Transcriptional response of the archaeal ammonia oxidizer

*Nitrosopumilus maritimus* to low and environmentally relevant ammonia concentrations

Running title: Transcription of AOA at low NH$_4^+$

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The ability of chemoautotrophic ammonia-oxidizing archaea to compete for ammonia among marine microorganisms at low ambient concentrations has been in part attributed to their extremely high affinity for ammonia, but as yet there is no mechanistic understanding of supporting metabolism. We examined transcription of selected genes for anabolic functions (CO₂ fixation, ammonia transport, and cell wall synthesis) and a central catabolic function (ammonia oxidation) in the thaumarchaeon *Nitrosopumilus maritimus* SCM1 growing at two ammonia concentrations, as measured by combined ammonia and ammonium, one well above the $K_m$ for ammonia oxidation (~ 500 μM) and the other well below the $K_m$ (<10 nM). Transcript levels were generally immediately and differentially repressed when cells transitioned from ammonia replete to ammonia limiting conditions. Transcript levels for ammonia oxidation, CO₂ fixation, and one of ammonia transports were approximately the same at high and low ammonia availability. Transcripts for all analyzed genes decreased with time in the complete absence of ammonia, but with varying rates of decay. The new steady state mRNA levels established are presumably more reflective of the natural physiological state of ammonia-oxidizing archaea and offer reference for interpreting message abundance patterns in the natural environment.

**INTRODUCTION**

The generally extremely low availability of ammonia in the open ocean, from below detection to low micromolar concentrations, suggests that free ammonia is a limiting resource for microbial growth (1-6). Thus, competition for ammonia among marine
microorganisms is intense. However, the generally high numbers of ammonia-oxidizing archaea (AOA) in these same environments indicates that they are not limited by ammonia (here defined as combined ammonia and ammonium) concentrations in the low nanomolar range. This inference is supported by the exceptionally low $K_m$ for ammonia oxidation of *Nitrosopumilus maritimus* ($K_m \approx 133$ nM combined $\text{NH}_4^+ + \text{NH}_3$) (7), by the direct determination of the $K_m$ for ammonia oxidation in a marine system dominated by AOA ($K_m \approx 100$ nM) (8), and environmental metatranscriptomic analyses showing significant abundance of transcripts for genes coding for the thaumarcheal ammonia monooxygenase (*amoA*), ammonia transport (*amt*), and cell wall S-layer proteins (*slp*) (9-14). Thus, the AOA may exert primary control over ammonia availability throughout marine water column, including the control of nitrification in marine oxygen minimum zones (1, 15-18).

However, there is as yet no reference for relating environmental measures of transcript abundance to the physiological state of the contributing populations. The constantly low concentrations of ammonia typical of the marine environment cannot be maintained using standard batch culture conditions. We therefore developed a simple system, based on growing *N. maritimus* in dialysis bags, as a mechanism to establish quasi-steady state growth at ammonia concentrations in the nanomolar range.

Once growth conditions were established, we examined transcript abundance patterns for genes coding key functions in ammonia acquisition and cell synthesis. Selected genes encompassed one subunit of the ammonia monooxygenase (*amoA*), two ammonia transporters (*amt1, amt2*), two S-layer proteins (*slp1, slp2*), and the 4-hydroxybutyryl-CoA dehydratase (*hcd*) in the pathway for CO$_2$ fixation (19). The transcripts for these genes in *N. maritimus* were examined under conditions of ammonia...
excess (batch culture), ammonia limitation (dialysis bag growth), extended ammonia starvation, and short-term fluctuations in ammonia availability.

**MATERIALS AND METHODS**

**Growth Conditions.** All studies were performed with *N. maritimus* SCM1 (20). The strain was maintained in 500 ml of HEPES-buffered synthetic *Crenarchaeota* medium (7) in 2-liter glass bottles at 30°C in the dark without shaking and transferred (1% inoculum) to fresh medium at late-exponential phase. The concentration of nitrite was determined colorimetrically with the Griess-Ilosvey reagent (quantitative limit, 1.0 µM) (21). The concentration of NH$_4^+$ was determined using a fluorescence microplate reader and the o-phthaldialdehyde (OPA) reagent (quantitative limit, 10 nM) (22).

