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**Influences of benthic infaunal burrows on community structure
and activity of ammonia-oxidizing bacteria in intertidal
sediments**

Running title: Community structure and activity of AOB in infaunal burrows

By

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ABSTRACT

Influences of benthic infaunal burrows constructed by the polychaete (*Tyllorrhynchus heterochaetus*) on O₂ concentrations and community structures and abundances of ammonia-oxidizing bacteria (AOB) and nitrite-oxidizing bacteria (NOB) in intertidal sediments were analyzed by the combined use of a 16S rRNA gene-based molecular approach and microelectrodes. The microelectrode measurements performed in an experimental system developed in an aquarium showed direct evidence of O₂ transport down to a depth of 350 mm of the sediment through a burrow. The 16S rRNA gene-cloning analysis revealed that the betaproteobacterial AOB communities in the sediment surface and the burrow walls were dominated by *Nitrosomonas* sp. Nm143-like sequences and most of the clones in *Nitrospira*-like NOB clone libraries of the sediment surface and the burrow walls were related to the *Nitrospira marina* lineage. Furthermore, we investigated vertical distributions of AOB and NOB in the infaunal burrow walls and the bulk sediments by real-time quantitative polymerase chain reaction (Q-PCR) assay. The AOB and *Nitrospira*-like NOB specific 16S rRNA gene copy numbers in the burrow walls were comparable with those in the sediment surfaces. These numbers in the burrow wall at a depth of 50-55 mm from the surface were, however, higher than those in the bulk sediment at the same depth. The microelectrode measurements showed higher NH₄⁺ consumption activity at the burrow wall as compared with those at the surrounding sediment. This result was consistent with the results of microcosm experiments showing that the consumption rates of NH₄⁺ and total inorganic nitrogen increased with increasing infaunal density in the sediment. These results clearly demonstrated that the infaunal burrows stimulated O₂ transport into the sediment in which otherwise reducing conditions prevailed, resulting in

development of high NH_4^+ consumption capacity. Consequently, the infaunal burrow became an important site for NH_4^+ consumption in the intertidal sediment.

INTRODUCTION

Benthic infaunal activities, such as burrow formation, burrow irrigation, defecation, and excretion of soluble and insoluble metabolites, increase the surface area across which solutes can diffuse into or out of the sediments and the substrate availability for inhabiting microorganisms (5, 13, 17, 24, 25, 32, 34). These burrows also provide a more stable physical environment compared to bulk sediment. Therefore, the sediment surrounding the infaunal burrows (i.e., burrow walls) could have markedly higher levels of microbial biomass, diversity, and activity compared with the bulk sediment. Previous studies have revealed that benthic infaunal activities resulted in changes in biogeochemical characteristics and microbial community structures in sediments (20, 22, 31). Although many studies demonstrated that presence of benthic infauna strongly affects the microbial ecology of estuarine sediments, surprisingly little attention has been paid to bioturbation effects on nitrification. In estuarine systems with high inputs of nitrogenous compounds, sediment is a major site for nitrification due to abundance of ammonia-oxidizing bacteria and their high activities (1). One such estuarine is the Niida River estuary in Hachinohe city, Japan, where the water quality deterioration (e.g. presence of organic carbon and NH_4^+) is evident, mainly due to the discharge of treated and untreated domestic and industrial wastewater and urban and agricultural run-off (26, 27).

The solute concentrations in an infaunal burrow have been measured by collecting the liquid samples in a burrow (17). This method was not completely satisfying because the

1 concentration measured was the mean value throughout the burrow. Application of
2 microelectrodes has enabled direct measurements of O₂ and nutrients in burrows without
3 sampling. Applications of O₂ microelectrodes revealed presence of O₂ in infaunal burrows
4 (5), in the burrows of freshwater insects (34), and in an actively ventilated polychaete
5 burrow (13). NH₄⁺ concentration profiles in freshwater sediments as influenced by insect
6 larvae were also measured by NH₄⁺ microelectrodes (1). These studies have demonstrated
7 evidence of enhanced mass transport through the burrows. However, the measurements
8 were limited in just a few cm depths from the sediment surface due to low accessibility of
9 the microelectrodes and the uncertainty of exact position of the burrow.

10 Microbial community structures in infaunal burrows and tubes have been investigated
11 by 16S rRNA gene-cloning analysis (20, 22) and ester-linked phospholipid fatty acid
12 analysis (31). These studies have provided a good understanding of the microbial
13 community structure and diversity in the burrow and sediment, and allowed comparison
14 with biogeochemical characteristics. One study revealed that the microbial community
15 structures in burrow walls were different from those in the bulk sediment (31). However,
16 community structure, abundance and *in situ* activity of nitrifying bacteria in infaunal
17 burrows and bulk sediment have not been analyzed, compared, and linked to available O₂
18 concentrations in the burrows.

19 Therefore, we have investigated the influences of infaunal burrows on microbial
20 community structures and the abundance of ammonia-oxidizing bacteria (AOB) and
21 nitrite-oxidizing bacteria (NOB) and *in situ* activities of AOB in intertidal sediments by
22 applying 16S rRNA gene-cloning analysis, real-time quantitative polymerase chain reaction
23 (Q-PCR) assay and microelectrodes. The Niida River estuarine sediment was selected, in

1 which a high number of infaunal burrows were constructed by the benthic infaunas,
2 *Tylorrhynchus heterochaetus*, which generally inhabited in the intertidal zone of the
3 Japanese estuary. To directly measure *in situ* O₂ concentration profiles in the burrows, we
4 have constructed a continuous flow aquarium with agar slits in a sideboard, through which
5 microelectrodes could be inserted into the burrow.

7 MATERIALS AND METHODS

8 *Sampling*

9 River water and sediment samples were collected as described below during the low
10 tide at an intertidal area of the Niida River, Hachinohe city, Japan, which was located
11 approximately 1.5 km from the river mouth (27). The samples were immediately transferred
12 to the laboratory and were analyzed within 12 h.

