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Enhancement of Nitrous Oxide Emissions in Soil Microbial Consortia via Copper Competition between Proteobacterial Methanotrophs and Denitrifiers

Jin Chang,[a,b] Daehyun D. Kim,[a] Jeremy D. Semrau,[b] Ju Yong Lee,[a] Hokwan Heo,[a] Wenyu Gu,[b]* Sukhwan Yoon[a]

[a]Department of Civil and Environmental Engineering, Korea Advanced Institute of Science and Technology, Daejeon, South Korea
[b]Department of Civil and Environmental Engineering, University of Michigan, Ann Arbor, Michigan, USA

ABSTRACT Unique means of copper scavenging have been identified in proteobacterial methanotrophs, particularly the use of methanobactin, a novel ribosomally synthesized, post-translationally modified polypeptide that binds copper with very high affinity. The possibility that copper sequestration strategies of methanotrophs may interfere with copper uptake of denitrifiers *in situ* and thereby enhance N_2O emissions was examined using a suite of laboratory experiments performed with rice paddy microbial consortia. Addition of purified methanobactin from *Methylosinus trichosporium* OB3b to denitrifying rice paddy soil microbial consortia resulted in substantially increased N_2O production, with more pronounced responses observed for soils with lower copper content. The N_2O emission-enhancing effect of the soil's native *mbnA*-expressing *Methylocystaceae* methanotrophs on the native denitrifiers was then experimentally verified with a *Methylocystaceae*-dominant chemostat culture prepared from a rice paddy microbial consortium as the inoculum. Finally, with microcosms amended with various cell numbers of methanobactin-producing *Methylosinus trichosporium* OB3b before CH_4 enrichment, microbiomes with different ratios of methanobactin-producing *Methylocystaceae* to gammaproteobacterial methanotrophs incapable of methanobactin production were simulated. Significant enhancement of N_2O production from denitrification was evident in both *Methylocystaceae*-dominant and *Methylococcaceae*-dominant enrichments, albeit to a greater extent in the former, signifying the comparative potency of methanobactin-mediated copper sequestration, while implying the presence of alternative copper abstraction mechanisms for *Methylococcaceae*. These observations support that copper-mediated methanotrophic enhancement of N_2O production from denitrification is plausible where methanotrophs and denitrifiers cohabit.

IMPORTANCE Proteobacterial methanotrophs—groups of microorganisms that utilize methane as a source of energy and carbon—have been known to employ unique mechanisms to scavenge copper, namely, utilization of methanobactin, a polypeptide that binds copper with high affinity and specificity. Previously the possibility that copper sequestration by methanotrophs may lead to alteration of cuproenzyme-mediated reactions in denitrifiers and consequently increase emission of potent greenhouse gas N_2O has been suggested in axenic and coculture experiments. Here, a suite of experiments with rice paddy soil slurry cultures with complex microbial compositions were performed to corroborate that such copper-mediated interplay may actually take place in environments cohabited by diverse methanotrophs and denitrifiers. As spatial and temporal heterogeneity allows for spatial coexistence of methanotrophy (aerobic) and denitrification (anaerobic) in soils, the results from this study suggest that this previously unidentified mechanism of N_2O production may account for a significant proportion of N_2O efflux from agricultural soils.

KEYWORDS copper, denitrification, methanobactin, methanotrophs, nitrous oxide

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Address correspondence to Sukhwan Yoon, syoon80@kaist.ac.kr.

* Present address: Wenyu Gu, Department of Civil & Environmental Engineering, Stanford University, Palo Alto, California, USA.

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Methane (CH_4) and nitrous oxide (N_2O) are the most influential greenhouse gases apart from CO_2, with estimated contributions of 16% and 6.2%, respectively, to global greenhouse gas emissions over a 100-year time frame (1). Biological sources and sinks significantly impact both atmospheric CH_4 and N_2O pools. That is, methanotrophs consume much (up to 100%) of CH_4 originating from methanogenesis, balancing the global CH_4 budget (2). Similarly, N_2O produced via microbially mediated nitrification and denitrification is offset by N_2O reduction mediated by microbes capable of expressing and utilizing nitrous oxide reductases (NosZ) (3–5). Collectively, the relative abundance and activity of different microbial groups are critical in determining if any particular environment is a net source or sink of these potent greenhouse gases.

Methanotrophy has become a rather comprehensive term following recent discovery of extremophilic verrucomicrobial methanotrophs, nitrite-reducing anaerobic NC-10 methanotrophs, and archaeal anaerobic methanotrophs (6–9); however, except for certain specialized extreme habitats, "conventional" aerobic proteobacterial methanotrophs often dominate methane oxidation *in situ*, especially in terrestrial systems (e.g., oxic-anoxic interfaces of rice paddy soils and landfill cover soils), as observed in recent metagenomic analyses (10, 11). Proteobacterial methanotrophs are phylogenetically subdivided into gammaproteobacterial and alphaproteobacterial subgroups (12). Both of these organismal groups utilize particulate and/or soluble methane monooxygenases (MMOs) encoded by *pmo* and *mmo* operons, respectively, and these MMOs are currently known as the only enzymes capable of preferentially catalyzing the oxidation of CH_4 to CH_3OH (13). Of the two forms of MMOs, the particulate methane monooxygenase (pMMO) has been regarded as the prevalent form in most terrestrial environments, and the majority of proteobacterial methanotroph genomes sequenced to date contain only the *pmo* operon(s) (14). The expression and activity of pMMO are strongly dependent on copper, although the exact role of copper in pMMO remains unanswered (12, 15, 16).