**Comparative analyses of alternative growth states: batch, dialysis bag, and starvation.** Batch culture experiments (three biological replicates) were carried out in 2-liter glass bottles without shaking in the dark (Fig. S1). One ml of medium was immediately quenched on ice for determinations of transcript abundance (*n* = 2), cell numbers (*n* = 1), and NH$_4^+$ and NO$_2^-$ concentrations (as described below).

The dialysis bag experiments were conducted using a 10-liter glass bottle containing approximately 9 liters of medium, without added NH$_4$Cl, pre-incubated overnight at 30°C. (Fig. S1). This medium contains approximately ~1.6 µM ammonia determined by the OPA reagent as described above, originating from the trace levels of ammonia present in the reagent grade chemicals used to prepare the synthetic medium. Dialysis tubing (12,000 ~ 14,000 Da molecular-weight-cutoff Spectra/Pro4, 6.4 mm diameter, Spectrum Labs, Rancho Dominguez, CA) cut to approximately 15.5 cm lengths was boiled in 800
ml of sodium bicarbonate buffer (790 ml of ultra pure water, 16 g NaHCO₃, 1.6 ml of 500 mM EDTA, pH 8.0) for 10 min. After rinsing the dialysis tubes in 800 ml of ultra pure water twice, the dialysis tubes were boiled in 800 ml of sterile ultra pure water for 10 min, and then washed in sterile pre-incubated 50 ml of SCM1 medium without NH₄Cl.

The rinsed dialysis tubing was filled with 3.5-ml of batch culture SCM1 at late exponential growth (day 6), and clamped at both ends. Dialysis bag cultures were transferred into a pre-incubated 10-liter glass bottle containing approximately 9 liters of medium, incubated at 30°C, and stirred with a stir bar in the dark. As shown in Fig. S1, replicate dialysis bag cultures were removed at 1, 2, 3, and 6-days of incubation, for a total of three biological replicates. Culture material was recovered from the dialysis bag with a 3 ml syringe, immediately quenched on ice, and then aliquoted for determinations of transcript abundance (1 ml), amoA gene abundance (1 ml), and cell counts (1 ml).

Cells were harvested by centrifugation (20,000 g) for 10 min at 4°C in 1.5 ml-PCR microcentrifuge tubes (Eppendorf, Hamburg, Germany). The supernatant was removed, immediately filtered with Millex-GV (pore size, 0.22 µm, Millipore, Billerica, MA, USA), and stored at -20°C for later determination of NH₄⁺ and NO₂⁻ concentrations. One ml of RNALater (QIAGEN, Valencia, CA) was added to each cell pellet, mixed well, and then centrifuged at 4°C, 20,000 g, for 10 min. After discarding 0.95 ml of supernatant, the fixed cells were stored at -80°C until used for qRT-PCR analysis (as described below).

Dialysis bag cultures (n = 6, for biological triplicates) for starvation experiments were established from 3.5-ml of mid-exponential growth SCM1 and placed in 10-liter glass bottles containing approximately 9-liters medium without added NH₄Cl (Fig. S1). After 1-day of incubation, 7 ml of medium recovered from two of the dialysis bag cultures was transferred to a sterile 15-ml conical tube (BD, Franklin Lakes, NJ) to initiate conditions...
of NH$_4^+$ starvation. The starvation experiments were carried out in the 15-ml tubes incubated at 30˚C without shaking in the dark (Fig. S1), removing 1 ml samples at different times ($n = 2$) to monitor transcript abundance and cell numbers ($n = 1$).

RNA extraction. Total RNA was extracted using an RNeasy Mini Kit (QIAGEN) in combination with an RNase-Free DNase set (QIAGEN) according to manufacturers instructions. The extracted RNA solutions were treated with DNase (TURBO DNA-free kit, Life Technologies, Carlsbad, CA), again according to manufacturers instruction. The absence of DNA contamination was confirmed by PCR.