14 *Microcosm experiments*

15 Microcosm experiments were carried out to determine the influence of infaunal
16 density on the consumption rates of NH₄⁺ and total inorganic nitrogen (N_i) that was defined
17 as the sum of the concentrations of NH₄⁺, NO₂⁻ and NO₃⁻. Grab samples of sediments were
18 obtained at the study site and passed through a 1-mm mesh to remove pebbles, large detritus
19 particles and indigenous infaunas. After thorough mixing the sediments were apportioned
20 into cylindrical sediment containers (11.4 cm in diameter and 30 cm height) to give a final
21 sediment height of 30 cm. Various numbers of *T. heterochaetus* were placed on the sediment
22 surface in each microcosm and allowed to burrow into the sediments. They generally
23 burrowed within a few minutes. The sediment surfaces in the microcosms were covered

with 1-mm meshes to prevent the infaunas from moving out of the microcosms. Then, the microcosms were buried in the sediment at the study site, where the surfaces of the microcosms were aligned with the surface of the natural sediment at the study site. The microcosms were allowed to stabilize for 2 to 3 weeks. Then each microcosm was brought to the laboratory and placed in an aquarium filled with 3 liters of the river water collected at the same site. NH_4Cl and NaNO_3 were added to the river water, resulting in final concentrations of approximately $360\ \mu\text{M}$ of NH_4^+ and NO_3^- , respectively. O_2 concentration of the overlaying water was kept ca. $210\ \mu\text{M}$ by continuously bubbling with air. The microcosms were incubated for 48 h in the dark. The changes in NH_4^+ , NO_2^- and NO_3^- concentrations in the overlying water were monitored with time. The consumption rates of NH_4^+ ($R(\text{NH}_4^+)$) and N_i ($R(\text{N}_i)$) were calculated from the decreases in NH_4^+ and N_i concentrations during the initial 12-h incubation, respectively. In total, 16 microcosm experiments were conducted with different infaunal densities.

Microelectrode measurements

The concentration profiles of O_2 and NH_4^+ in the sediment were measured in the laboratory using microelectrodes as described by Nakamura et al. (26). Clark-type microelectrodes for O_2 were prepared and calibrated as described by Revsbech (30). The LIX-type microelectrodes for NH_4^+ were constructed, calibrated, and used according to the protocol described by de Beer et al. (9) and Okabe et al. (28). To directly monitor O_2 concentrations inside an infaunal burrow along the depth, we constructed an aquarium with an acrylic plate (20 cm wide, 1 cm thick and 50 cm height) (see **Fig. S1** in the supplemental material). There were 45 slits ($0.5 \times 5\ \text{cm}$), which were filled with 3% agar plate, in one side

of the aquarium (see **Fig. S1C** in the supplemental material). By this means, we could determine the burrow structure and microelectrode position in the burrow. The aquarium was filled with the sediment collected in the same way as that for the microcosm experiments. An infauna (*T. heterochaetus*) was placed on the sediment surface and allowed to burrow. River water was continuously fed to the aquarium at a flow rate of 2 ml min⁻¹. The aquarium was maintained at 20°C in the dark. After 3 days, the infauna created a visible burrow. For the measurements of O₂ concentrations inside the infaunal burrow, the O₂ microelectrode was inserted into the burrow through the agar plate.

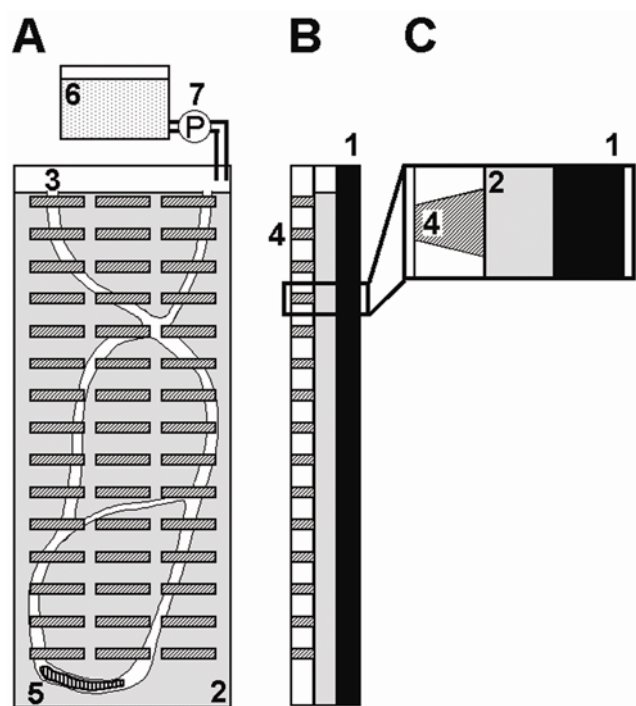


Fig. S1. Experimental apparatus for monitoring O₂ concentrations in an infaunal burrow: 1, sideboard; 2, sediment; 3, an infaunal burrow; 4, agar plates; 5, an infauna; 6, a tank filled with river water; 7, a pump. (A) A front view. (B) A side view. (C) A close-up view of the side view enclosed by the box in panel B.

In order to analyze NH_4^+ consumption rates in the burrow wall, the concentration profiles of O_2 and NH_4^+ were measured at a cross-section of the sediment in the aquarium. A synthetic medium was used to avoid interference with the LIX-type microelectrodes for NH_4^+ (26). The sediment was incubated in the medium at 20°C for more than 30 min before measurements to ensure that steady-state profiles were obtained. Three concentration profiles were measured for each chemical species and at each measuring point. The details of microelectrode measurements are described elsewhere (26, 27). Based on the O_2 and NH_4^+ concentration profiles measured, the total O_2 and NH_4^+ consumption rates were calculated using Fick's first law of diffusion (26, 27). The molecular diffusion coefficients used for the calculations were $2.09 \times 10^{-5} \text{ cm}^2 \text{ s}^{-1}$ for O_2 in liquid, $1.38 \times 10^{-5} \text{ cm}^2 \text{ s}^{-1}$ for NH_4^+ in liquid (3), and $2.2 \times 10^{-5} \text{ cm}^2 \text{ s}^{-1}$ for O_2 in 3% agar plate at 20°C (16). Differences between the rates were statistically analyzed by *t* test.

