

# Denitrifying Alphaproteobacteria from the Arabian Sea That Express *nosZ*, the Gene Encoding Nitrous Oxide Reductase, in Oxic and Suboxic Waters

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Marine ecosystems are significant sources of the powerful greenhouse gas nitrous oxide ( $N_2O$ ). A by-product of nitrification and an intermediate in the denitrification pathway,  $N_2O$  is formed primarily in oxygen-deficient waters and sediments. We describe the isolation of a group of alphaproteobacteria from the suboxic waters of the Arabian Sea that are phylogenetically affiliated with *Labrenzia* spp. and other denitrifiers. Quantitative PCR assays revealed that these organisms were very broadly distributed in this semienclosed ocean basin. Their biogeographical range extended from the productive, upwelling region off the Omani shelf to the clear, oligotrophic waters that are found much further south and also included the mesotrophic waters overlying the oxygen minimum zone (OMZ) in the northeastern sector of the Arabian Sea. These organisms actively expressed *NosZ* ( $N_2O$  reductase, the terminal step in the denitrification pathway) within the OMZ, an established region of pelagic denitrification. They were found in greatest numbers outside the OMZ, however, and *nosZ* mRNAs were also readily detected near the base of the upper mixed layer in nutrient-poor, oxic regions. Our findings provide firm molecular evidence of a potential sink for  $N_2O$  within well-ventilated, oceanic surface waters in this biogeochemically important region. We show that the *Labrenzia*-like denitrifiers and their close relatives are habitual colonizers of the pseudobenthic environment provided by *Trichodesmium* spp. We develop the conjecture that the  $O_2$ -depleted microzones that occur within the colonies of these filamentous, diazotrophic cyanobacteria might provide unexpected niches for the reduction of nitrogen oxides in tropical and subtropical surface waters.

Emissions of the greenhouse gas nitrous oxide ( $N_2O$ ) have increased steadily since the early part of the 19th century. Atmospheric  $N_2O$  concentrations are higher today than at any time during the past 650,000 years (1, 2). Apart from its significant warming potential ( $\sim 300$ -fold that of  $CO_2$  over a 100-year period), rising  $N_2O$  is of further environmental concern because it is presently the single most destructive source of emissions contributing to stratospheric ozone depletion (3). The growing inventory of atmospheric  $N_2O$  has occurred primarily as a result of an increase in emissions from the terrestrial environment owing to changes in agricultural practices, the combustion of fossil fuels, and other anthropogenically driven perturbations of the nitrogen cycle (2). The marine environment is also an important net source of  $N_2O$  to the atmosphere, however, and the unperturbed (non-anthropogenic) rates of emissions from coastal margins, shelf, and open waters are of the same order as those from land (1, 4, 5).

Significant feedbacks on marine  $N_2O$  emissions are anticipated over the coming decades as a result of increasing ocean acidification (5) and the expansion of hypoxic waters owing to surface warming and an acceleration in the prevailing rates of eutrophication from anthropogenic nutrient loading (1, 6). In oxic waters,  $N_2O$  is produced primarily as a by-product of nitrification, the oxidation of ammonium ( $NH_4^+$ ) to nitrite ( $NO_2^-$ ) and nitrate ( $NO_3^-$ ) carried out by ammonium-oxidizing bacteria (AOB) and archaea (AOA). Recent work has shown that ammonium oxidation rates (and hence  $N_2O$  production by AOB/AOA) are exquisitely sensitive to comparatively modest ( $\sim 0.1$ -unit) declines in pH (5) and show a mean reduction of  $\sim 20\%$  in response to near-future (20 to 30 years hence) ocean acidification. The pH of oceanic surface waters is set to decline by 0.3 to 0.4 units by the end of the present century (7) with the potential corollary of large nega-

tive feedbacks on marine nitrification rates and  $N_2O$  emissions in the coming decades (5).

The production of  $N_2O$  by nitrifiers is oxygen sensitive, however, and its emissions increase very substantially under hypoxic conditions (8). Over the past 50 years, the areal extent of hypoxic waters in coastal regions and at intermediate depths in the North Pacific and tropical oceans has expanded and shoaled significantly (9, 10), a trend that will only intensify as the global ocean continues to warm and lose oxygen over the next few decades and beyond (11). Approximately 10% of the contemporary ocean is either hypoxic ( $O_2$ ,  $<30\%$  saturation) or suboxic ( $O_2$ ,  $<1\%$  saturation), and even a modest expansion in the present volume of deoxygenated waters is likely to increase  $N_2O$  production by nitrifiers significantly (1, 6). The net impacts of increasing atmospheric  $CO_2$  on marine  $N_2O$  emissions in the years to come, therefore, will depend on the overall balance of both positive and negative feedbacks on ammonium oxidation rates and other climate-sensitive sources of  $N_2O$ .

The predominant source of  $N_2O$  emissions under suboxic conditions is not from ammonium oxidizers but from a taxonomically diverse group of mainly heterotrophic microbes known as denitrifiers (6). Denitrifiers use  $NO_3^-$  as an alternative respiratory

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electron acceptor to oxidize an electron donor (usually organic carbon) to generate metabolic energy during anaerobic growth. Denitrification leads ultimately to the liberation of dinitrogen gas ( $N_2$ ) in a four-step reductive pathway from  $NO_3^-$  that proceeds via  $NO_2^-$  and two obligate gaseous intermediates, nitric oxide (NO) and  $N_2O$ . Not all denitrifiers are capable of the final reduction of  $N_2O$  to  $N_2$ , however, while for others a sustained lag occurs during the induction of the necessary cellular machinery to carry out this terminal step in the process (12). Some denitrifiers produce  $N_2$  only in the complete absence of free oxygen but, nonetheless, are capable of partial denitrification and liberate  $N_2O$  under hypoxic conditions (13). Denitrification is both a sink and a potential source of emissions, therefore, and another key process in the marine nitrogen cycle known to be sensitive to the future expansion in the volume of deoxygenated waters in the ocean (1, 6).

The only known metabolic pathway for the consumption of  $N_2O$  is that which requires the copper-containing enzyme nitrous oxide reductase (NosZ). NosZ is found in all denitrifiers that are capable of reducing  $NO_3^-$  as far as  $N_2$  and also in a few nondenitrifying bacteria, such as *Wolinella* (*Vibrio succinogenes*), that can use  $N_2O$  as a terminal electron acceptor (14). The latter organisms are not themselves  $N_2O$  producers, however, because they lack the NO-producing nitrite reductase (NirK or NirS) that is the definitive biochemical feature of denitrifiers (15), the primary sink for this nitrogen oxide in the oceans (16).

Uncertainties over the future scale and climatic impact of marine  $N_2O$  emissions and, in particular, their sensitivities to acidification and deoxygenation (17) have led to calls for a better understanding of the biological sources and sinks of this trace gas in the oceans (1, 18). Key to constraining the marine  $N_2O$  budget is the development of a robust model of the distributions and activities of the organisms that produce and/or consume  $N_2O$  and their responses to stress induced by global environmental change. Approximately half of the annual emissions of  $N_2O$  in the contemporary ocean come from the three major oxygen minimum zones (OMZs) that are located in the eastern tropical North Pacific (ETNP), the eastern tropical South Pacific (ETSP), and the Arabian Sea (19, 20). The net contributions of different biological sources of  $N_2O$  in these regions are debated (see reference 18), but there is general agreement that the OMZs are “hot spots” within the ocean that are especially vulnerable to warming and deoxygenation over the coming decades (1, 17).

In the present study, we describe the isolation into culture of a novel group of pelagic denitrifying alphaproteobacteria from the suboxic waters of the Arabian Sea, one of the most intense regions of  $N_2O$  production in the world ocean (20). Employing a quantitative PCR protocol targeting the *nosZ* gene from these organisms, we show that they have a broad biogeographical distribution in this ocean basin, ranging from the upwelling region along the Omani shelf to the highly oligotrophic equatorial waters to the south. We further show that these organisms were expressing the cognate mRNA for *nosZ* in the suboxic intermediate waters of the OMZ located in the northeastern sector of the Arabian Sea and, quite unexpectedly, were also active at shallower depths within the upper mixed layer of waters well outside the region of the OMZ.

