages from the Arctic Ocean

Location	Water mass	Date	Depth (m)	Bottom depth (m)	Salinity	Temperature ( $^{\circ}\text{C}$ )	Chla ( $\mu\text{g l}^{-1}$ )	Prokaryotes ( $10^5$ cells per ml)	No. of reads	No of OTUs	Chao1	ACE
DAO_0002	Baffin Bay	12/07/2007	1000	1800	34.5	0.9	0.001	0.98	18780	328	344	346
DAO_0006	Beaufort Sea (Canada basin)	04/08/2007	1000	3820	34.9	0.0	0	0.50	16194	578	611	619
DAO_0008	Beaufort Sea (Canada basin)	04/08/2007	410	3820	34.8	0.8	0	0.73	19741	479	517	509
ACB_0009	Coastal Beaufort Sea	13/07/2007	2	50	20.0	2.5	0.33	20.9	27836	544	578	574
ACB_0010	Coastal Chukchi Sea	11/07/2007	2	100	35.0	2.0	0.06	7.48	29088	625	731	723
ACB_0011	Beaufort Sea	30/07/2004	8.4	50	29.9	5.2	0.64	9.59	20867	488	549	535
ACB_0012	Coastal Chukchi Sea	28/01/2008	2	100	35.0	-1.8	0.06	4.67	33152	593	629	626
ACB_0014	Amundsen Gulf (Franklin Bay)	17/01/2004	3	200	29.8	-1.6	0.044	3.73	29449	700	865	819

5'-CGRCGGCCATGCACCWC-3' and 1048arcR-minor 5'-CGRCRGCCATGYACCWC-3' under conditions described in Huber *et al.* (2007). The final 90-bp amplicon was sequenced with 454 Life Sciences GS20 sequencer at the Marine Biological Laboratory in Woods Hole, MA, USA. For each read from the sequencer the primer bases were trimmed from the beginning and the end, and low-quality sequences were removed Huse *et al.* (2007). Sequences were flagged as low quality when they were <50 nucleotides in length, the start of the sequence did not exactly match a primer sequence, sequences contained ambivalent nucleotide assignments with one or more Ns (unknown nucleotides) or if the first five nucleotides of a tag did not correspond to the expected five nucleotides run key (used to sort the pyrosequencing reads). All sequences obtained for this study will be made publicly available at <http://vamps.mbl.edu/>.

#### Identification of archaeal phylotypes

The taxonomical identification of the sequence reads (tags) followed the approach proposed by Sogin *et al.* (2006). The tags were compared by BLASTN to a reference database of hypervariable region tags based on the SILVA database (version 94) (Pruesse *et al.*, 2007), and the 100 best matches were aligned to the tag sequences using MUSCLE (Edgar 2004). A reference sequence or sequences were defined as those having the minimum global distance (number of insertions, deletions and mismatches divided by the length of the tag) to the tag sequence, and all reads showing the best match to the same reference V6 tag were grouped together as the same operational taxonomic unit (OTU) (Sogin *et al.*, 2006). Taxonomy was assigned to each reference sequence with the RDP Classifier (Wang *et al.*, 2007).

This automated pipeline was, however, not precise enough to classify Archaea below the phylum level. To enhance the precision of the taxonomical affiliation, a representative sequence from each OTU was sent to the SeqMatch tool of the RDP Release 10 (Cole *et al.*, 2009), and up to three sequences rendering the best seqmatch score to the pyrosequencing reads were used as additional reference sequences. Reference sequences were aligned using the CLUSTAL W package (Thompson *et al.*, 1994) and trimmed down to an overlapping region present in all sequences; the final length of the overlapping region was 600 bp. A Jukes–Cantor corrected distance matrix was generated from the reference sequences with DNADIST and a tree was constructed using the FITCH program from PHYLIP (Felsenstein 2004). The different clusters composing the distance tree were then identified.