Due to the dependency of pMMO activity on copper, pMMO-expressing methanotrophs have high demands for copper (17). Not surprisingly, methanotrophs have developed effective copper uptake mechanisms, presumably to cope with limited copper availability *in situ*. Some alphaproteobacterial methanotrophs of the *Methylocystaceae* family produce and excrete methanobactin, a modified peptide ~800 to 1,300 Da in size that chelates copper with the highest affinity among known metal chelators (empirical copper-binding constants range between 10^{18} and 10^{58} M^{-1}, depending on the experimental protocol used) (18). Copper-bound methanobactin is transported into the cell via a TonB-dependent transporter and presumably utilized for synthesis of a functional pMMO complex (19, 20). The genes encoding the methanobactin polypeptide precursor (*mbnA*) and the enzymes involved in its post-translational modifications have been identified in many alphaproteobacterial methanotroph genomes (roughly half of >40 genomes currently available in the NCBI database), and methanobactins isolated from seven distinct strains of *Methylocystaceae* methanotrophs have been chemically characterized, suggesting that the capability to synthesize and utilize methanobactin is a widespread, but not universal, trait for *Methylocystaceae* methanotrophs (18, 21–24).

In CH_4-rich, copper-depleted environments, these copper acquisition mechanisms of methanotrophs may interfere with other biogeochemical reactions catalyzed by cuproenzymes. Several key enzymes involved in the biological nitrogen cycle, including bacterial and archaeal ammonia monooxygenases (AMOs), copper-dependent NO_2^- reductases (NirK), and N_2O reductases (NosZ), require copper ions for their activity (25–27). The impact on NosZ expression and activity would be particularly consequential from an environmental perspective, as NosZ-catalyzed N_2O reduction is the only identified biological or chemical N_2O sink in the environment, and no copper-independent alternative pathway for N_2O reduction to N_2 has been identified to date (3, 28). In fact, methanobactin-mediated inhibition of NosZ activity was recently experimentally verified *in vitro* using simple, well-defined cocultures of *M. trichosporium* OB3b and several denitrifier strains possessing *nosZ*, suggesting the possibility of

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increased N_2O emissions *in situ* where methanotrophs and denitrifiers coexist (29). Oxic-anoxic interfaces and the vadose zones with fluctuating water content would provide settings in organic- and nitrogen-rich soils, where co-occurrence of obligately aerobic methanotrophy and obligately anaerobic denitrification is possible (30, 31). Whether such an N_2O production mechanism is truly relevant in the field, however, is not yet known, as neither methanobactin production nor its influence on denitrifiers has been observed in complex microbiomes such as agriculture soils.

As a follow-up to our previous study documenting the impact of methanobactin-producing methanotroph on denitrification and N_2O production in simple cocultures, the current study investigated further the potential ecological relevance of this methanotroph-denitrifier interaction by introducing microbial complexity and competition into the picture. The susceptibility of soil's complex denitrifying consortia to methanobactin-mediated alteration was examined with (i) denitrifying enrichments amended with exogenous addition of methanobactin and (ii) stimulation of native alphaproteobacterial methanotrophs of the *Methylocystaceae* family. Furthermore, the consequence of competition of alphaproteobacterial methanotrophs versus gammaproteobacterial methanotrophs on denitrification and associated N_2O production was examined with soil slurry microcosms augmented with various amounts of *M. trichosporium* OB3b before CH_4 enrichment.

RESULTS

The effect of methanobactin from *M. trichosporium* OB3b on N_2O production in denitrifying soil enrichments. The effect of the methanobactin isolated from *M. trichosporium* OB3b (MB-OB3b) on N_2O production was examined with NO_3^--reducing enrichments of five rice paddy soils with various physicochemical properties (Fig. 1; see Table S1 in the supplemental material). Without added MB-OB3b, the amount of accumulated N_2O-N did not exceed 0.2% of ~250 μmol NO_3^- initially added to the reaction vessels at any time in any of the enrichment cultures, despite four out of five soil samples being slightly acidic at $5.8 < pH < 6.3$ (Fig. 1A to E). In contrast, substantial N_2O accumulation was observed in soil slurry cultures amended with 10 μM MB-OB3b, with the exception of the soil with the highest copper content (Fig. 1F to J). Persistent N_2O accumulation was observed in the enrichments prepared with the soils with the lowest copper content (soil A, 1.14 $\pm$ 0.06 mg Cu/kg dry weight; soil B, 1.50 $\pm$ 0.02 mg Cu/kg dry weight), with 138 $\pm$ 2 and 102 $\pm$ 11 μmol N_2O-N produced from denitrification of 257 $\pm$ 29 and 275 $\pm$ 5 μmol NO_3^-, respectively. Although the non-stoichiometric N_2O production suggested partial N_2O reduction activity, no further N_2O reduction was observed for at least 24 h after completion of denitrification in either enrichment. Inhibition of NO_2^- reduction was also evident in the soil B enrichment (Fig. 1G). Persistent N_2O accumulation and delayed NO_2^- reduction were observed also with the enrichment prepared with soil C, which had a higher copper content (2.78 $\pm$ 0.28 mg Cu/kg dry weight) (Fig. 1H). The enrichments prepared with soils with the highest copper contents (soil D, 5.73 $\pm$ 0.38 mg Cu/kg dry weight; soil E, 8.05 $\pm$ 0.34 mg Cu/kg dry weight) did not persistently accumulate N_2O, even with 10 μM methanobactin added; however, the maximum amount of transiently accumulated N_2O was significantly higher ($P < 0.05$) with methanobactin than without. No significant NO_3^- or NO_2^- reduction was observed in any of the sterilized controls, and N_2O production over 120 h of abiotic incubation yielded <1.0 μmol N_2O-N, precluding the possibility that abiotic N_2O production contributed significantly to N_2O produced in the NO_3^--reducing soil enrichments (data not shown). These results demonstrated that methanobactin can inhibit N_2O and/or NO_2^- reduction in complex microbial consortia, but the effect may vary, depending on soil properties.