## MATERIALS AND METHODS

**Study site, sample collection, and nucleic acid purification.** Observations were made in September 2001 aboard RRS *Charles Darwin* during

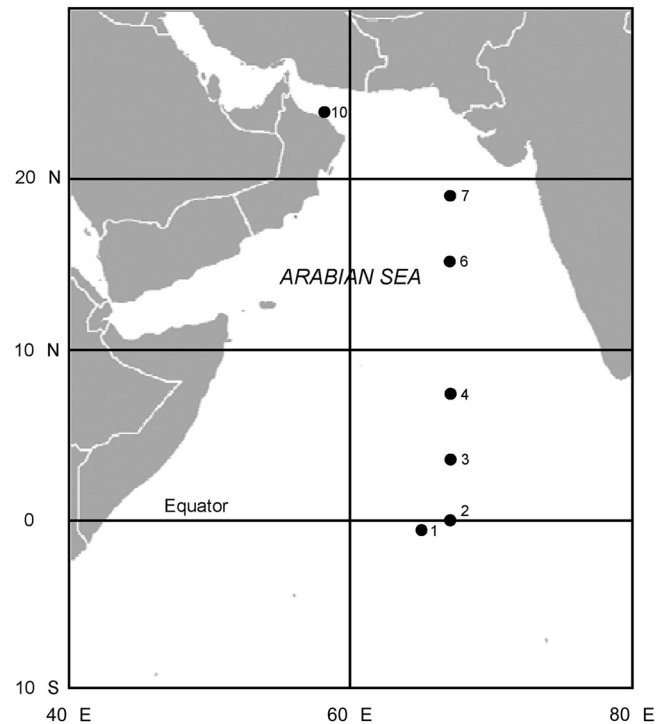


FIG 1 Study area in the Arabian Sea showing the locations and designated station numbers of the sites sampled during the south-to-north AMBITION cruise transect in September 2001.

the Natural Environment Research Council (NERC)-funded AMBITION cruise (CD132) in the Arabian Sea. Eleven stations were occupied along the length of a 5,150-km transect between Victoria, Seychelles, and Muscat, Oman (21). Hydrographical data were collected at each station with a Sea-Bird 911plus conductivity-temperature-depth (CTD) profiler (Sea-Bird Electronics, Inc., Bellevue, WA) configured with a Chelsea Aquatracker III fluorometer (Chelsea Instruments, West Molesey, United Kingdom) to measure chlorophyll fluorescence and auxiliary sensors for dissolved oxygen and photosynthetically active radiation (PAR).

Seawater samples were collected from discrete depths at selected stations (Fig. 1) with a rosette of 24 30-liter volume Niskin bottles mounted on the CTD profiler. Plankton samples were obtained by filtering 4 to 5 liters of seawater (unscreened) through 90-mm-diameter, 0.2- $\mu$ m-pore-size polycarbonate filters (Osmonics Inc., Minnetonka, MN) at a negative pressure of <20 mm Hg. Cell material collected on the filters was taken up in DNA isolation buffer (250 mM NaCl, 100 mM EGTA, 100 mM Tris-HCl [pH 8.0], and 1% [wt/vol] lithium dodecyl sulfate) and stored frozen at  $-70^{\circ}C$  or, for RNA samples, preserved in RNAlater (Applied Biosystems, Warrington, United Kingdom) and refrigerated at  $4^{\circ}C$ . At the end of the cruise, the preserved nucleic acid samples were shipped by air to the United Kingdom on dry ice and subsequently stored at  $-80^{\circ}C$  prior to the extraction and purification of DNA and RNA as described by Bird et al. (21).

**Plankton net hauls for the collection of *Trichodesmium* colonies and extraction of DNA.** Plankton samples were also collected from depths of 5 or 10 m at each station during short (10- to 15-min) horizontal hauls using a standard WP2 conical net (200- $\mu$ m mesh size) towed at a ship speed (through the water) of 1 knot. *Trichodesmium* colonies were sorted into sterile, filtered (0.2- $\mu$ m-pore-size polycarbonate membranes) surface seawater using a sterile, disposable inoculating loop, rinsed in two changes of sterile seawater, and transferred to 0.5 ml RNAlater prior to storage at  $4^{\circ}C$ . DNA was extracted from the sorted *Trichodesmium* colonies (~5 colonies per extraction) after rinsing twice in sterile artificial

TABLE 1 Oligonucleotide primers used in this study

Oligonucleotide	DNA sequence (5'–3')	Product size (bp)	Reference
nosZF1	GAYGTNCANTAYCARCCNGGNCA	612	This study
nosZR	CATYTCCAARTGNADNGCRTGRCA	612	This study
StanosF	GCTGAAACCCGGAGAATGAAC	252	This study
StanosR2	TGCATGTAGACACGCACCTT	252	This study
QnosF	GAAGTTCACGGTAAAGCAGG	98	This study
QnosR	CATAGTTGGACAGGCAGAAG	98	This study
Probe	ACGAAGTCACCATCTACGTCACCAACA	98	This study
nirS1F	CCTAYTGGCCGCCRCART	~890	24
nirS6R	CGTTGARCTTRCCGGT	~890	24

seawater (ASW) medium (22) to remove RNA later. The washed colonies were collected on 0.2- $\mu\text{m}$ -pore-size polycarbonate membranes and then lysed in TE buffer (10 mM Tris-HCl, pH 8.0, 1 mM EDTA) amended to 0.1% Triton X-100 and 0.2 mg ml<sup>-1</sup> lysozyme (Sigma catalog no. L7651) at 37°C for 15 min. The lysates were brought to 50  $\mu\text{g}$  ml<sup>-1</sup> proteinase K (Roche) and 0.2% SDS and incubated at 56°C for a further 10 min before purification of DNA using the Qiagen DNeasy kit according to the supplier's abridged protocol for the purification of DNA lysates.

**Enrichment culture for the isolation of nitrate-respiring (denitrifying) bacteria from the Arabian Sea.** Seawater was collected from a depth of 120 m in suboxic waters at station 10 (24°19'N, 58°10'E; depth, 2,863 m) (Fig. 1) and amended with 1.5 g liter<sup>-1</sup> tryptic soy broth (Thermo Fisher Scientific, Basingstoke, United Kingdom). The amended seawater was further supplemented with 1 $\times$  Guillard's (F/2) nutrient and vitamin solution (Sigma, Poole, United Kingdom) and incubated in filled, airtight, sterile bottles at 25°C. Following the visible establishment of bacterial growth after 3 to 5 days, subcultures were transferred aseptically to de-gassed (boiled), filter-sterilized seawater from the same station amended with 0.5 g liter<sup>-1</sup> NaNO<sub>3</sub>, 1.0 g liter<sup>-1</sup> NH<sub>4</sub>Cl, 0.006 g liter<sup>-1</sup> FeCl<sub>2</sub>, 0.001 g liter<sup>-1</sup> EDTA, 0.003 g liter<sup>-1</sup> K<sub>2</sub>HPO<sub>4</sub>, and 1 ml liter<sup>-1</sup> A5 trace metal mixture (23) and 1 g liter<sup>-1</sup> sodium acetate. The subcultures were incubated in filled, airtight, sterile bottles at 25°C until the medium turbidity increased (7 to 14 days). After two further rounds of subculture, the bacterial suspensions were transferred to streak plates of the same medium solidified with 10 g liter<sup>-1</sup> Difco Bacto agar (Becton, Dickinson and Co., Oxford, United Kingdom). The inoculated plates were incubated at 25°C in sealed, anaerobic jars in an oxygen-free nitrogen atmosphere. Individual colonies were isolated by repeated subculture on agar streak plates and maintained thereafter at 25°C under these same incubation conditions.