DNA extraction and PCR amplification

Three sediment samples (approximately 1 cm^3) were collected with sterile spatulas at different points corresponding to each sampling position (i.e., sediment surface, bulk sediment, and burrow walls at depths of 25-30 mm and 50-55 mm). DNA was extracted from each sample (approximately 0.2 cm^3) using a Fast DNA spin kit (Bio 101, Qiogene Inc., Carlsbad, Calif) as described in the manufacturer's instructions. The 16S rRNA gene fragments from the extracted total DNA were amplified with EX Taq DNA polymerase (TaKaRa Bio Inc., Ohtsu, Japan) by using the AOB specific primer set CTO189fA/B, CTO189fC and CTO654r (18) as well as the *Nitrospira*-like NOB specific primer set of Ntspa685 (15) and NTSPAf (26). The PCR conditions used for AOB and *Nitrospira*-like

NOB were described by Hermansson and Lindgren (14) and Nakamura et al. (26). PCR products were electrophoresed on a 1% (wt/vol) agarose gel. To reduce the possible bias caused by PCR amplification, the 16S rRNA gene was amplified in triplicate tubes for each sample, and then in total 9 PCR products were combined for the next cloning step.

Cloning and sequencing of the 16S rRNA gene and phylogenetic analysis

The purified PCR products were ligated into a pCR-XL-TOPO[®] vector and transformed into ONE SHOT *Escherichia coli* cells following the manufacturer's instructions (TOPO[®] XL PCR cloning; Invitrogen). Partial sequencing of 16S rRNA gene inserts (465 bp for AOB and 510 bp for *Nitrospira*-like NOB) was performed using an automatic sequencer (ABI Prism 3100 Avant Genetic Analyzer; Applied Biosystems) with a BigDye terminator Ready Reaction kit (Applied Biosystems). All sequences were checked for chimeric artifacts by the CHECK_CHIMERA program in the Ribosomal Database Project (21) and compared with similar sequences of the reference organisms by a BLAST search (2). Sequence data were aligned with the CLUSTAL W package (33). Clones with more than 97% sequence similarity were grouped into the same operational taxonomic unit (OTU), and their representative sequences were used for phylogenetic analysis.

Quantification of AOB and Nitrospira- and Nitrobacter-like NOB by Q-PCR.

Sediment samples were collected from sediment surface, burrow walls and bulk sediments at depths of 5-10 mm, 25-30 mm and 50-55 mm as described above. Total cell counts were performed after the diluted sediment samples on 0.2- μ m-membrane filters were stained with 6-diamidino-2-phenylindole (DAPI). At least 15 replicate analyses were

performed for each sample. Q-PCR assays were performed to quantify AOB and *Nitrospira*- and *Nitrobacter*-like NOB specific 16S rRNA genes. The Q-PCR assay for betaproteobacterial AOB and *Nitrospira*-like NOB was performed as described previously (14, 26). The Q-PCR assay for *Nitrobacter*-like NOB was performed in total volume of 25 μ l with 12.5 μ l of SYBR Green PCR Master Mix (Applied Biosystems), 7.5 pmol of each of the forward and reverse primers (FGPS872f and FGPS1269r) (10), 2.5 μ l of bovine serum albumin solution (Invitrogen) and either 0.1 pg of sample DNA or 10 to 10^5 copies per well of the standard bacterium DNA of *Nitrobacter winogradskyi* (NBRC 14297). All Q-PCRs were performed in MicroAmp Optical 96-well reaction plates with optical cap (PE Applied Biosystems). The template DNA in the reaction mixtures was amplified and monitored with an ABI prism 7000 Sequence Detection System (PE Applied Biosystems). The cycling regime for AOB and *Nitrospira*-like NOB was as follows: hold for 2 min at 50°C; hold for 10 min at 95°C; and 40 cycles of 15 sec at 95°C and 1 min at 60°C. The cycling regime for *Nitrobacter*-like NOB was as follows: hold for 2 min at 50°C; hold for 10 min at 95°C; and 80 cycles of 15 sec at 95°C and 1 min at 50°C. The detection limits for AOB, *Nitrospira*- and *Nitrobacter*-like NOB in this study were 2.7×10 , 1.6×10^2 and 5.4×10 copies per well, respectively, which correspond to 6.7×10^4 , 4.0×10^4 and 1.4×10^4 copies cm^{-3} when the sediment sample volume and DNA extraction step are taken into account. Four replicate analyses were performed for each sample. A *t* test was applied to evaluate differences of the total bacterial cells and AOB and *Nitrospira*-like NOB specific 16S rRNA gene copy numbers among the samples.

Analytical methods

The NH_4^+ concentrations were colorimetrically determined (6) and the NO_2^- and NO_3^- concentrations were determined using an ion chromatograph (HIC-6A; Shimadzu) equipped with a Shim-pack IC-AI column. The samples for NH_4^+ , NO_2^- and NO_3^- were filtered through 0.2- μm -membrane filters before the analysis. The O_2 concentration and pH in the overlaying water were directly determined using an O_2 and a pH electrode, respectively.

Nucleotide sequence accession numbers

The GenBank/EMBL/DDBJ accession numbers for the 16S rRNA gene sequences of representative 35 clones used for the phylogenetic analysis are AB239541, AB239545, AB239546, AB239560, AB239561, AB239566, and AB264558 to AB264586.

RESULTS AND DISCUSSION

Sediment characteristics

Many burrow openings with approximately 5 mm in diameter were found on the sediment surface at the study site during the low tide. Three types of benthic infaunas inhabited in the sediment. Numerically most abundant species was *T. heterochaetus* (84%) whereas the densities of *Notomastus* sp. (11%) and *Neanthes japonica* (5%) were low. Because the size of *T. heterochaetus* was much bigger (approximately 5 mm in diameter) than other two species (<1 mm in diameter), we speculated that all of the visible burrows were created by *T. heterochaetus*. The visible burrows extended down to a depth of at least 350 mm. Density of benthic infaunas in the upper 350 mm of the sediment was ca. 5700 individuals m^{-3} . Assuming a burrow of *T. heterochaetus* to be a straight tube with 5 mm in

diameter and 350 mm in length, the specific surface area of the burrow walls in the upper 350 mm of the sediment was calculated to be $26.4 \text{ m}^2 \text{ m}^{-3}$. The burrow walls were covered with thin oxidized-light brown layers. The color of the burrow wall was similar to that of the sediment surface. Average NH_4^+ , NO_2^- and NO_3^- concentrations ($\pm$ standard deviation) in the overlaying water were $44 \pm 39 \text{ } \mu\text{M}$, $1 \pm 5 \text{ } \mu\text{M}$ and $127 \pm 168 \text{ } \mu\text{M}$, respectively ($n = 57$). During the high tide, we occasionally found that suspended particles flew into the burrow and the fluffy biofilm developed at the burrow opening oscillated, indicating the overlaying water was introduced into the burrow. In contrast, the water in burrows stood during the low tide. Thus, we expected that the local environment in the burrows (i.e., the concentrations of O_2 , NH_4^+ , NO_2^- and NO_3^-) was dynamically changing with time as compared with that in the river water. Further information on the sediment at the study site and the physical and chemical parameters in river water can be found elsewhere (26, 27).