Denitrification and N_2O accumulation in a soil microbial consortium enriched with indigenous *Methylocystaceae* methanotrophs. To observe whether indigenous soil methanotrophs are capable of altering soil denitrification and enhancing N_2O production with methanobactin as the mediator, denitrification was observed with a *Methylocystaceae*-dominant quasi-steady-state culture extracted from a chemostat

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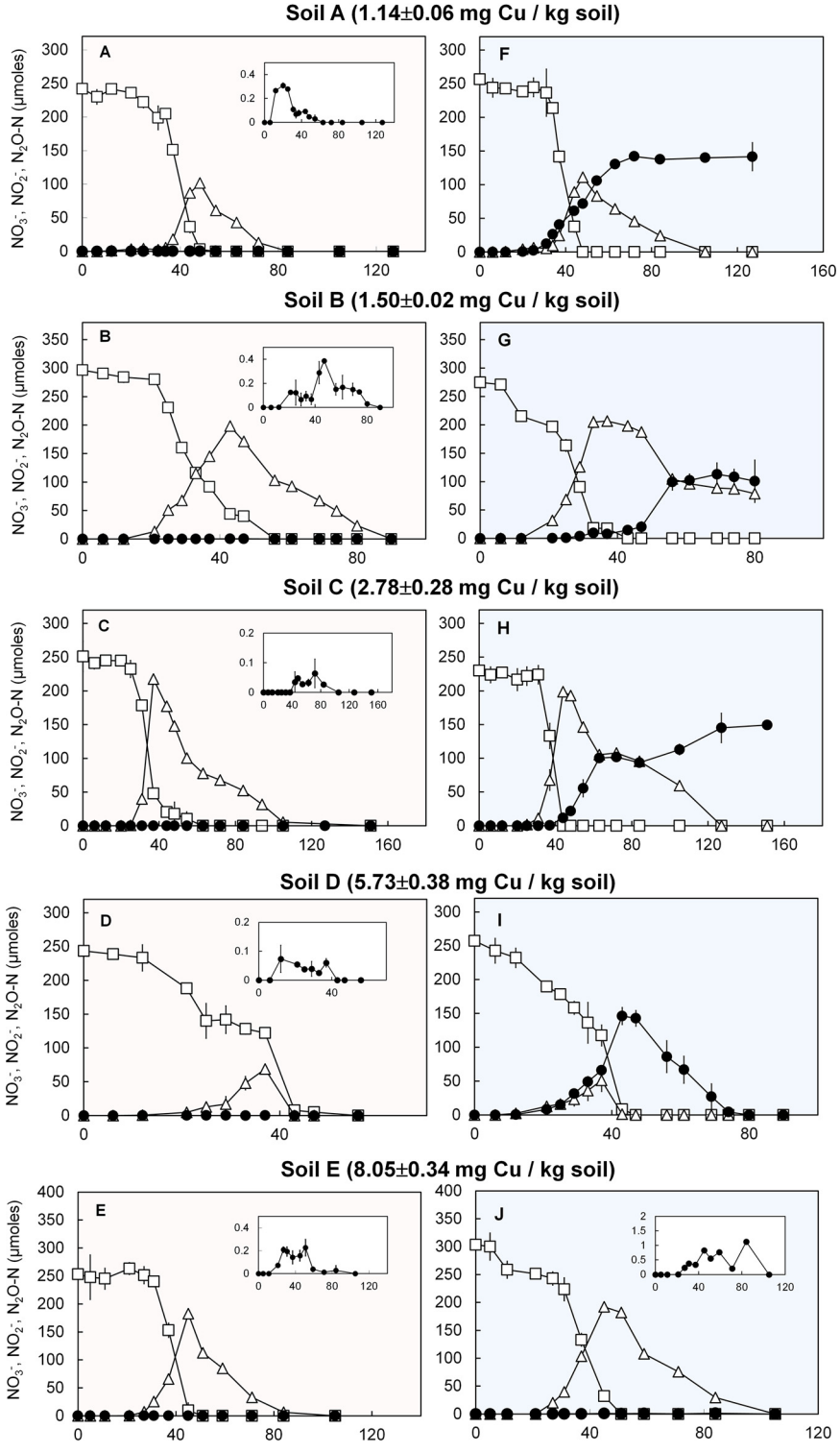

FIG 1 Denitrification of 250 μmol (5 mM) NO_3^- added to 50-ml rice paddy soil suspensions in 160-ml serum bottles amended without (A to E; shaded light red) and with (F to J; shaded blue) 10 μM OB3b-MB. The copper contents of the rice paddy soils used for preparation of the suspensions were 1.14 (A and F), 1.50 (B and G), 2.78 (C and H), 5.73 (D and I), and 8.05 (E and J) mg Cu/kg dry weight. The time series of the average amounts (μmol per vessel) of NO_3^- (□), NO_2^- (△), and N_2O (●) are presented with the error bars representing the standard deviations from triplicate samples. The inserts are magnifications of the N_2O monitoring data.

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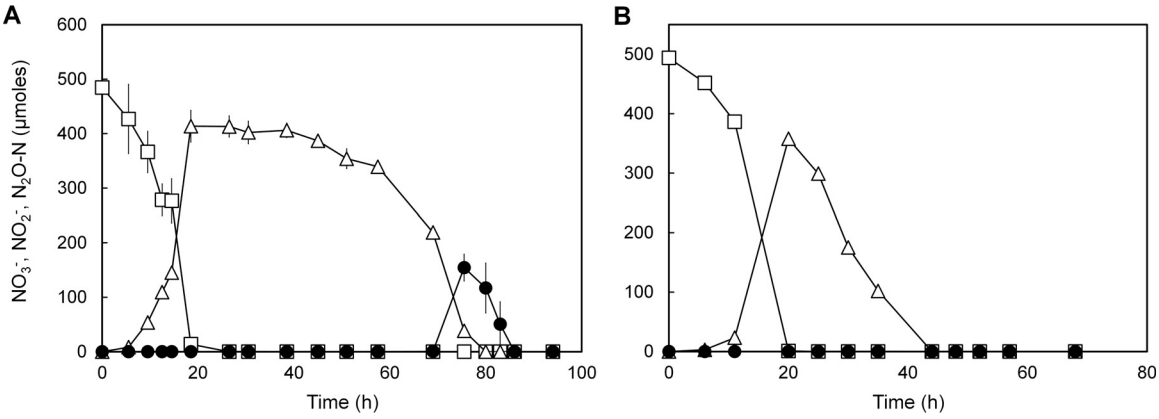