**PCR amplification of *nosZ*, *nirS*, and 16S rRNA genes from bacteria isolated from the Arabian Sea.** Independent isolates of putative, nitrate-respiring bacteria were grown aerobically at 25°C for 24 to 48 h in 3 ml ASW medium (22) amended with 10% (vol/vol) Luria-Bertani broth (10 g liter<sup>-1</sup> tryptone, 5 g liter<sup>-1</sup> yeast extract, 10 g liter<sup>-1</sup> NaCl) in an orbital incubator at 180 rpm. The cultures were harvested by centrifugation at 16,000  $\times$  g for 2 min, and DNA was isolated from the pelleted cell material using a DNeasy tissue kit according to the supplier's (Qiagen Ltd., Crawley, United Kingdom) recommended protocol for bacteria. The DNA samples were screened for the presence of *nosZ* by PCR using Thermo-prime DNA polymerase master mix (Fisher Scientific, Loughborough, United Kingdom) in reaction volumes of 25  $\mu\text{l}$  containing 2 mM MgCl<sub>2</sub>, 10 ng DNA, and 50 pmol each of the primers nosZF1 and nosZR (Table 1).

The degenerate primers nosZF1 and nosZR were designed to target the conserved motifs DV(H/Q)YQPGH and CHA(M/I/L)H(L/M)EM identified in the majority of complete *NosZ* sequences available from the GenBank database at the start of this study and correspond to nucleotide positions 1276 to 1298 and 1864 to 1887 of *nosZ* from *Pseudomonas stutzeri* (ZoBell strain) ATCC 14405 (GenBank accession no. CAA37714.2), respectively. After denaturation at 95°C for 2 min, the cycling conditions were 94°C for 30 s, 55°C for 30 s, and 72°C for 1 min for 25 cycles followed by a final extension at 72°C for 10 min.

The PCR products obtained were resolved in 1% (wt/vol) agarose gels and purified using the Wizard SV gel and PCR cleanup system (Promega Ltd., Southampton, United Kingdom) before TA cloning in the plasmid vector pCR-TOPO as recommended by the supplier (Invitrogen, Paisley, United Kingdom). The cloned PCR products from four independent isolates (designated 1N, 4N, 5N, and 8N) were DNA sequenced on both strands using M13 forward and reverse primers in parallel reactions performed with a DYEnamic ET terminator cycle sequencing kit (GE Healthcare Life Sciences, Little Chalfont, United Kingdom) and an ABI Prism 377 automated sequencer.

An ~890-bp fragment of the gene encoding NirS (nitrite reductase) was amplified from genomic DNA purified from the Arabian Sea isolate designated 4N using the primer pair nirS1F and nirS6R (24) (Table 1). The reactions were carried out as described above for *nosZ* except that the annealing temperature was increased to 58°C. Genomic DNA from this isolate was also used to amplify a fragment of the 16S rRNA gene with the universal primer pair 27F and 1492R (25) under similar reaction conditions but with a lower annealing temperature of 48°C and an extension step of 90 s. The *nirS* and 16S rRNA PCR products obtained were gel purified, TA cloned, and sequenced either in-house (16S rRNA) as described for *nosZ* above or, in the case of *nirS*, by a commercial provider (Source BioScience Lifesciences, Nottingham, United Kingdom) using M13 primers.

**Amplification of *nosZ* from *Trichodesmium* consortia by the PCR.** Aliquots of DNA (1  $\mu\text{l}$  from 100  $\mu\text{l}$  of extract) isolated from *Trichodesmium* colonies collected at three stations (stations 2, 3, and 4 [Fig. 1]) were interrogated for the presence of *nosZ* closely related to that of the Arabian Sea alphaproteobacteria described in this study (including the isolate designated 4N) using a targeted PCR protocol. Each 25- $\mu\text{l}$  reaction volume contained 1 $\times$  Thermoprimase DNA polymerase master mix (Thermo Fisher Scientific, Loughborough, United Kingdom) amended to 2 mM MgCl<sub>2</sub>, 0.1 mg ml<sup>-1</sup> bovine serum albumin (Promega Corp., Southampton, United Kingdom), and 50 pmol each of the primers StanosF and StanosR2 (Table 1). The primers target a 252-bp internal region of *nosZ* corresponding to nucleotides 97 to 348 of the trimmed sequence of the Arabian Sea isolate 4N (GenBank accession no. JN850958). The PCR cycling conditions consisted of an initial denaturation at 95°C for 2 min followed by 30 cycles at 94°C for 30 s, 52°C for 30 s, and 72°C for 45 s and a final extension at 72°C for 10 min.

The PCR products were resolved by electrophoresis through 1.5% (wt/vol) NuSieve 3:1 agarose gels (Lonza, Stratech Scientific, Newmarket, United Kingdom), and a fragment of the expected size from the station 3 sample was excised from the gel, purified, and TA cloned as described above. The inserts from 10 of the resulting recombinant clones were screened by digestion with the endonuclease SphI (which has a predicted recognition sequence centered at nucleotide position 105 within the *nosZ* fragment targeted by the primers), and all 10 clones were found to produce digestion products of the predicted size when resolved on 2.0% (wt/vol) NuSieve 3:1 agarose gels. One of these clones was sequenced on both strands by a commercial provider (Source BioScience Lifesciences, Nottingham, United Kingdom) to confirm its identity as described above.



To examine the wider distribution of *nosZ* in *Trichodesmium* consortia, DNA samples purified from colonies collected in surface waters near the Bahama Islands in 1991 and from a cultured isolate (strain IMS101) were also analyzed by PCR, and the products obtained were cloned and sequenced as described above.

**Detection and enumeration of *nosZ* DNA in natural samples of plankton from the Arabian Sea.** The depth and latitudinal distribution of bacteria in the Arabian Sea that harbored a *nosZ* gene identical to or highly similar in sequence to that amplified from the cultured isolates were investigated by quantitative PCR (QPCR). DNA samples obtained from four stations (designated 1, 4, 6, and 10 [Fig. 1]) occupied during the AMBITION cruise were assayed using a double dye-labeled oligonucleotide probe and the primer pair QnosF and QnosR (Table 1). The primers amplify a product of 98 bp and bind at nucleotide positions 376 to 395 and 454 to 473, respectively, within the original (trimmed) *nosZ* PCR products amplified from the cultured isolates. The probe, which was coupled with fluorescein at the 5' end and Elipse dark quencher at the 3' end (Eurofins MWG Operon, Ebersberg, Germany), binds nucleotide positions 399 to 425 bp starting 4 bp downstream of the 3' end of the forward primer QnosF.

QPCR assays were performed with a Stratagene Mx3000p thermocycler and Brilliant II reagents (Agilent Technologies Ltd., Wokingham, United Kingdom) in 25- $\mu$ l volumes containing 1.0  $\mu$ l of DNA, 25 pmol of each primer, and a 150 nM concentration of probe DNA. Following activation at 95°C for 10 min, reaction mixtures were cycled at 95°C for 30 s, 60°C for 30 s, and 72°C for 45 s for 40 cycles and data were collected at the end of the extension step. The threshold cycle ( $C_T$ ) was determined automatically by the instrument software, and the initial template quantity (*nosZ* copies) was calculated with reference to a standard curve constructed using a 10-fold dilution series of linear fragments of the cloned *nosZ* amplicon from the isolate 4N.

Standard DNA was amplified from the original plasmid clone (pnos4N) using the primer pair nosZF1 and nosZR as described above, and the concentration of the gel-purified 612-bp product was determined using a Picodrop 100 microspectrophotometer (Cambridge Bioscience, Cambridge, United Kingdom). Copy number was estimated from the DNA concentration of the standard by assuming that 1 bp has an average molecular mass of 650 Da and using the following expression: copy number =  $(N \times 6.022 \times 10^{23}) / (L \times 1 \times 10^9 \times 650)$ , where  $N$  is the DNA amount in ng and  $L$  is the length of the DNA fragment.