#### Diversity estimations and cluster analysis

Tags were aligned and a Jukes–Cantor corrected distance matrix was obtained using the DNADIST

program from PHYLIP (Felsenstein 2004). Rarefaction, the ACE and the Chao1 nonparametric richness estimators were then calculated with the program DOTUR (Schloss and Handelsman 2005). The 3% distance level between tag sequences was used for calculation of diversity estimators.

For cluster analysis, a distance matrix was computed from the abundance of OTU present in each community using Bray–Curtis similarity and a dendrogram was inferred with the unweighted pair-group average algorithm (UPGMA) as implemented in the program PAST (v 1.8, <http://folk.uio.no/ohammer/past/>).

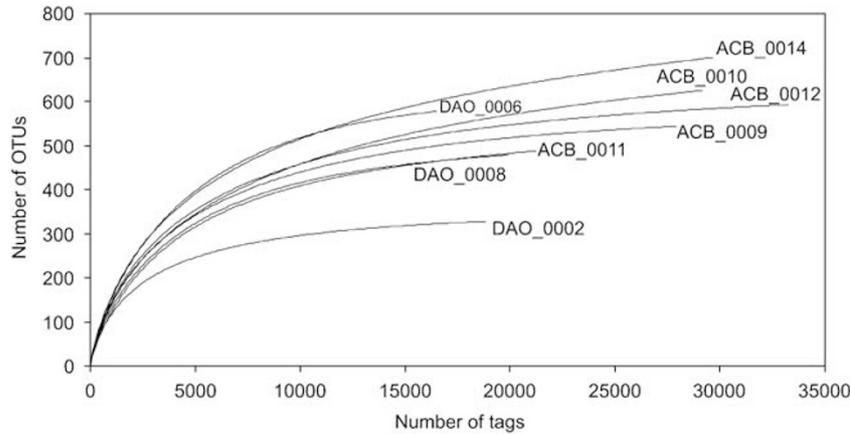
## Results

#### Characteristic of the water masses

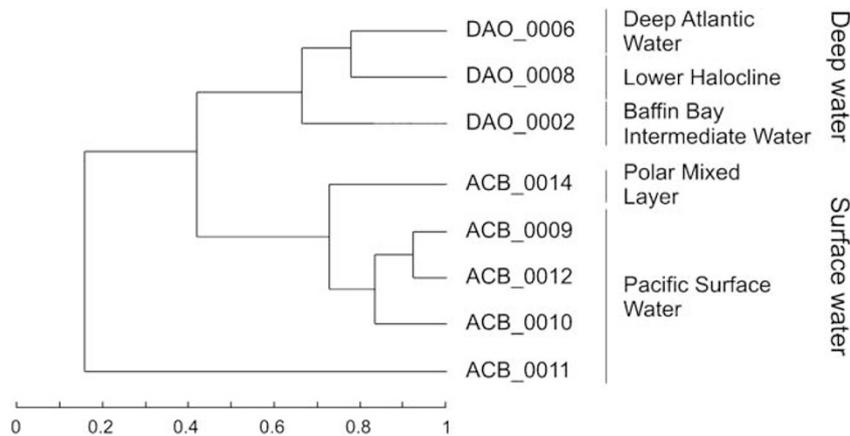
Sample DAO\_0002 was from the Baffin Bay intermediate water (Table 1), most often defined as having a temperature (T) > 0 °C and a salinity (S) > 34 (Tang *et al.*, 2004). Baffin Bay intermediate water originates from the mixing of the North Atlantic water flowing northward with the West Greenland current, and the Arctic water flowing southward through Smith Sound (Rudels *et al.*, 2004; Tang *et al.*, 2004). Samples DAO\_0006 was from the deep Atlantic water and DAO\_0008 from the lower halocline in the Canada Basin (Table 1). The Arctic Atlantic layer derives from the Norwegian Sea and ultimately from the North Atlantic. The Atlantic layer has  $T \geq 0$  °C and  $S \geq 34.5$  and is situated below the halocline. The halocline is defined as a pycnocline with higher temperature situated between the polar mixed layer (surface layer) and the Atlantic layer (Rudels *et al.*, 2004). Samples ACB\_0009, 0010, 0011 and 0012 were from coastal water sampled in the Beaufort Sea off Point Barrow (Alaska) and represent the Pacific surface water (Table 1). The Pacific water enters the Arctic through the Bering Strait, and the water salinity and temperature is altered through freshwater runoffs and ice melting and formation (McLaughlin *et al.*, 2004). Sample ACB\_0014 was from the polar mixed layer taken in Franklin Bay, Amundsen Gulf (Table 1). The polar mixed layer is a low salinity layer influenced by water of the Pacific origin and freshwater from the Mackenzie River (Garneau *et al.*, 2008).