FIG 2 (A) Progression of denitrification in the 50-ml enrichment cultures (in 160-ml serum bottles with N_2 headspace) extracted from the *Methylocystaceae*-dominant chemostat. One milliliter of separately prepared denitrifier enrichment culture was added to the enrichment cultures at $t = 0$. To a set of controls (B), $CuCl_2$ was added to a concentration of 2 μM before incubation. The changes to the amounts of NO_3^- (□), NO_2^- (△), and N_2O (●) in the culture vessels were monitored. The error bars represent the standard deviations from triplicate samples.

(Fig. 2; see Table S2 in the supplemental material). The copy number of eubacterial 16S rRNA genes in the quasi-steady-state reactor culture was (1.52 ± 0.02) × 10^6 copies ml^{-1}. Alphaproteobacterial methanotrophs of the *Methylocystaceae* family were the dominant bacterial population, with 78% of the 16S rRNA reads assigned to operational taxonomic units (OTUs) affiliated with this taxon, while the OTUs assigned to the gammaproteobacterial methanotrophs (the *Methylococcaceae* family) constituted a minority group, with 1.5% relative abundance. Nonmethanotrophic taxa identified in the reactor culture included *Chitinophagaceae* (7.3%), *Pseudomonadaceae* (1.4%), and *Mycobacteriaceae* (1.3%). The most abundant *nirK* and *nosZ* genes recovered from the metagenome of the chemostat culture were most closely affiliated in terms of translated amino acid sequences to *Methylocystis* sp. strain Rockwell (69% of recovered *nirK* genes) and *Methylocystis* sp. strain SC2 (93% of recovered *nosZ* genes), both of which are the only *nirK* and *nosZ* genes found in sequenced alphaproteobacterial methanotroph genomes (see Table S3 in the supplemental material). The other *nirK* genes recovered with >1% relative abundance (among the recovered *nirK* sequences) included those most closely affiliated with the genera *Mesorhizobium* (16%), *Bauldia* (3.5%), *Panacibacter* (3.1%), *Pseudomonas* (1.8%), and *Hypomicrobium* (1.4%). The most abundant non-methanotrophic *nosZ* genes included those affiliated with the genera *Flavobacterium* (3.4%; clade II) and *Pseudomonadas* (2.5%; clade I). The coverage of *nirS* genes was substantially lower than that of *nirK* genes (2.1 × 10^{-3} *nirS*/*recA* compared to 0.133 total *nirK*/*recA* and 4.2 × 10^{-2} non-*Methylocystis nirK*/*recA*). The recovered *nirS* genes included those affiliated with the genera *Bradyrhizobium* (29.0% of recovered *nirS* genes), *Zoogloea* (41.0%), and *Acidovorax* (31.9%).

The sole unique *mbnA* sequence assembled from the shotgun metagenome reads exhibited high similarity to *mbnA* sequences of organisms affiliated with the *Methylocystis* genus (see Fig. S1 in the supplemental material). The quantitative PCR (qPCR) quantification targeting the *Methylocystaceae mbnA* genes estimated 8.1 ± 1.8 × 10^4 copies ml^{-1}, which translated to an *mbnA*/16S rRNA ratio of 0.054. Furthermore, the *mbnA* transcript/gene ratio was 19.5 ± 4.9, as determined using reverse transcription-qPCR (RT-qPCR), indicating that the *mbnA* gene was actively transcribed during quasi-steady-state operation of the reactor.

Reduction of NO_3^- was observed immediately after addition of the denitrifying inoculum to the degassed CH_4-enriched culture extracted from the chemostat and was unaffected by Cu^{2+} amendment (Fig. 2B). Repression of NO_2^- reduction was apparent in the culture without Cu^{2+} amendment, as no significant decrease in NO_2^- concentration