The lower detection limit of *nosZ* standard DNA was routinely between 10 and 100 copies or better, and PCR efficiency was 98.3% ( $r^2 = 0.998$ ). The absence of PCR inhibitors in the natural sample DNA preparations was verified by spiking parallel reaction mixtures with  $10^5$  copies of the target DNA fragment. Those samples in which the  $C_T$  of the spiked controls was  $>1.5$  cycles greater than that predicted were excluded from further analysis.

**Quantitative reverse transcriptase PCR (QRT-PCR).** RNA samples (0.5  $\mu$ g) were treated with Ambion Turbo DNase (Applied Biosystems, Warrington, United Kingdom) and reverse transcribed using a Quantitect reverse transcriptase (RT) kit (Qiagen Ltd., Crawley, United Kingdom) and random hexamer primers following treatment with Genomic Wipe-out reagent as recommended by the suppliers. cDNAs originating from *nosZ* mRNA were quantified by QPCR as described above using 1-in-50 dilutions of the RT reaction mixtures and run in parallel with minus-RT controls and no-template blanks. PCR efficiency was 97.7% ( $r^2 = 0.999$ ).

To verify the specificity of the assays, the RT-PCR products amplified from the 50-m depth sample from station 2 were resolved through a 2.5% (wt/vol) NuSieve 3:1 agarose gel and the 98-bp product was purified using the Wizard SV gel and PCR cleanup system. The purified product was screened with the endonucleases EcoRV and NcoI (which have recognition sequences centered at nucleotide positions 58 and 75, respectively, within the 98-bp amplicon) and found to produce digestion products of the expected sizes. The RT-PCR amplicons were TA cloned as described above, and three randomly selected clones were DNA sequenced. The

cloned inserts were found to contain the internal region of *nosZ* targeted by the QPCR primers and to share 98 to 100% identity with *nosZ* from the Arabian Sea isolate 4N.

Experiments ( $n = 3$ ) were also conducted to examine the effect of  $pO_2$  on *nosZ* mRNA abundance in cultures of the Arabian Sea isolate 4N grown at 25°C in the acetate-containing medium described above under aerobic conditions ( $O_2$  saturation,  $\geq 95\%$ ) or after transfer to a suboxic ( $O_2$  saturation,  $\leq 1\%$ ) atmosphere for 30 min prior to sampling. RNA was extracted, reverse transcribed, and assayed for *nosZ* by QPCR as described for the natural samples, and the products were run out on 2.5% (wt/vol) NuSieve 3:1 agarose gels. Dissolved oxygen concentrations were determined using a calibrated oxygen electrode (Hansatech Instruments Ltd., Kings Lynn, United Kingdom).

**Phylogenetic reconstruction.** The DNA sequences of *nosZ* and *nirS* from the Arabian Sea isolate 4N were trimmed of primer sequences and translated *in silico*. Peptide sequences closely related to each target gene were identified using the Basic Local Alignment Search Tool (BLAST) at NCBI (26) and aligned in MUSCLE with those from the isolate 4N using the unweighted-pair group method using arithmetic (UPGMA) clustering method (27). Evolutionary analyses were performed in MEGA5 (28) using the maximum likelihood method based on the Dayhoff matrix model. The bootstrap consensus trees produced in each case were based on 500 replicate samples.

**Nucleotide sequence accession numbers.** The DNA sequences of *nirS*, *nosZ*, and the 16S rRNA gene from the Arabian Sea isolate designated 4N have been deposited in GenBank under the accession numbers JX827176 (*nirS*), JN850958 (*nosZ*), and JN850957 (16S rRNA). The DNA sequence of the *nosZ* fragment amplified from *Trichodesmium* colonies collected at station 3 is deposited in GenBank under accession number JX827177, and that from the cultured isolate strain IMS101 is deposited under accession number KC205088.

## RESULTS AND DISCUSSION

### Isolation of putative denitrifying bacteria from the Arabian Sea and characterization of the representative isolate, strain 4N.

Liquid enrichment cultures of bacteria from the Arabian Sea that were capable of utilizing acetate as their sole carbon source and nitrate as an alternative electron acceptor were successfully established under suboxic growth conditions. This enrichment culture medium was designed to select for the growth of nitrate-respiring bacteria (and denitrifiers, in particular). With the notable exception of methanogens (29), acetate is a nonfermentable carbon substrate for other bacteria. After transfer to a solid medium under an oxygen-free atmosphere, the growth of individual colonies of putative, nitrate-respiring (denitrifying) bacteria was observed after 7 to 10 days. Four of these were selected for further experimentation and brought into axenic, clonal culture using the same medium and growth conditions.

Following their establishment, all isolates produced very similarly sized, round, smooth, shiny colonies that developed a pink/reddish-brown coloration at their center as they matured. All of the isolates were found to be Gram negative and displayed an elevated requirement for  $Na^+$  and  $Cl^-$  but not for either  $Mg^{2+}$  or  $Ca^{2+}$ , i.e., the isolates were halophytic rather than obligately marine (30). In broth culture, the isolates grew primarily as individual, actively motile cells that produced gas (nitrogen) when grown under anaerobic conditions in the acetate-nitrate enrichment medium described above. The ability of the isolate designated 4N to denitrify was subsequently confirmed by the acetylene block technique (15) using the same medium and incubation conditions.

PCR primers were designed to screen DNA extracted from all four isolates for the presence of *nosZ* (and *nirS*, see below). The forward nosZF1 primer targets a motif within region H4 of ni-

trous oxide reductase that includes a highly conserved histidine residue, while the reverse primer, nosZR, targets a region close to the C terminus that includes several ligands for the Cu<sub>A</sub> center of the enzyme (31) and overlaps the coding region targeted by the nondegenerate primer Nos2230R designed by Scala and Kerkhof (32). The primers should amplify DNA from the majority of nosZ sequences deposited in the GenBank database to date, but they are not completely universal. In particular, they are unlikely to recognize nosZ from those bacteria in which a serine rather than a histidine codon occurs adjacent to the 3' end of the reverse primer target. This substitution occurs most frequently in a subset of NosZ proteins that carry a C-terminal extension of about 200 residues and which also show variability in the conserved motif targeted by the forward primer (33, 34). In this instance, however, a PCR product of the expected size (~612 bp) was amplified from DNA extracted from all four axenic Arabian Sea isolates screened.

The DNA sequences of the cloned amplicons from all four isolates were virtually identical (>99%), and a BLASTN search of the GenBank database confirmed that they contained the specific region of nosZ targeted by the primers. The most closely related DNA sequence (~95% nucleotide identity) to that from all four Arabian Sea clones was from the cultured isolate *Labrenzia aggregata* IAM 12614, a rosette-forming, aerobic, denitrifying, marine heterotroph isolated from a sediment sample from the western Baltic Sea (35). All of the nucleotide differences (29/565 bases) that occur between nosZ in this species and that from the isolate designated 4N (which we studied in the most detail) were located in the third codon position, however (data not shown), and their derived peptide sequences were identical.

Besides *L. aggregata* IAM 12614, phylogenetic analysis of the peptide sequence from the Arabian Sea isolate 4N showed that it also clustered with NosZ from several other known marine, denitrifying alphaproteobacteria (Fig. 2a). These included *Nisaea denitrificans*, originally isolated from the Mediterranean Sea (36), and also *Ruegeria pomeroyi*, *Roseobacter* sp. strain SK209-2-6, and *Roseobacter denitrificans* among members of the widely distributed *Roseobacter* clade (37, 38). The two most closely related environmental sequences (clones 5\_16 and 5\_75) originated from the same coastal sediment in the East China Sea (Fig. 2a) and shared a similar level of amino acid identity (between 86 and 90%) with *L. aggregata* IAM 12614 as they did with *Polymorphum gilvum*, a hydrocarbon-utilizing halophile that shares significant genome synteny with members of the genus *Labrenzia* (39).