Microcosm experiments

The consumption rates of NH_4^+ ($\text{R}(\text{NH}_4^+)$) and N_i ($\text{R}(\text{N}_i)$) of the sediments with various benthic infaunal densities (i.e., *T. heterochaetus*) were determined in the microcosm experiments (**Fig. 1**). Mean values ($\pm$ standard deviation) of $\text{R}(\text{NH}_4^+)$ and $\text{R}(\text{N}_i)$ of the sediment without the infauna were $670 \pm 540 \text{ } \mu\text{mol m}^{-2} \text{ h}^{-1}$ and $1260 \pm 770 \text{ } \mu\text{mol m}^{-2} \text{ h}^{-1}$, respectively. Both rates increased as infaunal density increased. The increase in the $\text{R}(\text{N}_i)$ was more significant as compared with the $\text{R}(\text{NH}_4^+)$.

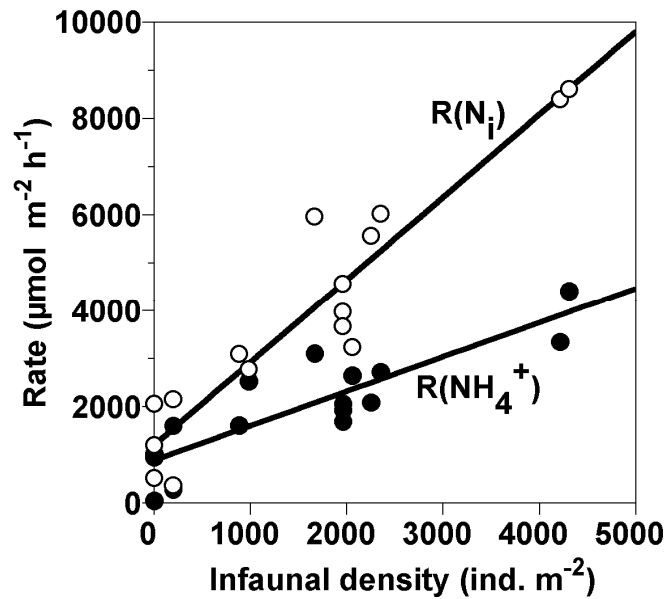


Fig. 1. The consumption rates of NH_4^+ ($R(\text{NH}_4^+)$) and total inorganic nitrogen ($R(\text{N}_i)$) of the sediment as a function of the density of *T. heterochaetus* in the microcosm experiments. The solid lines indicate linear regression of the data. The equations of the straight lines were $y = 0.72x + 890$ with $r^2 = 0.75$ (NH_4^+ consumption) and $y = 1.72x + 1200$ with $r^2 = 0.88$ (total inorganic nitrogen consumption).

Community structures of AOB and NOB

Three 16S rRNA gene clone libraries of AOB belonging to the *Betaproteobacteria* were constructed from three sediment samples taken from the sediment surface (SS) and the burrow walls at depths of 25-30 mm (BW-25) and 50-55 mm (BW-50) (Table 1 and Fig. S2 in the supplemental material). Sixty-seven, forty-seven and fifty-two clones were randomly selected from the SS, BW-25 and BW-50 clone libraries, respectively, and the partial sequences of 465 bp were analyzed. In total, the clones were grouped into 27 OTUs, and their representative sequences were used for phylogenetic analysis (Fig. S2). According to Purkhold et al. (29), we classified the *betaproteobacterial* AOB into seven stable lineages

(*Nitrosomonas oligotropha*, *Nitrosomonas marina*, *Nitrosomonas cryotoleransa*,
Nitrosomonas europaea/*Nitrosococcus mobilis*, *Nitrosomonas communis*, *Nitrosomonas* sp.
Nm143, and *Nitrospira briensis*). In all three samples the most frequently detected clones
were affiliated with the *Nitrosomonas* sp. Nm143 lineage with 93-99% sequence similarity
(**Table 1**). These clones represented 30, 62 and 44% of the total clones recovered from the
SS, BW-25 and BW-50 samples, respectively. *Nitrosomonas* sp. Nm143-like sequences
have been found at intermediate salinity sites of other estuaries (4, 12). The second
frequently detected clones recovered from the SS and BW-25 samples were affiliated with
the *Nitrosomonas marina* lineage (the detection frequency of 9 and 19%, respectively),
whereas the second frequently detected clones were affiliated with the *Nitrospira briensis*
lineage (the detection frequency of 17%) in the BW-50 samples. *Nitrosomonas marina*-like
sequences have also been detected in relatively high-salinity environments (7, 12), because
the *N. marina* lineage comprises obligate halophilic and salt-tolerant species. Thus, the
AOB community structures in the burrow walls were similar to that in the sediment surface.
This reflected the transport of overlying water into the burrows as confirmed by the
microelectrode measurements. In this study, we focused only on the community structure of
aerobic AOB affiliated with the *betaproteobacteria* whereas other ammonia-oxidizing
microorganisms (e.g. anaerobic ammonium-oxidizing bacteria (i.e., ANAMMOX bacteria),
the genus *Nitrosococcus* affiliated with the *gammaproteobacteria* and ammonia-oxidizing
Crenarchaea) have not been analyzed. Contribution of these AOB to NH_4^+ oxidation in the
sediment should be examined in the future study.



Fig. S2. Phylogenetic tree for the betaproteobacterial AOB, showing the positions of the clones obtained from three different points in the sediment. The tree was generated by

1 using 465 bp of the 16S rRNA genes and the neighbor-joining method. Scale bar = 5%
2 sequence divergence. Parsimony bootstrap values of 70 or greater are presented at the
3 nodes (from 100 replicates). The *N. marina* sequence (X82559) served as the outgroup for
4 rooting the tree. The numbers in parentheses indicate the frequencies of appearance of
5 identical clones in the total clones analyzed.