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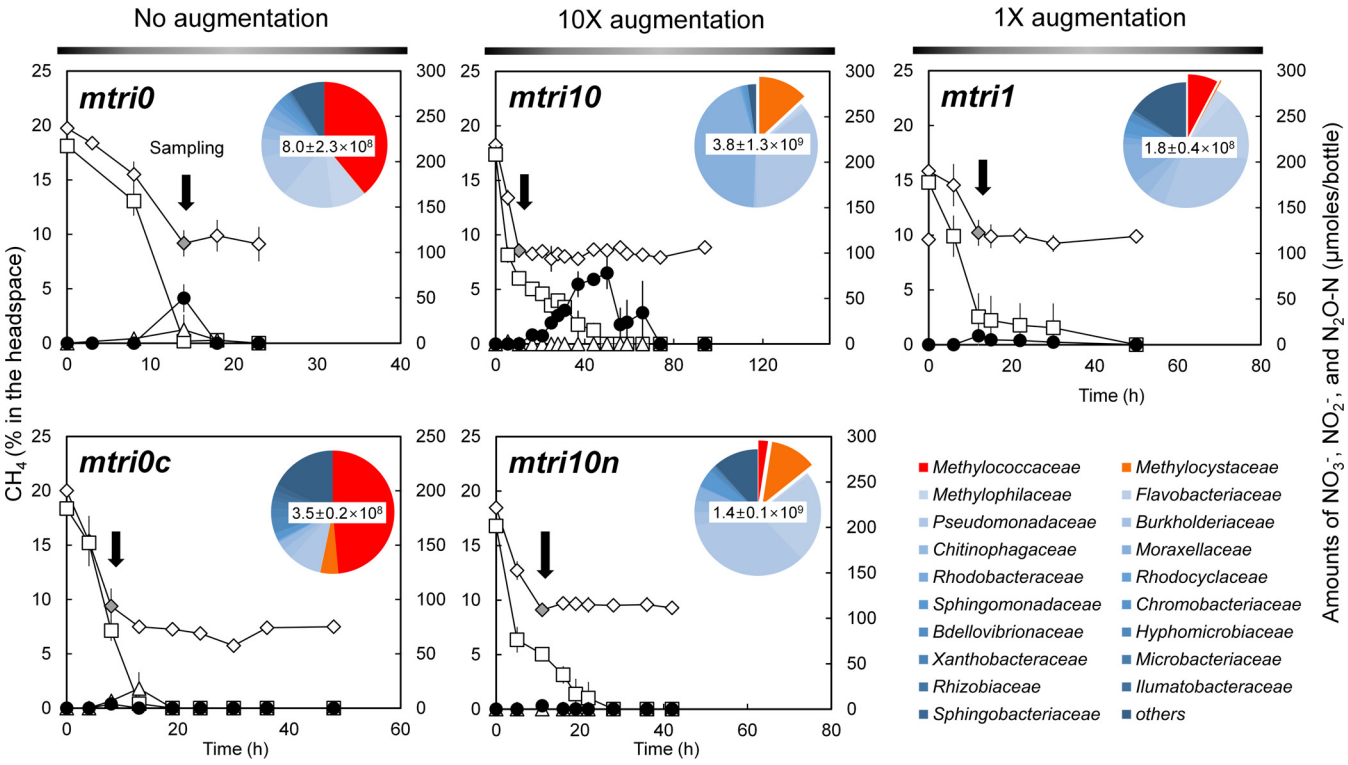

FIG 3 Monitoring of CH_4 oxidation and subsequent denitrificationand N_2O production in rice paddy soil suspensions augmented with *M. trichosporium* OB3b (or Δ*mbnAN* mutant) cells prior to incubation to quantities with their *pmoA* copy numbers matching 0 (mtri0 and mtri0c), 1 (mtri1), and 10 (mtri10 and mtri10n) times the estimated *Methylococcaceae* 16S rRNA gene copy numbers in the suspensions. The Cu-replete control, to which 2 μM $CuCl_2$ was added (mtri1c), and the control augmented with Δ*mbnAN* mutant of *M. trichosporium* (mtri10n) were included. The time series of the amounts (μmol per vessel) of NO_3^- (□), NO_2^- (△), and N_2O (●) and the headspace concentrations (%) of CH_4 (◇) are presented with the error bars representing the standard deviations of triplicate samples. The pie charts embedded in the panels illustrate the microbial community compositions of the enrichment cultures at the time points indicated by the black arrows. The numbers in the center of the pie charts are the estimates of bacterial 16S rRNA copy numbers per ml of culture at the indicated time points. The detailed microbial community compositions are presented in Table S4.

was observed for ~20 h after maximum NO_2^- accumulation was observed (414 ± 29 μmol NO_2 at t = 18.5 h). Eventual reduction of NO_2^- led to transient accumulation of N_2O, and the maximum amount of N_2O observed before it was presumably reduced to N_2 was 155 ± 25 μmol N_2O-N (~32% of added NO_3^--N). Such apparent partial repression of NO_2^- and N_2O reduction was absent in the samples amended with 2 μM Cu^{2+}. NO_2^- reduction progressed without any apparent lag, and the amount of N_2O in the vessel did not increase higher than 0.09 ± 0.01 μmol N_2O-N before it was consumed. These observations suggest that alteration of copper availability by methanobactin-producing *Methylocystaceae* methanotrophs had a substantial effect on denitrification and N_2O reduction in the broader microbial community.

Effects of altered community compositions of methanotrophic enrichments on denitrification and N_2O production upon transition to anoxia. Methanotroph-enriched microbiomes with different ratios of methanobactin-producing *Methylocystaceae* to gammaproteobacterial methanotrophs were mimicked in soil slurry microcosms by adding various amounts of *M. trichosporium* OB3b precultures to rice paddy soil slurries before batch enrichment with 20% (vol/vol) CH_4 (Fig. 3). The 16S rRNA gene copy number of *Methylococcaceae* methanotrophs in the soil sample was estimated to be (6.5 ± 1.2) × 10^6 copies g wet weight $soil^{-1}$ from their relative abundance (0.41%) and the total bacterial 16S rRNA copy number [(1.6 ± 0.3) × 10^9 bacterial 16S rRNA copies g wet weight $soil^{-1}$]. *Methylosinus trichosporium* OB3b *pmoA* copy numbers in the CH_4-enriched slurries were 0, (1.2 ± 0.2) × 10^6, and (3.3 ± 0.7) × 10^8 per ml in the cultures that had been augmented with *M. trichosporium* OB3b *pmoA* copy numbers in quantities matching 0, 1, and 10 times the estimated 16S rRNA gene copy numbers affiliated with *Methylococcaceae* methanotrophs (referred to as mtri0, mtri1, and mtri10, respectively). The 16S rRNA

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amplicon sequencing of the same samples estimated the relative abundances of *Methylocystaceae* methanotrophs to be 0.15, 0.35 and 13.0%, respectively (see Table S4 in the supplemental material). The relative abundances of *Methylococcaceae* methanotrophs were 39, 7.7, and 0%, respectively, in the same samples.