qRT-PCR. The mRNAs of amoA, ($\text{Nmar}_1500$), amt1 ($\text{Nmar}_0588$), amt2 ($\text{Nmar}_1698$), hcd ($\text{Nmar}_0207$), slp1 ($\text{Nmar}_1201$), and slp2 ($\text{Nmar}_1547$) genes were quantified by real-time PCR in a LightCycler system (Roche Diagnostics, Mannheim, Germany) with LightCycler RNA Master SYBR Green I kit (Roche Diagnostics). Each reaction of the one-step reverse transcription-PCR (RT-PCR) was conducted in a 10-µl volume containing 0.5 µl of template RNA solution, 0.5 µM of each primer, and 3.75 µl of LightCycler RNA Master SYBR Green I, and 3.25 mM Mn(OAc)$_2$. The PCR primer sets in Table S1 and PCR cycling in Table S2 were used for quantitative RT-PCR. The standard curves for each mRNA were generated using an RNA standard in a ten-fold dilution series of $10^2$-$10^9$ copies per reaction. The amplification efficiencies of quantitative RT-PCRs averaged 70.8% ($r^2 = 0.999$), 80.3% ($r^2 = 1.000$), 72.2% ($r^2 = 0.999$), 76.8% ($r^2 = 0.999$), 74.1% ($r^2 = 0.999$), and 81.0% ($r^2 = 0.999$) for amoA, amt1, amt2, hcd, slp1, and slp2 transcripts, respectively.
Cell counts. Cells were fixed in 2% paraformaldehyde for at least 1 h. The fixed cells were diluted in phosphate-buffered saline to adjust the density from 10 to 100 cells per field for the cell counting, and filtered on 0.2-µm-pore-size polycarbonate GTBP membrane filters (Millipore, Billerica, MA, USA), and then stained with 2 µl of Abnova 4’,6’-diamidino-2-phenylindole (DAPI) (1.5 mg ml\(^{-1}\), Abnova, Taipei, Taiwan). The cell numbers were determined for each sample by epifluorescence microscopy, counting cells in each of 20 random fields.

Response to short-term ammonia starvation and ammonia re-addition. Rinsed dialysis tubing was filled with 3.5-ml batch culture SCM1 at late exponential growth (day 6), clamped at both ends, and used for studies of growth under conditions of low ammonia availability and in response to short term changes in ammonia availability. Conditions of ammonia limitation were established by transferring the dialysis bag cultures to 10-liter glass bottles containing 9-liter synthetic medium at 1.6 µM NH\(_4^+\). This concentration of ammonium originates from the trace levels present in the reagent grade chemicals used to prepare the synthetic medium. After incubation for 8 h in low ammonia medium, replicate dialysis bag cultures were transferred to a 10-liter bottle containing approximately 9 liters of medium without added NH\(_4\)Cl (Fig. S2). Ammonia starvation was initiated at specified times by transferring individual replicates to a sterile petri dish without media in a plastic bag containing a wetted filter strip and incubating at 30°C. Dialysis bags were transferred between bottles containing media without ammonia and sterile petri dishes without media for analyses of transcriptional response to short term fluctuations in ammonia availability (Fig. S2). Culture material was removed
using a 3 ml syringe and immediately quenched on ice prior to analysis of transcript abundance, and NH$_4^+$ and NO$_2^-$ concentrations.

RESULTS AND DISCUSSION

*N. maritimus* retained within dialysis bags suspended in a large volume of medium poised at ~2 μM total ammonia grew at near maximum growth rates while maintaining the concentration of ammonia at or below 10 nM, the analytical limit of detection (Fig. 1). Since cells showed no tendency to attach to the dialysis bag walls under these growth conditions, the cells on average were experiencing ammonia concentrations in the very low nanomolar range. At these low ammonium concentrations, oxygen was not a limiting resource. Ammonia in batch cultures was depleted below the detection limit of 10 nM after *N. maritimus* entered stationary phase as described previously (7). The concentration of ammonia in the dialysis bags decreased following transfer to the ammonia limited medium (Figs. 1A and 1B) and was maintained below 10 nM by active consumption by *N. maritimus* throughout the experiments. In contrast, the NO$_2^-$ concentration within the dialysis bags was slighter higher than the bulk medium, reflecting a production rate exceeding the rate of equilibration between compartments. *N. maritimus* cell numbers within the dialysis bags increased throughout the experimental period, confirming their ability for continuous growth at low nanomolar concentrations of ammonia (Fig. 1C and Table 1). The abundances of *amoA* genes determined by quantitative PCR were smaller than those obtained by direct cell count (Table 1). The inefficiencies of cell harvesting and extraction likely caused this discrepancy. However, since this loss is relatively low and is expected to be comparable for DNA and RNA, this does not alter our conclusions.
Prior to transfer to the dialysis bags, the highest per cell transcript values for those genes evaluated were for those coding for ammonia oxidation (amoA), ammonia uptake (amt2), and one of two closely related S-layer proteins (slp2) (Table 2). This is consistent with published metatranscriptomic analyses of natural marine systems, finding higher-level transcripts for amoA, amt2 and slp2. Thus, transcripts of these genes, coding for central energy generating and anabolic functions are expected to provide useful signatures for active cell growth in environmental samples (9-14). The per-cell values for these transcripts varied between a high value for slp2 (~ 60 transcripts/cell) and a low value for hcd (~ 0.08 transcripts/cell). Since the amoA gene counts were comparable to cell counts (Table 1), the low abundances of the per-cell transcripts except for slp2 likely reflect a physiological state in which some cells in a population are not actively transcribing those genes. The per cell abundance of all transcripts decreased during dialysis bag incubation, approaching new steady-state values by day 12.