6
7 In contrast to the AOB community structure, information about community structure
8 and abundance of NOB in intertidal sediments is scarce. Although *Nitrobacter* is the most
9 commonly isolated and studied NOB from water environments, recent studies demonstrated
10 the presence of *Nitrospira* as dominate NOB in sediments (1, 7, 12, 26). In this study, the
11 *Nitrospira*-like NOB specific 16S rRNA gene copy numbers were one to three orders of
12 magnitude higher than those of the *Nitrobacter*-like NOB (**Table 2**). Hence, the community
13 structures of *Nitrospira*-like NOB in the sediment were further analyzed. Three 16S rRNA
14 gene clone libraries of *Nitrospira*-like NOB were constructed from the SS, BW-25 and
15 BW-50 samples (**Fig. 2** and **Table S1** in the supplemental material). Partial sequences of
16 510 bp were analyzed from 46, 44, and 46 clones randomly selected from the SS, BW-25
17 and BW-50 clone libraries, respectively. The diversity of the *Nitrospira*-like NOB clone
18 libraries was very low and more than 97% of the clones analyzed were related to the
19 *Nitrospira marina* lineage with 96-99% sequence similarity. Other clones were related to
20 the *Nitrospira defluvii* lineage with 97-99% sequence similarity (**Table S1**). Thus, the
21 *Nitrospira*-like NOB community structures in the burrow walls were also similar to that in
22 the sediment surface. Similarly, presence of *Nitrospira marina*-like NOB in estuarine
23 sediment was indicated by stable isotope probing analysis (12). *Nitrospira marina*-like

NOB were also detected in a submerged filter treating a high salinity industrial wastewater containing NH_4^+ and phenol (8) and in biofilms developed in freshwater or seawater aquaria (15).

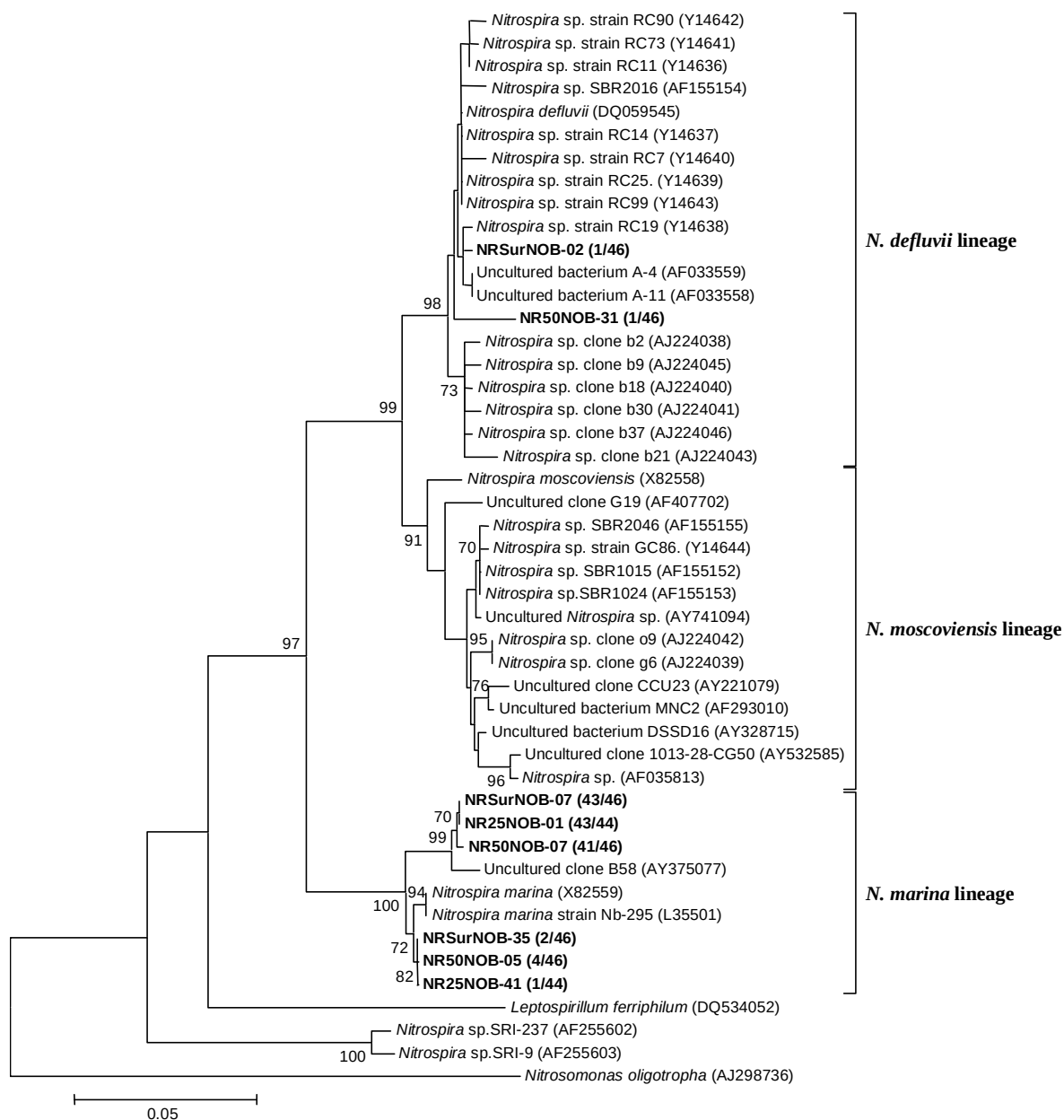


Fig. 2. Phylogenetic tree for *Nitrospira*-like NOB, showing the positions of the clones obtained from three different points in the sediment. The tree was generated by using 510 bp of the 16S rRNA genes and the neighbor-joining method. Scale bar = 5% sequence

divergence. Parsimony bootstrap values of 70 or greater are presented at the nodes (from 100 replicates). The *N. oligotropha* sequence (AJ298736) served as the outgroup for rooting the tree. The numbers in parentheses indicate the frequencies of appearance of identical clones in the total clones analyzed.

Microbial density

Total microbial cell counts were performed on different layers of the bulk sediment and the burrow wall samples (**Table 2**). The lateral average of total bacterial cell numbers was highest (6.5×10^9 cells cm^{-3}) at the sediment surface and slightly decreased with depth down to 4.0×10^9 cells cm^{-3} at a depth of 50-55 mm of the sediment. The cell numbers in the burrow wall were lower (1.0 to 1.9×10^9 cells cm^{-3}) than those of the bulk sediment samples ($P < 0.0001$, $n=15$) probably because the burrow walls were composed of loosely packed sediment (biofilms).