#### Diversity of the archaeal assemblages

After removing low quality tags, the sequencing effort yielded a total of 195 107 reads for the eight samples analyzed, with on average 24 388 tags per sample (Table 1). The tags had a mean length of 69 nucleotides after trimming the primers. The coverage of the libraries was high with rarefaction curves reaching an asymptote at the 3% difference level (Figure 2). For six of the eight communities analyzed, the number of OTU detected was close to the total number of OTU estimated by Chao1 and



**Figure 2** Rarefaction curves for eight archaeal communities from the Arctic Ocean at a 3% difference level between 16S rRNA gene fragments.



**Figure 3** Dendrogram representing the similarity between tag composition of eight archaeal assemblages from surface (ACB) and deep (DAO) water masses of the Arctic Ocean described in Table 1. Clustering is on the basis of a distance matrix computed with Bray–Curtis index of similarity. The dendrogram was inferred with the unweighted pair-group average algorithm (UPGMA).

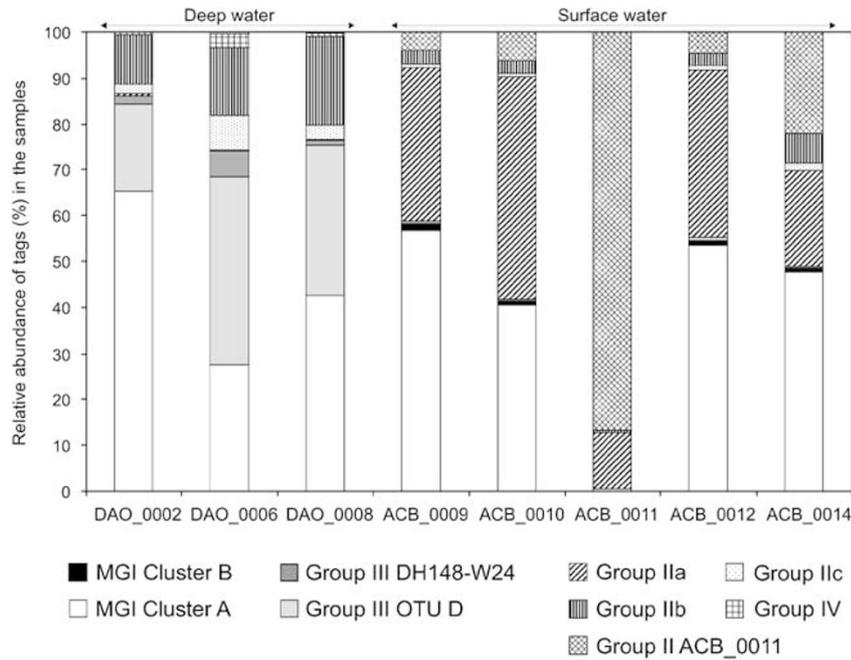
ACE diversity indices, additional evidence that the natural communities were well covered by the sequencing effort (Table 1). Surface samples and deep-water samples were distinct from each other and shared only 28% of OTUs in common. Overall, evenness was low and a small number of sequences were highly represented in the samples. In the surface samples, just three sequences accounted for >80% of all sequences recovered in the dataset. In the deep samples, six sequences accounted for >80% of all sequences.

Samples from coastal surface waters had generally higher diversity than samples from deeper water masses, and the polar mixed layer from the surface of Franklin Bay had overall the highest estimated richness (Figure 1, Table 1). The Baffin Bay intermediate water sample had the lowest richness, as its diversity was nearly half of that estimated for surface Beaufort Sea waters (Table 1).