Relatively small changes in transcript abundance for amoA, amt2, hcd, and slp2 genes were observed following transition to growth at the low ammonia concentration maintained in the dialysis bags (Fig. 2). Reduced per cell abundances were about 54% (hcd), 28% (amt2), 21% (amoA), and 10% (slp2) of levels determined prior to transfer to the dialysis bags. These data indicate that the cellular investment in central systems for energy generation and anabolism do not vary greatly over a wide range of ammonia concentrations, in this study varying from below 10 nM to 500 μM.

In contrast to those genes for which transcript numbers remained relatively high in a low ammonia environment, transition to nanomolar ammonia growth conditions resulted in a much greater reduction in transcripts for an alternative S-layer gene (slp1) and ammonia transporter (amt1), reduced to 3% and 0.4% of initial abundances, respectively.
Such a rapid decrease is suggestive of active message decay. Also, since these genes code for variants of the ammonia transporter and S-layer proteins, they may function primarily during growth at higher ammonia concentrations, the Amt2 possibly functioning as a lower affinity transporter and the Slp2 supporting cell wall synthesis associated with higher growth rates. It is also possible that varying stoichiometry of S-layer proteins with varying ammonia availability may be of functional significance. For example, recent structural analyses of the S-layer protein of the methanogenic archaeon *Methanosarcina acetivorans* revealed a highly negatively charged pore surface, suggesting a possible role of this S-layer in selective ion uptake (23). Therefore, it is essential to investigate the structure of S-layer of the *N. maritimus* in order to evaluate if it serves a similar function in the collection of ammonia, or ammonium, at extremely low concentrations.

The dual roles of ammonia – assimilated for cell synthesis or oxidized as a source of energy and reductant - also presents a possible conflict for this organism. Ammonia/ammonium collected at the cell surface must be partitioned between anabolic and catabolic pathways, necessitating a balance in the affinities of enzyme systems controlling the flux of ammonia through these alternative pathways. Studies of ammonia assimilation by *Escherichia coli* have shown that active ammonia transport by its AmtB ammonia permease is essential for growth below ~20 μM NH₄⁺ (24). At higher ammonia concentrations diffusion of ammonia through the membrane is a major source of ammonia for biosynthesis. In *N. maritimus*, transcripts for two ammonia permease genes are abundant at the higher ammonia concentration, with transcripts for one (*amt2*) present in about 10 fold higher abundance than the second (*amt1*). The rapid depletion of *amt1* transcripts with transition from high to low ammonia concentrations...
(Fig. 2B) suggests this permease has a higher $K_m$ (lower affinity) than that of the second

(amt2). Thus, the Amt2 permease presumably supports ammonia assimilation at
concentrations in the very low nanomolar range.

All transcripts decreased following short-term elimination of ammonia, albeit having
significantly different rates of decay, with transcripts for amoA and amtI being rapidly
depleted in the absence of ammonia (Fig. 3). However, cells were also immediately
responsive to the resupply of nanomolar amounts of ammonia, and within 1-2 hours
returned to near pre-starvation transcript levels. Thus, these results point to an
extremely rapid adaptive response to changing ammonia availability, and further enforce
the importance of rapid preservation of cells harvested for metatranscriptomic analyses
(10).