Phylogenetic analysis of NirS peptide sequences showed that the closest known sequence to that obtained from the isolate 4N was also from *L. aggregata* IAM 2614 (Fig. 2c), with which it shares 93.2% and 98% nucleotide and amino acid identity, respectively. Both the *L. aggregata* IAM 2614 and the isolate 4N NirS sequences were very closely related to two environmental clones (designated SdnirS9 and AgnirS-1) originally retrieved from shallow water (20-m depth) sponges in the East China Sea and (somewhat more distantly) to numerous benthic environmental sequences that clustered with *R. pomeroyi* and other members of the *Roseobacter* clade (Fig. 2c). Two clones of pelagic origin (clones M60-80 and M60-134) that were obtained from the Baltic Sea (35) were also paired within this wider cluster of sequences but shared only 90 to 91% amino acid similarity with NirS from isolate 4N.

To confirm the most likely taxonomic affiliation of the Arabian Sea isolate 4N, a BLASTN search of the GenBank database was also conducted with the 16S rRNA gene (1,405 bp) from this isolate.

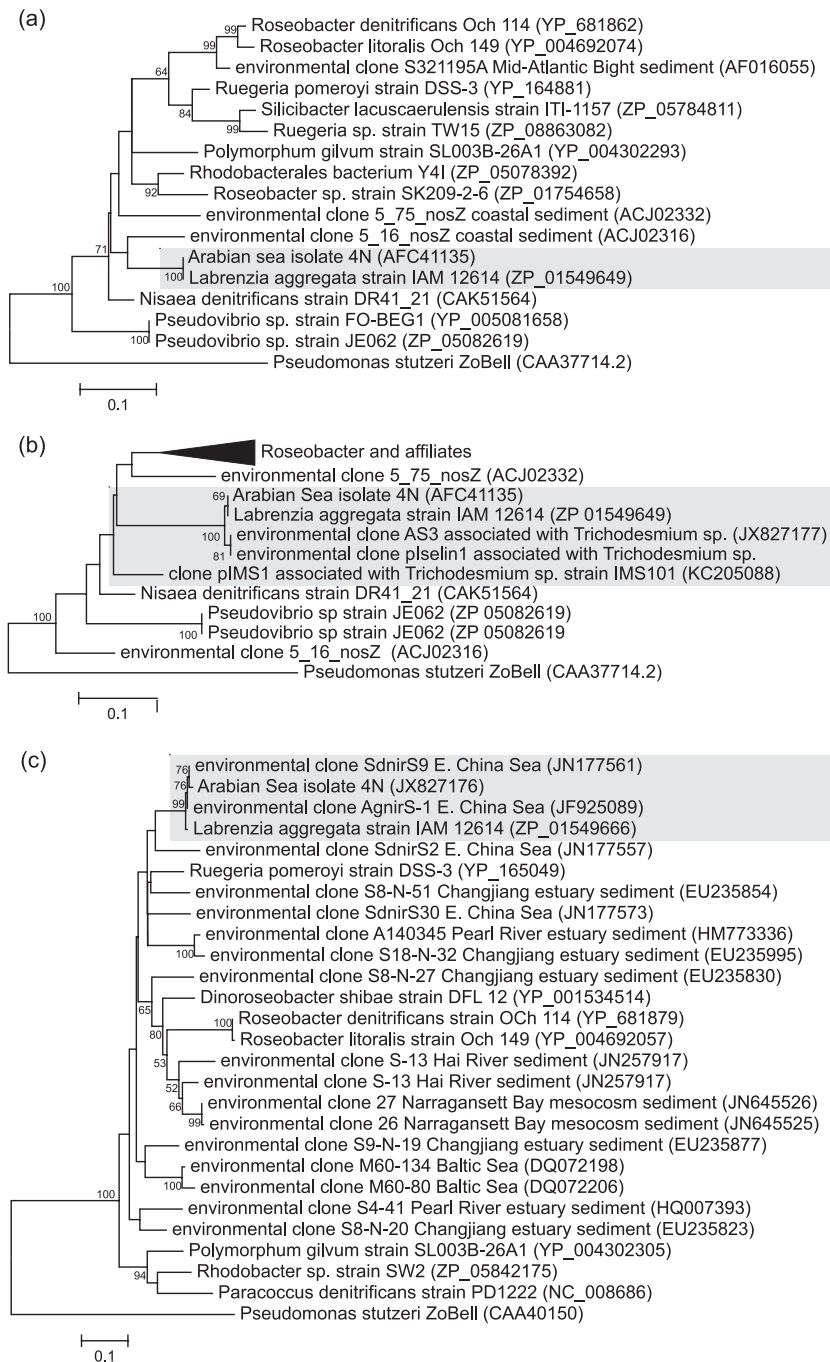
This search revealed that it was >99% identical to 16S rRNA gene sequences from several *Labrenzia* strains isolated from a range of saline habitats and also to some unclassified symbiotic or epibiotic bacteria associated with marine invertebrates or phytoplankton (Table 2). Among the top-scoring matches was the 16S rRNA gene from *L. aggregata* strain 2PR58-2, which its GenBank entry (accession no. EU440961) descriptor reveals was isolated from the deep water column of the southwest Indian Ocean. The remaining high-scoring *Labrenzia* 16S rRNA gene sequences were mostly from isolates obtained from benthic habitats, as was the case for the source of the majority of the most closely related environmental NirS sequences also (see above).

The consensus view of the combined molecular data, therefore, is that the Arabian Sea isolate 4N (and, in all probability, isolates 1N, 5N, and 8N) is most likely to be a new member of the genus *Labrenzia* (40), although this proposed taxonomic affiliation awaits further detailed systematic investigation.

**Spatial distribution of *Labrenzia*-like denitrifying bacteria in the Arabian Sea.** The depth distribution of bacteria closely related to the new Arabian Sea isolates was assayed by QPCR at four stations along the AMBITION cruise track (Fig. 1). The stations were located in waters of markedly contrasting biogeochemical regimes that included highly oligotrophic waters at the southern end of the transect (stations 1 and 4), where primary production was dominated by *Prochlorococcus* spp.; a more mesotrophic transition zone north of 8°N (station 6), in which *Synechococcus* spp. predominated (41); and the nutrient-rich shelf waters along the southeastern coast of Oman (station 10), where the biota was impacted by seasonal upwelling. Monsoonal winds that blow from the southwest between June and September each year lead to a pronounced shoaling of the nutricline in these more northerly waters. At station 10, the nutricline was located at a depth of 25 to 30 m, whereas in the waters further to the south it was typically found at depths of 50 to 70 m (42). Mirroring the latitudinal gradient in nutrient inventories, peak chlorophyll concentrations increased from south to north and were in excess of 6 mg m<sup>-3</sup> at station 10 (Fig. 3).

Oxygen-depleted waters were encountered below the surface mixed layer at all stations (Fig. 3), and the OMZ (<5 μM O<sub>2</sub>) was located between 11 and 24°N at depths below ~100 m (43). Nitrite maxima coincident with the depths of the chlorophyll maximum were found at all stations surveyed, whereas a secondary nitrite maximum (SNM) (>3 μM), located at ~150 m and below, was present only in the central region between 11 and 19°N (M. Woodward, personal communication). The occurrence of a secondary nitrite maximum is often considered to be indicative of active denitrification (44). A recent report has challenged this orthodoxy, however, and its authors (45) measured only low, sporadic N losses within the SNM of the Arabian Sea OMZ at a time of year (September/October 2007) similar to that in the present study.