*Composition and distribution of archaeal assemblages*  
Clustering analysis of the archaeal community composition separated the samples in two main

clusters: one contained all samples from deep-water masses and the other, all surface samples (Figure 3). Within surface water communities, sample ACB\_0011 formed a separate branch indicative of a distinct archaeal assemblage. Sample ACB\_0014 from the polar mixed layer in Franklin Bay was close to the other samples from the Pacific surface waters (Figure 3). Among deep-water samples, the two samples from the deep Atlantic water and lower halocline in the Canada Basin grouped together separately from Baffin Bay intermediate water communities (Figure 3).

MGI Crenarchaeota was overall the most abundant group in the Arctic Ocean and comprised between 27% and 63% of all tags. The deep Baffin Bay sample had the highest proportion of MGI Crenarchaeota, whereas the deep central Arctic had the lowest (Figure 4). The most abundant MGI sequences, representing >40% of the identified tags, belonged to the same cluster (MGI cluster A, Supplementary Figure S1). Within that cluster, the most abundant tag was not only 100% similar to the 16S rRNA sequence of *Nitrosopumilus maritimus*, but also to environmental sequences retrieved from



**Figure 4** Composition of eight archaeal assemblages from the Arctic Ocean. Naming of the different archaeal groups follows the clusters defined in Supplementary Figure S1.

various ecosystems, ranging from marine sediments and deep oceanic water masses to a freshwater rhizosphere. Sequences had best BLAST matches among others to OTUA, OTUB and OTUC clusters defined earlier by Martin-Cuadrado *et al.* (2008). Within MGI cluster A, a relatively abundant (up to 5% of the tags) sub-cluster (KM3–69, Supplementary Figure S1) was found specifically in the deep Arctic and was almost absent from surface waters (<0.2% of the tags). That cluster had 96% similarity to the target V6 region of 16S rRNA of *N. maritimus*. A second, less abundant cluster of MGI (MGI cluster B, Supplementary Figure S1) was detected mainly in the Pacific surface waters. Cluster MGI cluster B contained mainly sequences from sea sediments and was closely related to *N. maritimus* (98% sequence similarity). A few representative sequences of the newly defined Crenarchaeota group 1A (DeLong *et al.*, 2006) were also detected in deep Arctic samples, but at very low abundance (<200 tags). Only one group 1A tag was detected from surface samples.

Group III Euryarchaeota was the most abundant archaeal group in sample DAO\_0006 from the deep Atlantic water, and represented the second most abundant tags, after group I Crenarchaeota, in the lower halocline (DAO\_0008) and Baffin Bay intermediate waters (DAO\_0002) (Figure 4). Group III Euryarchaeotal sequences grouped into two clusters. The most abundant sequences grouped into the OTU D cluster (Supplementary Figure S1), following the designation by Martin-Cuadrado *et al.* (2008). V6 tags belonging to OTU D were not only 100% similar to sequences from the deep Southern Ocean, the Mediterranean and the Pacific Ocean (Mehta *et al.*, 2005;

Martin-Cuadrado *et al.*, 2008), but also from shallower (200 m) waters of the Pacific (Mincer *et al.*, 2007). A second less abundant V6 tag belonged to a cluster of group III Euryarchaeota, which we named cluster DH148-W24 after its first representative (López-García *et al.*, 2001). The tag was 100% similar to sequences retrieved from deep-water masses of the North and South Atlantic and the Pacific Ocean (DeLong *et al.*, 2006; López-García *et al.*, 2001). Very few group III Euryarchaeota sequence tags were detected in surface waters (<0.4 % of all tags); they all belonged to OTU D cluster.