Betaproteobacterial AOB and *Nitrospira*- and *Nitrobacter*-like NOB specific 16S rRNA gene copy numbers were quantified by Q-PCR assay (**Table 2**). The AOB and *Nitrospira*-like NOB specific 16S rRNA gene copy numbers were in the range of 10^7 copies cm^{-3} except those at a depth of 50-55 mm of the bulk sediment. Presence of aerobic AOB in anoxic parts of the sediment could be explained by direct transport of AOB from the sediment surface by mixing, persistence of AOB, and capability of anoxic respiration of AOB. In contrast, the *Nitrobacter*-like NOB specific 16S rRNA gene copy numbers were one to three orders of magnitude lower than the *Nitrospira*-like NOB specific 16S rRNA gene copy numbers. Thus, *Nitrospira*-like NOB might be numerically dominant NOB in the intertidal sediment. The AOB and *Nitrospira*-like NOB specific 16S rRNA gene copy

numbers in the burrow walls (1.2 to 4.2×10^7 and 1.0 to 3.3×10^7 gene copies cm^{-3} for AOB and *Nitrospira*-like NOB, respectively) were comparable with those at the sediment surface (2.4 and 2.8×10^7 gene copies cm^{-3} for AOB and *Nitrospira*-like NOB, respectively) ($P > 0.05$, $n = 4$). These copy numbers slightly increased with depth in the burrow wall, whereas in the bulk sediment these copy numbers decreased with depth. Therefore, the AOB and *Nitrospira*-like NOB specific 16S rRNA gene copy numbers became higher in the burrow wall than in the bulk sediment at a depth of 50-55 mm ($P < 0.01$, $n = 4$).

Microelectrode measurements

The concentration profiles of O_2 and NH_4^+ were measured at a cross-section of the sediment (**Fig. 3A**). The microelectrodes were inserted into 4 points on the cross-section; the burrow wall at depths of 5 mm (point 1), 25 mm (point 2) and 50 mm (point 3), and the bulk sediment at a depth of 50 mm (point 4), as indicated in **Fig. 3B**. O_2 penetration depths were in the range of 0.6 to 1.2 mm at the cross-section of the sediment (**Fig. 3C**). The total O_2 consumption rates at the points 1, 2, 3 (i.e., in the burrow wall) and 4 (i.e., in the bulk sediment) were calculated to be 0.19, 0.21, 0.26 and 0.23 $\mu\text{mol cm}^{-2} \text{h}^{-1}$, respectively. Moreover, the O_2 concentrations at the sediment surface and bulk sediments at 5 mm and 25 mm were measured (data not shown) and the total O_2 consumption rates were calculated to be 0.20, 0.23 and 0.21 $\mu\text{mol cm}^{-2} \text{h}^{-1}$, respectively. NH_4^+ was consumed in the upper parts of the sediment at the points 1, 2 and 3, indicating high NH_4^+ consumption activity at the burrow wall (**Fig. 3D**). The total NH_4^+ consumption rates at the points 1, 2 and 3 were calculated to be 0.050, 0.026 and 0.033 $\mu\text{mol cm}^{-2} \text{h}^{-1}$, respectively. The total NH_4^+ consumption rates at the sediment surface and at the bulk sediment at a depth of 50 mm were

1 0.028 and 0.003 $\mu\text{mol cm}^{-2} \text{h}^{-1}$, respectively. Although O_2 consumption rates in the burrow

2 wall were similar to those in the sediment surface ($P > 0.05$), the total NH_4^+ consumption

3 rate at the point 1 was higher than that at the sediment surface ($P < 0.05$) and the rates at the

4 points 2 and 3 were comparable to that at the sediment surface. NH_4^+ was produced in the

5 deeper part of sediments at the points 3 and 4.