In the mtri0 culture, the maximum amount of accumulated N_2O was 49.6 $\pm$ 14.5 μmol N_2O-N and the duration of N_2O accumulation was shorter than 10 h (Fig. 3). The maximum observed N_2O accumulation (78.2 $\pm$ 16.4 μmol N_2O-N at $t = 50$ h) was significantly higher ($P < 0.05$) and the duration of N_2O accumulation substantially longer (68.5 h) in the mtri10 cultures, in which *Methylocystaceae* methanotrophs outnumbered *Methylococcaceae* methanotrophs. The decrease in the amount of NO_3^- observed during active CH_4 oxidation ($t < 10.5$ h) was presumably due to assimilation, as denitrification was unlikely to occur before O_2 depletion. Thus, N_2O accumulation resulting from dissimilatory reduction of NO_3^- between 10.5 and 94 h was near stoichiometric, albeit transient, in the mtri10 cultures. In the mtri1 cultures, the maximum amounts of accumulated N_2O were significantly lower ($P < 0.05$) than those in either the mtri0 or mtri10 cultures, in line with the substantially lower methanotroph population.

In the negative-control experiment performed with the Δ*mbnAN* mutant strain added to the soil suspension in place of the wild-type *M. trichosporium* OB3b (referred to as mtri10n), the maximum amount of N_2O accumulated in the vessel was limited to 0.27 $\pm$ 0.09 μmol N_2O-N, despite the *Methylocystaceae* dominance of the methanotrophic population, as indicated by the high *M. trichosporium* OB3b *pmoA* copy numbers [(1.2 $\pm$ 0.7) $\times$ 10^8 per ml] and the low *Methylococcaceae*/*Methylocystaceae* population ratio (0.21) in the mtri10n culture (Fig. 3). The results of this control experiment confirmed that the enhanced N_2O accumulation of the *Methylocysteceae*-dominant enrichment was due to copper sequestration by methanobactin produced by *M. trichosporium* OB3b. Also notable was the obvious dissimilarity observed in the microbial community compositions between the enrichments with the OB3b wild-type strain and the Δ*mbnAN* mutant strain (Fig. 3; Table S4). The additional set of control experiments performed with 2.0 μM $CuCl_2$ added to the soil-only, and thus *Methylococcaceae*-dominant, samples (referred to as mtri0c), suggested that the instantaneous N_2O accumulation observed in the *Methylococcaceae*-dominant culture could also be explained as the effect of Cu^{2+} sequestration by the methanotrophic population. The maximum N_2O production (3.75 $\pm$ 1.25 μmol N_2O-N at $t = 106$ h) was ~13 times lower with Cu^{2+} added than without, despite the abundance of *Methylococcaceae* family of methanotrophs [48.5% of (3.5 $\pm$ 0.2) $\times$ 10^8 eubacterial 16S rRNA copies ml^{-1}].

DISCUSSION

Copper deficiency has been suggested as a potential cause for N_2O emission from denitrification taking place in terrestrial and aquatic environments (32–34). In a previous study, Chang et al. demonstrated, using simplified pure-culture and coculture experiments, that MB-OB3b lowered the copper availability of the medium such that N_2O reduction was substantially repressed in denitrifiers (29). Whether methanobactin-mediated N_2O emission enhancement is relevant to actual soil environments with much more intricate microbiome complexity remained to be resolved, however. The results of the experiments in this study, although performed under well-controlled laboratory conditions, showed that methanobactin-enhanced N_2O emissions may actually occur in soil environments with complex microbiomes. Methanobactin addition exerted significant influence on denitrification carried out by the soils' indigenous microbial consortia, demonstrating that the diverse denitrifying community was still susceptible to methanobactin-mediated copper deprivation. Furthermore, the N_2O emission enhancement observed with the *Methylocystaceae* methanotroph-enriched reactor culture provided unprecedented experimental evidence of indigenous alphaproteobacterial methanotrophs influencing N_2O emission from denitrifiers. Observation of substantial *mbnA* transcription and the absence of this N_2O emission enhancement effect in the copper-amended sample supported that methanobactin produced by the indigenous

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Methylocystaceae methanotrophs was likely the cause of the increased N_2O production from denitrification. Additionally, the increased N_2O production observed in soil enrichments with broadly varied *Methylocystaceae*/*Methlycoccaceae* ratios suggested that methanotrophic population composition *in situ* may be a major determinant of the copper-mediated N_2O emission enhancement, while also suggesting the existence of an additional mechanism via which methanotrophs affect N_2O reduction in denitrifiers.

Evidence of the effect of methanobactin on the soil microbial consortia apart from N_2O emission enhancement could also be discerned. Delays in NO_2^- reduction during the progression of denitrification were observed in some, but not all, of the denitrifying enrichments incubated in the presence of OB3b-MB capable of producing methanobactin. At least two distinct forms of nitrite reductases are known to mediate NO_2^- to NO reduction in denitrifiers: the copper-dependent nitrite reductase NirK and copper-independent cytochrome cd_1 nitrite reductase NirS (35). In the previous investigation by Chang et al., indeed, NO_2^- reduction by NirK-utilizing *Shewanella loihica*, but not NirS-utilizing denitrifiers, was affected by methanobactin-mediated Cu deprivation (29). The environmental conditions that select for enrichment of either NirK- or NirS-utilizing organisms remain unclear, and abundances of *nirS*- or *nirK*-possessing organisms vary in rice paddy soils (36). Possibly, the relative abundances and activities of the NirK- and NirS-utilizing denitrifiers may have varied in the denitrifying consortia prepared with soils A to E, explaining the nonuniform responses of the denitrifiers in the consortia to copper deprivation.