In Pacific surface waters, the second most abundant archaeal group was group II Euryarchaeota. Group II contained the most abundant group of sequences in samples ACB\_0010, ACB\_0011 and ACB\_0014 (Figure 4). Group II Euryarchaeota sequences are grouped under three main clusters (Supplementary Figure S1). Most of the group II sequences from surface waters belonged to group II.a (Figure 4). The most abundant cluster II.a tags had the highest similarity (98%) to sequences from marine sediments (Clementino *et al.*, 2007) and surface Pacific waters (Frigaard *et al.*, 2006). In deep waters, the most abundant group II sequences belonged to cluster II.b (Figure 4). They had the highest similarity (100%) to sequences retrieved from deep waters of the Atlantic and the Pacific Ocean. Other deep-water group II sequences grouped within cluster II.c, earlier named OTU F (Martin-Cuadrado *et al.*, 2008) containing sequences from deep-water masses (Supplementary Figure S1).

The sample from the Beaufort Sea (Sample ACB\_0011) had a distinct community composition from all other samples. The community was

composed almost entirely of group II Euryarchaeota (>99% of the sequences), whereas MGI Crenarchaeota were practically absent (0.40 % of the sequences). One single group II tag represented >85% of the sequences. It was 100% similar to a sequence retrieved from surface waters of the Californian coast (AF257278).

Group IV sequences were detected in six of the eight samples, but were 50–200 times more abundant in deep-water samples from the central Arctic than in surface waters (Figure 4). Group IV Euryarchaeota were 100% similar to DH148-Y16 from deep-water masses of the South Atlantic (López-García *et al.*, 2001).

## Discussion

The importance of microorganisms in regulating nutrient and biogeochemical cycles is well recognized (Arrigo 2005). The temporal dynamics and distribution of marine microbial diversity remains, however, far from being completely understood, as entire oceanic regions such as the deep Arctic Ocean have never been explored. Here, we applied massively parallel tag sequencing as a tool to enhance our understanding of the diversity and distribution of archaeal assemblages in the Arctic Ocean, a poorly studied though globally important oceanic province. We were able to associate tags of the 16S rRNA gene hypervariable V6 region to taxonomical clusters within the uncultured Archaea, at a resolution precise enough to detect patterns in population distribution. We could thus identify the composition of archaeal assemblages from different water masses and unveiled unique patterns of diversity in the Arctic Ocean.

A surprising finding was that group III Euryarchaeota dominated the deep Atlantic water masses from the central Arctic (>40% of tag sequences) and was the second most abundant group in the two other deep-water samples. This is the first time group III is reported as dominating an archaeal assemblage. Group III Euryarchaeota were first identified in the Northeast Pacific (Fuhrman and Davis 1997) and was later detected in different parts of the Mediterranean Sea (Massana *et al.*, 2000; Martín-Cuadrado *et al.*, 2007, 2008), the South Atlantic (Martín-Cuadrado *et al.*, 2007; López-García *et al.*, 2001) and in the North Pacific Ocean (DeLong *et al.*, 2006). However, group III is usually rarely represented in marine ecosystems, in which MGI Crenarchaeota and group II Euryarchaeota are the most abundant groups (Massana *et al.*, 2000). The depths from which group III has been retrieved suggest that it is specific to the deep ocean. In fact, group III was absent or rare in clone libraries and Denaturing Gradient Gel Electrophoresis analysis from the upper layers of the central Arctic (Bano *et al.*, 2004) and coastal Beaufort Sea (Galand *et al.*, 2006, 2008a). One recent report from 1000 m depth

in the South Atlantic described an archaeal assemblage, in which group III was the second most abundant group of sequences retrieved (Martín-Cuadrado *et al.*, 2008). Remarkably, the same authors did not detect group III in nearby waters from the Antarctic Polar Front, suggesting a water mass-related occurrence of the group. Thus, the high abundances of group III in the deep Arctic Ocean and in one sample of the South Atlantic could indicate that the group successfully thrives in deep high latitude water masses, and that it deserves specific attention in future polar studies using quantitative methods such as FISH or qPCR.