A similar, rapid decay of amoA transcripts in *Nitrosomonas europaea* was observed
under short-term ammonium-starvation (25). In contrast, amoC transcript numbers
remained elevated under these conditions (26). The response of ammonia-starved *N.
europaena* and *Nitrospira briensis* to ammonia addition was also similar to *N. maritimus*,
as evidenced by a rapid increase in amoA transcripts following readdition of 15 or 50 mM
ammonia to *N. europaea* and 5 mM ammonia to *N. briensis* (27, 28). Thus, the AOA and
ammonia-oxidizing bacteria (AOB) may share similar regulatory strategies to cope with
changing ammonia concentrations, although differing vastly in the range of ammonia
concentrations to which they are responsive. The $K_m$ of *N. maritimus* for ammonia
oxidation (133 nM) is much lower than that of AOB, *N. europaea* ($K_m = 553$ µM) (7),
*Nitrosoococcus oceanii* ($K_m = 101$ µM) (7), and *N. briensis* ($K_m = 3$ µM) (28).

Although additional studies are essential to establish differences in the ammonia
affinity of the two permeases, and to evaluate the suggested variation in S-layer protein
stoichiometry associated with changing ammonia concentrations, our results provide important context for interpreting transcript abundance patterns observed in analyses of natural marine populations. More generally, the strategy used in this study for growing marine oligotrophs under nutrient limitation may assist in the study of other microorganisms living at the extremes of energy limitation common to much of the biosphere (29, 30).

ACKNOWLEDGMENTS

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REFERENCES


TABLE 1 Changes in cell numbers, growth rates, and amoA gene abundances of *N. maritimus* SCM1 during batch culture and dialysis bag growth, and in response to ammonia starvation.

<table>
<thead>
<tr>
<th>Day</th>
<th>Cell number (cells / mL)</th>
<th>Growth rate (cell / mL / day)</th>
<th>amoA gene (copies / mL)</th>
<th>amoA gene in a cell (copies / cell)</th>
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<td>0</td>
<td>8.2 × 10^5</td>
<td>8.9 × 10^5</td>
<td>4.9 × 10^5</td>
<td>0.6</td>
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<td>2</td>
<td>2.6 × 10^6</td>
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<td>9.6 × 10^7</td>
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<td>7</td>
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Cell number data are presented as means ± standard deviation. Growth rates were calculated as the slope of the linear regression of cell number data. amoA gene abundances are presented as means ± standard deviation.
### TABLE 2  Changes in the per-cell transcript and gene abundance of *N. maritimus* SCM1 during batch culture and dialysis bag growth, and in response to ammonia starvation.

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<th>7</th>
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</table>
Figure legends

FIG 1 Changes in concentration of (A) NH$_4^+$ and (B) NO$_2^-$, and (C) cell numbers of *N. maritimus* SCM1 during batch culture (circles), dialysis bag growth (open squares, inner dialysis bag; closed squares, outer dialysis bag), and in response to ammonia starvation (open triangles, inner dialysis bag and the batch culture under ammonia starvation; closed triangles, outer dialysis bag for ammonia starvation). The gray zone shows ammonia detection limit (~10 nM). Black arrows indicate times when cells were transferred to dialysis bags. Open black arrows show times when cells were removed from dialysis bags for starvation experiments. Bars are standard errors ($n = 3$).

FIG 2 Changes in transcript abundance (*amoA*, *amt1*, *amt2*, *hcd*, *slp1*, and *slp2*) associated with changing ammonia availability of *N. maritimus* SCM1 during batch culture (circles), dialysis bag growth (squares), and in response to ammonia starvation (triangles). Black arrows show times of transfer from batch culture to dialysis bags. Open black arrows show times of transfer of cells from dialysis bags for starvation analyses. Bars are standard errors ($n = 3$).

FIG 3 Changes in *N. maritimus* transcript abundance associated with short-term ammonia starvation (*amoA*, *amt1*, *amt2*, *hcd*, *slp1*, and *slp2*). Short-term ammonia starvation followed by readdition (circles). Continuous dialysis bag growth (squares). Gray bars correspond to periods of ammonia-starvation.