The shotgun metagenome analyses of the methanotrophic chemostat culture identified organisms affiliated with the *Methylocystaceae* family as the dominant *nirK*-possessing organisms. Furthermore, the *nosZ* gene affiliated with *Methylocystis* sp. strain SC2 was the dominant *nosZ* gene in the chemostat culture (37). These observations are certainly interesting, as the only *Methylocystaceae nirK* and *nosZ* sequences available in the database are of the *nirK* gene found in the genome of *Methylocystis* sp. strain Rockwell and the *nosZ* gene found in a plasmid of *Methylocystis* sp. strain SC2 (37, 38). Both genes were unique, in the sense that they both shared $<$75% translated amino acid sequence identity with any other NirK or NosZ sequences in the NCBI database, and no previous study has reported recovery of these *Methylocystaceae nirK* and *nosZ* sequences in metagenome/metatranscriptome analyses. Despite the potential significance of the discovery, the possibility that the *Methylocystaceae* methanotrophs harboring these genes might have significantly affected NO_2^- reduction and N_2O production in the anoxic batch incubations was highly unlikely. Both *Methylocystis* sp. strain Rockwell and *Methylocystis* sp. strain SC2 have been confirmed to have the inability to grow anaerobically, and neither strain has been confirmed to be capable of reducing NO_2^- or N_2O utilizing NirK or NosZ, respectively (37, 38). Therefore, although the collected data are insufficient to completely rule out the possibility that these *Methylocystaceae* populations significantly contributed to denitrification and N_2O production and consumption, it is more plausible that the N_2O dynamics in the anoxic cultures were largely determined by nonmethanotrophic facultatively anaerobic organisms carrying *nirK*, *nirS*, and/or *nosZ* genes.

In this metagenome analysis, many of the nonmethanotrophic organisms putatively harboring *nirK* or *nirS* and those putatively harboring *nosZ* belonged to different taxa, suggesting that N_2O production and N_2O consumption may have been carried out by distinct organismal groups in the anoxic batch experiments performed with this chemostat culture. That is, the dominant non-*Methylocystaceae nirK* and *nosZ* genes recovered from the chemostat metagenome were those affiliated with *Rhizobiales* and *Bacteriodetes*, respectively, and only *nirK* and *nosZ* affiliated with *Pseudomonaceae* were recovered with similar coverage. This metagenomic observation may imply that the alteration of copper availability brought about by methanobactin-producing methanotrophs influence N_2O emissions from denitrification occurring in modular manner, which may be the more likely case in complex environmental microbiomes (39, 40).

The distinctive contrast between the microbial compositions of the mtri10 and

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mtri10n enrichments implied that methanobactin had a pronounced impact on the overall microbial community. The microbial community formed from enrichment on CH_4 and acetate after augmentation with the wild-type strain OB3b (mtri10) carried a distinctively large proportion of *Moraxellaceae* (44.7%). Such abundance of *Moraxellaceae* was not observed in the enrichment with augmented Δ*mbnAN* mutant cells (mtri10n), suggesting that this enrichment of *Moraxellaceae* was due to the presence of methanobactin produced by *M. trichosporium* OB3b. The most probable explanations are a selective bactericidal property and/or reduced copper bioavailability to competing organisms with high demands for copper (18, 22). The relative abundance of other phylogenetic groups, including *Methylophilaceae* and *Flavobactericeae*, also varied substantially across the treatments. Unlike the case for *Moraxellaceae*, however, the experimental evidence was not sufficient to attribute these alterations to the effect of methanobactin.

The observed N_2O emission enhancement in methanotroph-enriched cultures dominated by the *Methylococcaceae* family (i.e., the mtri0 enrichment, albeit to a level lower than that observed in the mtri10 enrichment), was unanticipated, as none of the sequenced genomes of the methanotrophs belonging to this phylogenetic group has been reported to produce methanobactin encoded by *mbnA* (18). What is evident from the experimental results, however, is that copper competition was central to N_2O production in these enrichments, as copper amendment removed the N_2O-accumulating phenotype, as observed in the mtri0c enrichment. Indications that *Methylomicrobium album* BG8 and *Methylococcus capsulatus* Bath may utilize methanobactin-like copper chelators had been previously reported (41, 42). These *Methylococcaceae* strains tested positive on the Cu-CAS (chromo azurol S) plate assays, and the putative methanobactin-like compound of ~1,000 Da in size isolated from the spent medium of *M. capsulatus* Bath bound Cu with 1:1 stoichiometry; however, the genomic basis for synthesis of these compounds in *Methylococcaceae* methanotrophs has not yet been elucidated. Another copper uptake mechanism involving copper-binding periplasmic membrane proteins MopE and CorA has been identified in *M. capsulatus* Bath and *M. album* BG8, respectively (43, 44). The estimated binding constants of these putative copper chelators are tens of orders of magnitudes lower than those reported for methanobactin from *Methylocystaceae* methanotrophs; however, if present at large concentrations, the copper chelators may still exert a significant impact on the Cu availability. Which, if any, of these mechanisms was responsible for copper withholding from denitrifiers and N_2O reducers in the *Methylococcaceae*-dominant enrichments cannot be determined with the current data and warrants future investigation.

One of the most prominent characteristics of soil environments is their spatial and temporal heterogeneity, in terms of both physicochemical makeup and microbial composition (45, 46). In CH_4-enriched soil environments such as rice paddy and landfill soils, proteobacterial methanotrophs are often reported to be abundant, with *pmoA* copy numbers ranging between 10^7 and 10^{10} gene copies g dry soil^{-1} (47–49). These numbers are of the same magnitude as the methanotrophic populations in the enrichment cultures observed to induce N_2O production from the cohabiting denitrifying population in the laboratory experiments performed here. Methanotroph population density at local microsites may even be higher, especially at the oxic-anoxic interfaces, where CH_4 and O_2, the essential substrates of proteobacterial methanotrophs, are both available (48). Thus, it would not be surprising to find local concentrations of methanobactin-producing *Methylocystaceae* methanotroph communities in microniches within the soil sufficiently dense as to cause copper deficiency to cohabiting NosZ-utilizing N_2O reducers. At a microscopic scale, temporal oxic-to-anoxic shifts or *vice versa* would constantly occur at the oxic-anoxic interfaces, allowing for spatial coexistence of O_2-dependent CH_4 oxidation and O_2-inhibited denitrification (31). The substrates of denitrification, NO_3^- and NO_2^-, may be transported from oxic surface soils or supplied via oxidation of organic N or NH_4^+ *in situ* at the oxic-anoxic interface with intermittently available O_2 (30). Periodic oxic-anoxic transitions may also take place in the vicinity of