The *Labrenzia*-like nosZ gene was detected at all stations sampled along the length of the transect, but unexpectedly, the highest mean concentration (4.9 × 10<sup>5</sup> copies liter<sup>-1</sup>) was found at a depth of 300 m at station 1, well outside the OMZ (Fig. 3). More typically, nosZ from these organisms was present at mean concentrations of ~1 × 10<sup>5</sup> copies liter<sup>-1</sup> or fewer at this and the other three stations sampled. Best estimates suggest, therefore, that these denitrifiers were only moderately abundant at the time of sampling. Nonetheless, their very broad biogeographical distribu-



**FIG 2** Phylograms for *nosZ* (a and b) and *nirS* (c) based on partial peptide sequences (188, 83, and 283 residues, respectively) from the Arabian Sea isolate 4N and closely related alphaproteobacterial genes. The maximum likelihood trees were inferred from 500 bootstrap replicates in MEGA5 (28) and rooted with the appropriate orthologue from the ZoBell strain of *Pseudomonas stutzeri* (ATCC 14405). Those partitions that received greater than 50% support in the bootstrap are indicated at their corresponding nodes. The GenBank accession numbers of the sequences are indicated after the sequence names, and the distance scale is shown to the bottom left of each phylogram. The shaded regions highlight the clusters within which genes from the Arabian Sea isolate 4N are found in each phylogram. Note that the highlighted sequence designated pIselin1 in panel b was obtained from *Trichodesmium thiebautii* DNA collected in 1991 from near-surface waters in the Bahama Islands aboard RV *Columbus Iselin* during cruise 91-11 (84) and was identical to that of clone AS3 obtained from station 3 in the Arabian Sea (GenBank accession number JX827177). The template used to amplify the *nosZ* clone pIMS1 from *Trichodesmium* strain IMS101 was from a DNA sample originating from the laboratory of Jon Zehr (University of California, Santa Cruz) and was supplied by Jon G. Kramer of the Center of Marine Biotechnology, University of Maryland, in late 1997.

tion is indicative of a lifestyle that is highly adaptable given the extremes in physical forcing that are the signature feature of the Arabian Sea region (46).

Paradoxically, the overall abundance of the *Labrenzia*-like

*nosZ* gene detected within the OMZ region at station 6 was lower than that encountered elsewhere. Two maxima ( $\sim 1 \times 10^4$  copies liter<sup>-1</sup>) were present: the first located in the oxygenated, surface mixed layer at  $\sim 45$  m and coincident with the chlorophyll maxi-



**TABLE 2** Taxonomic affiliations, environmental sources, and percent identities of 16S rRNA genes from cultivated and environmental sources closest in sequence to that of the Arabian Sea isolate clone 4N

Organism	Source	Identity, no. of identical nucleotides/total no. of nucleotides (%)	Accession no.
<i>Labrenzia</i> sp. strain PEB02	Microbial mat, Ebro delta, Spain	1,401/1,405 (99.72)	GU213158
<i>Labrenzia</i> sp. strain w-2-2	<i>Karlodinium micrum</i> (Dinophyceae)	1,401/1,405 (99.72)	GQ495021
<i>L. aggregata</i> strain 2PR58-2	Southwest Indian Ocean deep-sea water	1,401/1,405 (99.72)	EU440961
Bacterium K2-26	Lake Kauhako, Hawaii	1,401/1,405 (99.72)	AY345441
<i>Labrenzia</i> sp. strain PEB05	Microbial mat, Ebro delta, Spain	1,400/1,405 (99.64)	GU213161
Symbiont CV812-530	Eastern oyster ( <i>Crassostrea virginica</i> )	1,390/1,394 (99.71)	AF246614
<i>L. aggregata</i> IAM 12614	Baltic Sea, sediment	1,397/1,405 (99.43)	D88520
<i>Labrenzia</i> sp. strain MK16	Salt marsh, southwest South Korea	1,386/1,395 (99.35)	AY690722
Epibiont alphaproteobacterium PM05	<i>Pseudo-nitzschia multiseriata</i> (Bacillariophyceae)	1,365/1,377 (99.12)	AY548766

imum and a second that occurred within the SNM at a depth of 200 m (Fig. 3). The total numbers of heterotrophic denitrifiers within the SNM of the Arabian Sea OMZ can exceed  $1 \times 10^8$  cells liter<sup>-1</sup> (47), although more modest estimates of their concentrations ( $\sim 1.5 \times 10^4$  liter<sup>-1</sup>) have been reported recently (45). Such wide variability in population size is most probably related to the episodic nature of the supply of organic carbon to support denitrification (45, 47), and it might be that the *Labrenzia*-like organisms are more numerous and dynamic contributors to the denitrifying assemblage within the OMZ at other times of the year.

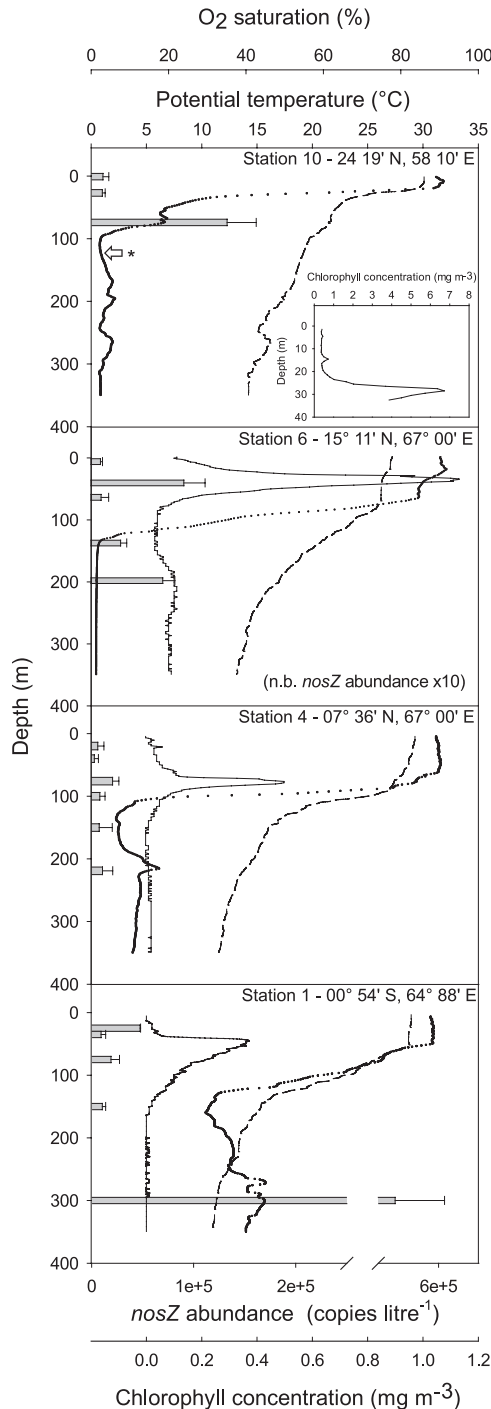
**Transcription of *nosZ* by *Labrenzia*-like denitrifying bacteria within the OMZ of the Arabian Sea.** The distribution of these organisms at station 6 (and the others sampled outside the OMZ) suggests that they do not operate exclusively as denitrifiers, since they were clearly not confined to the suboxic intermediate waters of the OMZ where active denitrification has been demonstrated experimentally (47). Not surprisingly, therefore, *nosZ* mRNAs from these organisms were below the limit of detection by QRT-PCR at three depths (4, 15, and 35 m) within the upper mixed layer at this station (data not shown). Deeper RNA samples from within the core of the OMZ were not available from this particular station, but *nosZ* transcripts were readily detected ( $9.844 \times 10^3 \pm 1.56 \times 10^3$  copies liter<sup>-1</sup>;  $n = 3$ ) in a 220-m-depth sample from a site (station 7) with a similar oxygen deficit ( $O_2$  at 220-m depth =  $3.1 \mu\text{mol liter}^{-1}$ ; 1.28% saturation) located immediately to the north (Fig. 1). At least in this mesotrophic region of the Arabian Sea, therefore, the evidence to hand is that the active consumption of  $N_2O$  by the *Labrenzia*-like alphaproteobacteria was most probably confined to the deeper, suboxic waters.