Unfortunately, there are no cultivated representatives of group III Euryarchaeota to date and the metabolism and ecological role of those organisms remain unknown. Recently, however, metagenomic analysis revealed the presence in group III DNA fragments of genes that resembled those found in ammonia-oxidizing bacteria, suggesting that at least some members of group III Archaea may oxidize ammonia (Martín-Cuadrado *et al.*, 2008). If this is confirmed, the abundant presence of group III Euryarchaeota in the deep Arctic water could imply an important role for that group in the nitrogen cycle at high latitudes, and raises the question of possible competition with ammonia-oxidizing Crenarchaeota.

We found the highest proportion of group I Crenarchaeota (MGI) (65% of the sequences) in waters from the Baffin Bay intermediate layers. MGI Archaea were detected earlier in various oceanic regions and are believed to constitute the most abundant microbial group in deep waters (Karner *et al.*, 2001; DeLong *et al.*, 2006). In the Arctic, however, high proportions of MGI Crenarchaeota were found in Archaea clone libraries from surface layers of the Beaufort Sea (Galand *et al.*, 2008a,b), in the upper layers of the Central Arctic (Bano *et al.*, 2004) and in the western part of the Arctic Ocean, where almost no Euryarchaeota were detected with FISH (Kirchman *et al.*, 2007). Generally, Arctic MGI Crenarchaeota seemed to be more abundant in oligotrophic offshore waters (Galand *et al.*, 2008b).

One single representative of MGI has been isolated so far, *N. maritimus* (Konneke *et al.*, 2005), and the genome of the marine sponge symbiont *Cenarchaeum symbiosum* has been described (Hallam *et al.*, 2006a). The ability of those strains to oxidize ammonia and the wide abundance of archaeal genes putatively encoding the ammonia monooxygenase subunit A (*amoA*) all point to an important role of Crenarchaeota in the global nitrogen cycle (Wuchter *et al.*, 2006; Lam *et al.*, 2007), but recent findings have balanced this view suggesting that not all MGI Crenarchaeota are ammonia oxidizers. Recent data showed strong vertical and latitudinal gradients in the ratio of archaeal *amoA* to crenarchaeal 16S rRNA genes in the Atlantic Ocean (Agogue *et al.*, 2008). Similarly a

low *amoA* to 16S rRNA ratio was reported from an Arctic meromictic lake (Pouliot *et al.*, 2009) and other data suggest that some Crenarchaeota are heterotrophic (Teira *et al.*, 2006; Kirchman *et al.*, 2007). However, the archaeal *amoA* gene is abundant in the Arctic, indicating that Crenarchaeota contribute to nitrogen cycling in the Arctic Ocean (Galand *et al.*, 2009). In fact, the most common MGI tag (MGI cluster A), detected in both surface and deep-water mass samples had 100% similarity to the 16S rRNA gene of *N. maritimus*, suggesting that the detected Crenarchaeota were indeed ammonia oxidizers. On the other hand, we noted that the same tag also had a 100% match to sequences belonging to three other MGI clusters (OTU A, B and C in (Martin-Cuadrado *et al.*, 2008). As the tag was identical to sequences belonging to different crenarchaeotal groups, its affiliation could not be precisely resolved. This result clearly shows that the V6 region is relatively conserved among several MGI clusters. Longer reads, including additional hypervariable 16S rRNA gene regions, may be needed to increase taxonomical resolution. The tag approach, however, successfully separated other MGI clusters. Cluster KM3–69, for instance, was specifically detected in deep-water layers, whereas cluster MGI cluster B was specifically found in surface waters. MGI cluster B tags were similar to sequences retrieved from sediments, which may indicate that the presence of the group in coastal waters may be influenced by sediment re-suspension or discharge from riverine runoffs.

Group II Euryarchaeota was the second most abundant group after Crenarchaeota in the surface layers. Group II is generally thought to be more abundant in surface waters (Karner *et al.*, 2001; Herndl *et al.*, 2005; DeLong *et al.*, 2006), but in the Arctic, contrasting distributions have been reported. Group II Euryarchaeota were almost absent in FISH samples from the Western Arctic (Kirchman *et al.*, 2007), which is heavily influenced by Pacific waters, but this group was abundant in central Arctic clone libraries (Bano *et al.*, 2004) and dominated clone sequences from surface waters on the Mackenzie Shelf (Galand *et al.*, 2006). The discrepancy between studies may be because of the use of different probes, primers or extraction methods to detect or quantify Archaea (Casamayor *et al.*, 2002). A standardized approach combining quantitative and qualitative methods, and covering a vast range of water masses would allow a more precise description of archaeal distributions in the Arctic.