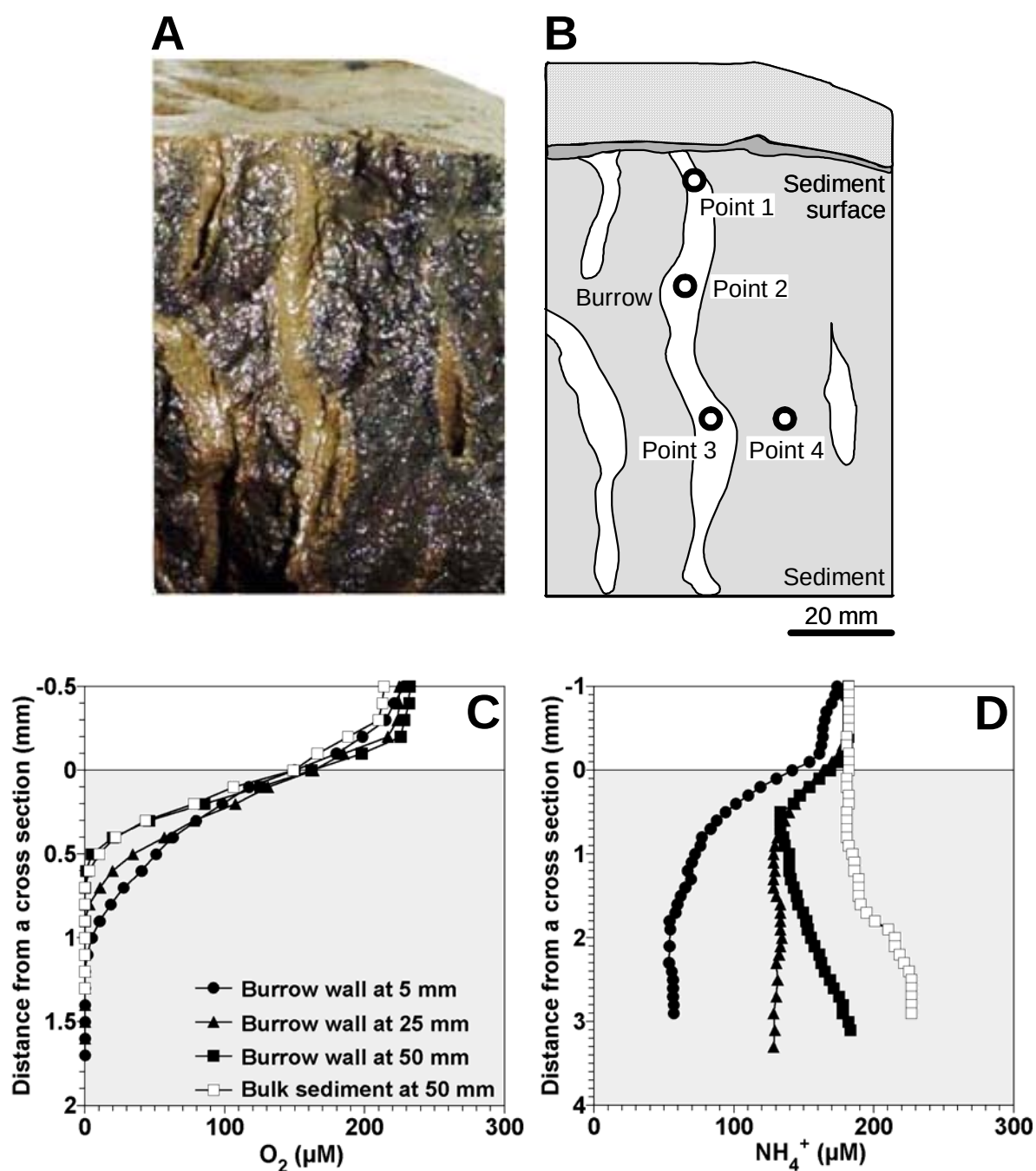


Fig. 3. Concentration profiles of O_2 and NH_4^+ at a cross-section of the sediment. (A) Photograph of the cross-section of the sediment. (B) A drawing of the cross-section of the sediment indicated in panel A. The points 1 to 4 in panel B indicate the points where the microelectrodes were inserted. (C and D) Concentration profiles of O_2 and NH_4^+ measured at each point, respectively. The profiles are average values ($n = 3$) and the standard deviations were less than 10% of the average values. Zero on the horizontal axis corresponds to the surface of the cross-section. The legends of panel D were the same as those in panel C.

Microelectrode measurements as well as Q-PCR assay demonstrated that the 16S rRNA gene copy numbers of AOB and *Nitrospira*-like NOB and the NH_4^+ consumption rates in the burrow walls were comparable with or higher than those in the sediment surface and in the bulk sediment (especially at a depth of 50-55 mm). These results indicate that infaunal burrows supported higher abundance of nitrifying bacteria and *in situ* NH_4^+ consumption activity in the burrow wall. These results were also consistent with the result of the microcosm experiments showing that increasing benthic infaunal density in the sediment resulted in the increases in NH_4^+ and total inorganic nitrogen consumption rates (**Fig. 1**). Mayer et al. showed that nitrification potentials in tube or burrow walls of 6 types of benthic macrofaunas were significantly greater than those of the surrounding bulk sediments (23). Furthermore, Dollhopf et al. found that coupled nitrification-denitrification was enhanced by macrofaunal burrowing activity in salt marsh sediments (11).

Many studies have aimed to measure solute concentrations in burrows (5, 13, 24, 25, 32, 34). However, all of the data were limited in the upper parts (a few cm depth) of the

sediments. In this study, we constructed and used the aquarium with agar slits in a sideboard to overcome this limitation (see **Fig. S1** in the supplemental material). The O_2 microelectrode was inserted into the center of the burrow at different depths through the agar slits and an O_2 concentration profile along the burrow structure was determined (**Fig. 4A**). A typical horizontal O_2 concentration profile was depicted in **Fig. 4B**. The O_2 concentration in the burrow decreased from 190 μM in the overlying water to 120 μM at a depth of 80 mm, but below which the decrease in the O_2 concentration was moderate (**Fig. 4A**). Thus, approximately 70 μM of O_2 still existed even at a depth of 350 mm. In contrast, O_2 penetration depth was only less than 1 mm below the sediment surface without infaunal burrows (data not shown). This was probably attributed to the infaunal irrigation activity to exchange the water with low concentrations of O_2 and nutrients with fresh water (19).

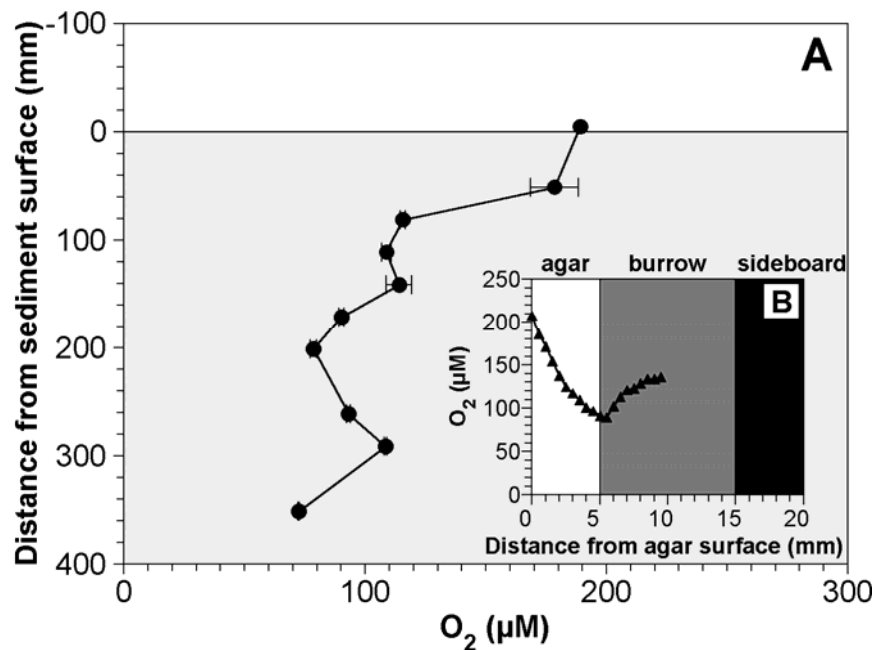


Fig. 4. A vertical O_2 concentration profile along an infaunal burrow measured in the aquarium with slits filled with agar plates in a sideboard (A). Data points were obtained

1 from the horizontal O₂ concentration profiles, as depicted in panel B, at different depths.
2 The values indicate the average O₂ concentrations at the center of the burrow (i.e., 10 mm
3 from agar surface in panel B). The error bars indicate the standard deviations of the
4 measurements for 10 min at each position. A typical horizontal O₂ concentration profile
5 in an infaunal burrow (B). An O₂ microelectrode was inserted into the center of a burrow
6 through agar plate. Zero, 5 mm and 10 mm on the horizontal axis correspond to the agar
7 surface, the agar-burrow interface and the center of the burrow, respectively.