Comparable studies of *nosZ* expression in natural populations of denitrifying bacteria in open waters are sparse. A recent investigation, however, did find *nosZ* mRNAs within the metatranscriptome of organisms obtained from suboxic depths within the OMZ off the coast of northern Chile but not in samples from shallower waters (see Table S6 in reference 48). Like the *Labrenzia*-like bacteria in the present study, organisms harboring *nosZ* were present at these shallower, oxic depths, but their metabolism was probably fueled by aerobic respiration outside the OMZ core because transcripts originating from *nirS/K* were also close to the limits of detection (48). Indeed, it is thought that many of the organisms that inhabit OMZs are metabolically versatile and that they are involved in other biologically mediated transformations of nitrogen and various other elemental cycles (49). *Labrenzia* species, for example, have been shown to be capable of CO oxidation and mixotrophic CO<sub>2</sub> fixation (50) as well as the production of dimethylsulfide from dimethylsulfoniopropionate (51).

**Depth distribution of *Labrenzia*-like denitrifying bacteria transcribing *nosZ* outside the Arabian Sea OMZ.** Nevertheless, some denitrifiers are able to reduce nitrogen oxides to  $N_2$  and/or  $NO_x$  under elevated  $O_2$  concentrations (52, 53), and it is thought that some water column denitrification must occur outside the major OMZs (54, 55). Indeed, firm evidence for aerobic denitrification in the wider marine environment is accumulating (56). Accordingly, RNA samples that were obtained from the oligotrophic stations 1 and 2 were also analyzed by QRT-PCR for the *Labrenzia*-like *nosZ* mRNA even though the water column at these equatorial latitudes was not especially oxygen depleted, i.e.,  $\geq 30\%$  saturation throughout the upper 300 m (Fig. 3 and 4). Most unexpectedly, transcripts originating from the *Labrenzia*-like *nosZ* gene were readily detected in the upper water column at both stations (Fig. 4). Transcript abundance was greatest, however, within the chlorophyll maximum located toward the base of the euphotic zone at both stations.

The highest *nosZ* mRNA concentrations ( $[0.49 \pm 0.12] \times 10^5$  and  $[0.89 \pm 0.27] \times 10^5$  copies liter<sup>-1</sup> at 74-m and 50-m depths at stations 1 and 2, respectively) were centered on the 23.5 isopycnal ( $\text{kg m}^{-3}$ ) and were also coincident with the primary nitrite maximum (PNM) at each station. The nitrite concentrations ( $>0.5 \mu\text{M}$ ) within the PNM at these stations were somewhat above the range (0.01 to 0.4  $\mu\text{M}$ ) commonly encountered elsewhere (57). The source of nitrite within the PNM is thought to derive from the incomplete assimilation of nitrate by phytoplankton and/or the activities of nitrifying bacteria (57, 58, 59). The predominance of *Prochlorococcus* spp. at these stations (41), which, in general, do not utilize nitrate, would suggest that nitrifiers, rather than phytoplankton, were the most significant nitrite producers in these waters. The latter organisms, rather than denitrifiers, are also thought to be the most likely sources of  $N_2O$  found within the oxycline associated with the PNM (18).

The occurrence of *nosZ* mRNAs in oxic waters is perplexing since the activity of the gene product, nitrous oxide reductase, is considered to be the most oxygen-sensitive step in the denitrification pathway (60). Constitutive expression of *nosZ* (and the preferential reduction of  $N_2O$  over  $NO_2^-$ ) has been shown under aerobic conditions in the denitrifier *Pseudomonas stutzeri* strain TR2 (53). In the Arabian Sea isolate 4N, by contrast, the abundance of *nosZ* mRNA is close to the limit of detection when grown in well-aerated cultures but is upregulated rapidly (within 30 min) in denitrifying medium following the removal of oxygen (Fig. 4). When placed in the field context of the low abundances of *Labrenzia*-like *nosZ* mRNAs detected in near-surface waters and below the upper mixed layer at stations 1 and 2, these experimental find-



**FIG 3** Depth distributions of chlorophyll (solid lines), potential temperature (dashed lines), oxygen saturation (dotted lines), and the concentration (mean  $\pm$  standard error,  $n = 3$ ) of the *Labrenzia*-like *nosZ* gene (bars) at four stations in the Arabian Sea. The abundance of *nosZ* at station 6 is shown at 10-fold the actual concentration, and the depth distribution of chlorophyll at station 10 is shown in the inset. The arrow shows the depth (120 m) from which the seawater was obtained for the original enrichment cultures at station 10. \*, PCR inhibitors detected.

ings are consistent with the specific induction of *nosZ* within the localized environment of the PNM.

It has been recognized for more than 25 years that particulate materials such as marine snow and fecal pellets are potential mi-

crostrites within the oxygenated water column that might provide the reduced niches required to sustain suboxic processes such as denitrification and methanogenesis (61, 62, 63). These materials, which are readily trapped on density surfaces within the pycnocline (64, 65), support an active, sessile, microbial community that includes *Roseobacter* spp. (66), with which the genus *Labrenzia* is closely affiliated and with which some species share a particle-associated lifestyle (50). *Labrenzia* spp. are also known to form close (symbiotic) associations with a range of other organisms that include large diatoms and dinoflagellates (67, 68) and other algae (69). It may not be entirely coincidental, therefore, that the greatest concentrations of *Labrenzia*-like *nosZ* transcripts that were detected at stations 1 and 2 were located within the chlorophyll maximum close to the pycnocline.

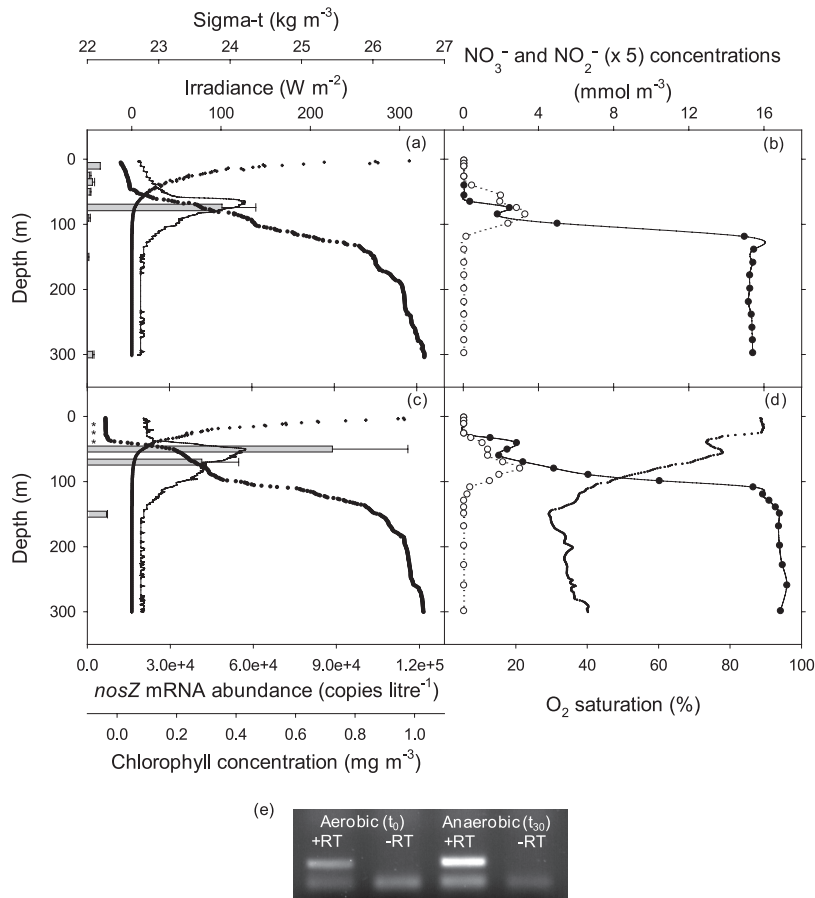
***Trichodesmium* spp. as potential hosts for *Labrenzia*-like denitrifying bacteria in oxic waters of the Arabian Sea.** The phytoplankton present at these oligotrophic stations were dominated by the small-size classes (*Prochlorococcus* spp. and, to a lesser extent, picoeukaryotic algae [41, 42, 70]). The very low sinking velocities of these organisms would militate against the production of significant numbers of potentially suboxic aggregates within the water column (71). Perhaps tellingly, however, the diazotrophic cyanobacterium *Trichodesmium* was also present at these southern stations but not in the nutrient-enriched waters to the north of station 4 (21). *Trichodesmium* spp. form macroscopic colonies that are known to provide a so-called pseudobenthic habitat for many other organisms. These include metazoa such as small copepods, diatoms, protists and other eukaryotic microbes; viruses; and also autotrophic and heterotrophic prokaryotes (72, 73, 74, 75, 76).