An overview of archaeal clone distribution in the southern Beaufort Sea and Amundsen Gulf suggests that coastal water masses, influenced by particle laden river runoff, would be a more suitable habitat for group II Euryarchaeota (Galand *et al.*, 2008b). Group IIa sequences found in the Pacific surface waters had high similarity to sequences retrieved from sediments, suggesting that the occurrence of

this group is related to sediment and particle loads exported by Arctic rivers to the adjacent Beaufort Sea. A recent metagenomic study showed that the genome of group II Euryarchaeota seemed enriched in putative anaerobic respiration components, suggesting that anaerobic pathways for energy production are present in the group (Martin-Cuadrado *et al.*, 2008). In the light of this finding and these data showing abundant group IIa sequences in coastal waters, we speculate that group IIa Euryarchaeota could be involved in the anaerobic degradation of organic matter coming from river runoff. The most abundant group II sequences found in the deep-water masses (cluster b and c) were distinct from the one detected in surface waters. The presence of different group II sequences in different environments may indicate that within group II Euryarchaeota ecological differences may favor different phylogenetic clusters or even different metabolic functions within the separate clusters.

Group IV Euryarchaeota were mainly found in deep Atlantic water, but always at very low abundance. Group IV was originally described from the Antarctic Polar Front, and further detected in the North Atlantic and Mediterranean Sea (López-García *et al.*, 2001). Group IV has always been detected in deep-water masses with the exception of one clone retrieved from upper Arctic mixed layers (Bano *et al.*, 2004). Our results confirm group IV as specifically inhabiting deep-water masses at a low abundance. Nothing is known at present about the metabolism of this group.

Finally, archaeal communities in the Arctic Ocean had a relatively low diversity, as the rarefaction curves reached an asymptote with a maximum estimated richness of 344–865 OTUs at a 97% level. Those results are in agreement with earlier reports, indicating that Archaea are generally less diverse than Bacteria, as shown through clone library surveys (Aller and Kemp 2008) and pyrosequencing of hydrothermal vents samples (Huber *et al.*, 2007).

In summary, we have shown that archaeal assemblages from Arctic surface layers were different from those in deep-water masses, with great differences in the amount of group II and group III Euryarchaeota. However, there were also large differences between the composition of the coastal Beaufort Sea community (sample ACB\_0011) and the other samples from Pacific surface water. MGI was essentially absent from the ACB\_0011 community, in which a single group II tag represented most of the sequences (85%). The difference in community composition was reflected in the physicochemical characteristics of the water. The warmer, less saline water of ACB\_0011 indicates the intrusion of a different water mass at that depth. The predominance of one single sequence tag may also indicate a bloom triggered, for example, by higher concentration of phytoplankton as indicated by high chlorophyll a values (Table 1).

For the first time, we successfully applied massively parallel tag sequencing to describe the taxonomic composition of archaeal communities. Our data revealed unique assemblages of Archaea in the Arctic Ocean, with group III Euryarchaeota dominating the deep Atlantic water mass. Those results open new perspectives on the possible competition between group III Euryarchaeota and MGI Crenarchaeota for putative energy sources such as ammonia oxidation. We also showed that the distribution of archaeal diversity varied with water masses, suggesting biogeographical traits for archaeal communities in the Arctic. Pyrosequencing gave a high coverage of these assemblages and offered a good taxonomical assignment for most of the groups as shown by the general agreement between the present V6 data and earlier clone library descriptions. The methodological approach had, however, some limitation, as the most abundant MGI tag could not be precisely affiliated. The low taxonomical resolution for that particular group is probably because of the presence of a more conserved V6 regions.

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