8
9 The advantages of this method were to be able to know the exact position of burrows
10 in sediment and to directly measure the concentration profiles in the burrows at the deeper
11 parts of the sediment. However, O₂ diffused into the burrow and sediment through the agar
12 plate, as indicated by the O₂ concentration gradients in the agar plate (**Fig. 4B**). The O₂
13 transport rates through the agar plate were, therefore, calculated from the O₂ concentration
14 gradients in the agar plate based on Fick's first law of diffusion. The diffusion coefficient for
15 O₂ in the 3% agar used for the calculation was $2.2 \times 10^{-5} \text{ cm}^2 \text{ s}^{-1}$ (16). The mean O₂ transport
16 rate was calculated to be $0.012 \pm 0.001 \text{ } \mu\text{mol cm}^{-2} \text{ h}^{-1}$. This value corresponded to ca. 5% of
17 the total O₂ consumption rates in the burrow walls ($0.19 \text{ to } 0.26 \text{ } \mu\text{mol cm}^{-2} \text{ h}^{-1}$). Thus, O₂
18 concentrations in the burrow would be slightly overestimated in this study. However, this
19 does not negate the general trend in the results presented here.

20 In summary, combination of the 16S rRNA gene-based molecular approach and
21 microelectrode measurements clearly demonstrated that the infaunal burrow facilitated O₂
22 transport into the sediment, which supported the higher abundance and *in situ* activity of
23 nitrifying bacteria in the burrow walls in the intertidal sediment. Further studies with more

quantitative techniques such as fluorescent *in situ* hybridization (FISH) and microelectrodes for N_2O , NO_2^- and NO_3^- are desired to fully understand nitrogen cycling in the bioturbated sediment.

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TABLE 1. Detection frequency and phylogenetic relatives of the AOB clones analyzed at sediment surface (SS) and burrow walls (BW) at 25-30 mm and 50-55 mm from the sediment surface.

Lineage Closest relative	No. of OTUs (No. of clones) in the following layer			
	SS	BW-25	BW-50	Similarity
Total	8 (67)	7 (47)	12 (52)	
<i>Nitrosomonas oligotropha</i> lineage				
<i>Nitrosomonas</i> sp. Is79A3 (AJ621026)			1 (2)	95-98
Unidentified <i>Betaproteobacterium</i> Vm4 (AJ003756)	1 (2)			99
<i>Nitrosomonas marina</i> lineage				
<i>Nitrosomonas</i> sp. Is343 (AJ621032)	1 (5)	1 (4)	1 (4)	97-99
Uncultured bacterium clone AZP2-9 (AY186223)	1 (1)	3 (5)	1 (3)	95-98
<i>Nitrosomonas europaea</i> / <i>Nitrosococcus mobilis</i> lineage				
<i>Nitrosomonas europaea</i> (BX321856)	1 (1)			99
<i>Nitrosomonas communis</i> lineage				
<i>Nitrosomonas communis</i> (AJ298732)			1 (2)	95
<i>Nitrosomonas</i> sp. Nm143 lineage				
Uncultured AOB clone B8s180r (AB212150)	1 (18)	1 (27)	1 (12)	95-99
<i>Nitrosomonas</i> sp. Nm143 (AY123794)	1 (2)		3 (7)	94-96
<i>Nitrosomonas</i> sp. NS20 (AB212171)		2 (2)	1 (4)	93-98
<i>Nitrospira briensis</i> lineage				
<i>Nitrospira multiformis</i> (AY123807)	2 (2)		3(9)	94-97

TABLE 2. Summary of total microbial cell counts and 16S rRNA gene copy numbers of AOB and *Nitrospira*- and *Nitrobacter*-like NOB in the bulk sediment and burrow wall samples.

Sediment depth	Total cell (10^9 cells cm^{-3}) ^a		AOB (10^7 copies cm^{-3}) ^a		<i>Nitrospira</i> (10^7 copies cm^{-3}) ^a		<i>Nitrobacter</i> (10^7 copies cm^{-3}) ^a	
	Sediment	Burrow	Sediment	Burrow	Sediment	Burrow	Sediment	Burrow
0-5 mm (surface)	6.5 ± 0.7	N.D. ^b	2.4 ± 0.3	N.D. ^b	2.8 ± 1.8	N.D. ^b	0.1 ± 0.0	N.D. ^b
5-10 mm	5.4 ± 1.3	1.3 ± 0.3	2.8 ± 1.4	1.2 ± 0.8	5.7 ± 4.1	1.0 ± 0.9	0.1 ± 0.1	0.3 ± 0.0
25-30 mm	4.7 ± 1.0	1.0 ± 0.1	3.0 ± 0.9	3.0 ± 0.5	1.1 ± 0.8	1.4 ± 0.1	0.2 ± 0.1	0.3 ± 0.1
50-55 mm	4.0 ± 0.5	1.9 ± 0.7	0.7 ± 0.2	4.2 ± 1.7	0.6 ± 0.2	3.3 ± 0.2	0.004 ± 0.003	0.2 ± 0.1

^a The values are averages $\pm$ standard deviations.

^b N.D. : not determined.

TABLE S1. Detection frequency and phylogenetic relatives of the *Nitrospira*-like NOB clones analyzed at sediment surface (SS) and burrow walls (BW) at 25-30 mm and 50-55 mm from the sediment surface.

Lineage	No. of OTUs (No. of clones)			
Closest relative	SS	BW-25	BW-50	Similarity
Total	3 (46)	2 (44)	3 (46)	
<i>Nitrospira marina</i> lineage				
Uncultured bacterium clone B58 (AY375077)	1 (43)	1 (43)	1 (41)	96-99
<i>Nitrospira marina</i> (X82559)	1 (2)	1 (1)	1 (4)	98-99
<i>Nitrospira defluvii</i> lineage				
<i>Nitrospira defluvii</i> (DQ059545)	1 (1)			99
<i>Nitrospira</i> sp. clone b2 (AJ224038)			1 (1)	97