As part of a parallel study examining diazotroph diversity in this region, we collected *Trichodesmium* colonies at the southern stations 2, 3, and 4 and, by serendipity, were able to interrogate DNA isolated from the colonies for the presence of the *nosZ* gene from the *Labrenzia*-like Arabian Sea isolates. For this purpose, a different primer pair (StanosF and StanosR2 [Table 1]) from that used for the QPCR experiments was utilized in order to amplify a larger (252-bp), more phylogenetically informative fragment of *nosZ* from the target organisms.

PCR products of the expected size were readily obtained from DNA samples purified from the *Trichodesmium* colonies collected at all three southern stations. The amplicon obtained from station 3 was selected for further investigation and found to contain the fragment of *nosZ* targeted by the primers. Compared to *nosZ* from the isolate 4N, its DNA sequence (GenBank accession number JX827177) was found to contain a nucleotide substitution (a G/A transition) at position 204 within the 252-bp region amplified, leading to a single amino acid change (Y to C) at amino acid position 68 within the translated peptide. The peptide sequence was otherwise  $\sim$ 99% identical to those of the Arabian Sea isolate 4N and *L. aggregata* IAM 12614 and, unsurprisingly, formed part of the same tight cluster when analyzed phylogenetically (Fig. 2b).

While their wider distribution in more northerly waters (and in the deeper waters at station 1) does not suggest an obligate relationship, at least some component of the *Labrenzia*-like population sampled at the southern stations was part of the epibiotic community associated with *Trichodesmium* spp. Given the particle-associated and epibiotic lifestyle of both cultured close relatives and the range of host organisms from which so many of the phylogenetically related environmental clone sequences origi-





**FIG 4** (a to d) Depth distribution of chlorophyll (solid line), PAR (crosses), density (closed circles), and the concentration (mean  $\pm$  standard error,  $n = 3$ ) of *Labrenzia*-like *nosZ* mRNA (bars; \*, not detected) at stations 1 (a) and 2 (c). The corresponding depth distributions of nitrate (closed circles) and nitrite ( $\times 5$ , open circles) concentrations at stations 1 (b) and 2 (d) are shown in the right-hand panels. The additional dotted line shown in panel d is the oxygen saturation profile for station 2. (e) Electrophoretogram of *nosZ* RT-PCR products from the Arabian Sea isolate 4N grown in denitrification medium under an aerobic atmosphere ( $t_0$ ) and following transfer to suboxic conditions for 30 min ( $t_{30}$ ). Equal quantities of total RNA were used for the reverse transcriptase (+RT) reactions. The absence of contaminating DNA was verified in parallel reactions in which the enzyme was omitted (-RT). The mean concentration ( $n = 3$ ) of the *nosZ* PCR product at  $t_{30}$  determined by QPCR was 4.23-fold that at  $t_0$ . The lower band in each lane is a mixture of QPCR primers and the fluorescein-labeled *nosZ* probe.

nated (Fig. 2; Table 2), this finding was not unexpected. It acquires some significance, however, because oxygen-depleted microzones are known to occur within the interior of *Trichodesmium* colonies (77). Owing to their high endogenous respiratory rates, they may become virtually anoxic in subdued light and, in particular, at night (78). The low-oxygen environment that can develop within *Trichodesmium* colonies, therefore, might provide a potentially novel niche for N<sub>2</sub>O reduction by colony-associated, *Labrenzia*-like bacteria and, perhaps, also for earlier steps in the denitrification pathway from NO<sub>3</sub><sup>-</sup> and NO<sub>2</sub><sup>-</sup>.

It should not be forgotten, though, that the *Trichodesmium* colonies sampled in the present study were from well-illuminated waters where NO<sub>3</sub><sup>-</sup> and NO<sub>2</sub><sup>-</sup> concentrations were in the low-nanomolar range (Fig. 4) (42). Nevertheless, with this caveat in mind a rationale can be developed to explain why *nosZ* transcripts from the *Labrenzia*-like bacteria were found to be most abundant close to the chlorophyll maximum within dimly lit waters near the pycnocline. *Trichodesmium* colonies can undergo rapid (>10-m hour<sup>-1</sup>) vertical migrations through the water column (79) and may spend extended periods of the day in the lower reaches of the

upper mixed layer (80, 81). At these depths, they are physically closer to the sources of nitrogen oxides (NO<sub>2</sub><sup>-</sup> and N<sub>2</sub>O) produced *in situ* within the PNM (18) and also to the reservoir of NO<sub>3</sub><sup>-</sup> advected from deeper waters.

Even during the peak daylight hours, endogenous photosynthetic oxygen evolution is likely to be well below the compensation point in *Trichodesmium* colonies residing at these poorly illuminated but more nitrogen-replete depths (78). The foci of suboxia that develop within the colonies under these conditions are not only within easy reach of a ready supply of nitrogen oxides from surrounding waters but also are bathed in the locally produced fixed carbon (and nitrogen) that is exuded from within the *Trichodesmium* colony (e.g., see references 82 and 83). This beneficial combination of circumstances should provide the ideal conditions for the rapid induction of *nosZ* by epibiotic *Labrenzia*-like organisms associated with *Trichodesmium*, as was observed for the cultured isolate 4N under suboxia in the laboratory (Fig. 4).

Our observations provide a plausible mechanistic explanation for the expression of one of the signature genes (*nosZ*) of the

denitrification pathway in an oxygenated water column and have pinpointed a potential biological sink for the  $N_2O$  produced by nitrifiers (and denitrifiers?) in low-latitude near-surface waters. Much further work is required to establish how widespread such close associations between *Trichodesmium* spp. and epibiotic denitrifiers are, but during the present study a closely related (99% nucleotide identity) *nosZ* sequence was also amplified from *Trichodesmium thiebautii* colonies collected off the Bahama Islands in 1991 (84), and a further *nosZ* gene fragment was recovered from DNA purified from the cultured isolate *Trichodesmium erythraeum* strain IMS101 (Fig. 2b). We note, furthermore, that 16S rRNA sequences belonging to alphaproteobacteria that are very similar to those of *Labrenzia* spp. (e.g., clone BATS14\_155) and *Nisaea denitrificans* (e.g., clone BATS14\_163), a closely related denitrifier (Fig. 2), were among those recovered from the epibiotic community of *Trichodesmium* spp. collected from the Sargasso Sea (see Table S1 in reference 76).

Close biogeographical coupling between pelagic nitrogen fixation and denitrification in nitrogen-deficient ocean waters has been proposed before (85) but not, to date, at the microscopic spatial scales implied by the findings reported here.

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