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the water table in upland soils, as precipitation and drying cause fluctuations in the elevation of the water table (50). In such settings, aerobic microbial processes of nitrification and methanotrophy and anaerobic microbial processes of denitrification and methanogenesis may alternate at a larger scale. Snapshot views of physicochemical and biological states of soils may cast doubt on the likelihood of the methanotroph-enhanced N_2O emissions occurring in actual soil environments; however, with these spatial and temporal shifts in consideration, the suggested mechanism is plausible in any terrestrial environment with high organic and nitrogen content. Thus, in approximating greenhouse gas budgets from environments such as rice paddy soils, landfill cover soils, wetland soils, and upland agricultural soils, N_2O arising from the copper-mediated interaction between methanotrophs and N_2O reducers needs to be considered for development of a more accurate prediction model for N_2O emissions.

MATERIALS AND METHODS

Soil sampling and characterization. Soil samples were collected from an experimental rice paddy located at Gyeonggido Agricultural Research & Extension Services in Hwaseong, South Korea (37°13′21″N, 127°02′35″E), in August 2017 and four rice paddies near Daejeon, South Korea (36°22′41″N, 127°19′50″E), in December 2018 (Table S1). Samples were collected from the top layer of soil (0 to 20 cm below the overlying water). After removal of plant material, soil samples were stored in sterilized jars, which were then filled to the brim with rice paddy water. The samples were immediately transported to the laboratory in a cooler filled with ice and stored at 4°C until use. A small portion (~50 g) of each collected soil was stored at −80°C for DNA analyses.

The physicochemical characteristics of these soil samples were analyzed directly after sampling. The soil pH was measured by suspending 1 g wet weight soil in 5 ml Milli-Q water. Total nitrogen and carbon content of air-dried soil samples were analyzed with a Flash EA 1112 elemental analyzer (Thermo Fisher Scientific, Waltham, MA). For measurements of NH_4^+, NO_3^-, and NO_2^- content in soil samples, 1 g air-dried soil was suspended in 5 ml 2 M KCl solution and shaken at 200 rpm for an hour. After settling the suspension for 10 min, the supernatant was passed through a 0.2-μm-pore membrane filter (Advantec MFS, Inc., Tokyo, Japan). The filtrate was analyzed using colorimetric quantification methods. The total copper content of soil samples was measured with an Agilent ICP-MS 7700S inductively coupled plasma mass spectrometer (Santa Clara, CA) after pretreatment with boiling aqua regia (51).

Media and culture conditions. Unless otherwise mentioned, modified nitrate mineral salts (NMS) medium was used for enrichment of methanotrophs in soil microbial consortia and incubation of axenic cultures of the wild-type and Δ*mbnAN* mutant strains of *M. trichosporium* OB3b (52). The medium contained per liter, 1 g $MgSO_4{\cdot}7H_2O$ (4.06 mM), 0.5 g KNO_3 (4.95 mM), 0.2 g $CaCl_2{\cdot}2H_2O$ (1.36 mM), 0.1 ml of 3.8% (wt/vol) Fe-EDTA solution (Sigma-Aldrich, St. Louis, MO), 0.5 ml of 0.02% (wt/vol) $Na_2MoO_4{\cdot}2H_2O$ solution, and 0.1 ml of the 10,000× trace element stock solution prepared with 5 g liter^{-1} $FeSO_4{\cdot}7H_2O$, 4 g liter^{-1} $ZnSO_4{\cdot}7H_2O$, 200 mg liter^{-1} $MnCl_2{\cdot}7H_2O$, 500 mg liter^{-1} $CoCl_2{\cdot}6H_2O$, 100 mg liter^{-1} $NiCl_2{\cdot}6H_2O$, 150 mg liter^{-1} H_3BO_3, and 2.5 g liter^{-1} EDTA. All glassware used in this study was equilibrated in a 5 N HNO_3 acid bath overnight before use to reduce background contamination of copper to $<0.01\ \mu$M in media. Forty-milliliter aliquots of the medium were distributed into 250-ml serum bottles, and the bottles were sealed with butyl rubber stoppers (Geo-Microbial Technologies, Inc., Ochelata, OK) and aluminum caps. After autoclaving, the media were amended with 0.2 ml of 200× Wolin vitamin stock solution, and 500 mM pH 7.0 KH_2PO_4–Na_2HPO_4 solution was added to a final concentration of 5 mM (53). High-purity CH_4 (>99.95%; Deokyang Co., Ulsan, South Korea) replaced 20% of the headspace air. After inoculation, culture bottles were incubated in dark at 30°C with shaking at 140 rpm.

Analytical procedures. Headspace CH_4 concentrations were measured using a GCMS-QP2020 gas chromatograph-mass spectrometer (Shimadzu Cooperation, Kyoto, Japan) equipped with an SH-Rt-Q-BOND column (30 m in length by 0.32-mm inner diameter with 10-μm film thickness). The injector and oven temperatures were set to 150 and 100°C, respectively, and helium was used as the carrier gas. Headspace N_2O concentrations were measured with an HP 6890 series gas chromatograph equipped with a HP-PLOT/Q column and an electron capture detector (Agilent, Palo Alto, CA). The injector, oven, and detector temperatures were set to 200, 85, and 250°C, respectively. For each CH_4 or N_2O measurement, 50 or 100 μl of gas sampled using a Hamilton 1700-series gas-tight syringe (Reno, NV) was manually injected into the gas chromatographs. The dissolved N_2O concentrations were calculated from the headspace concentrations using the dimensionless Henry's law constant of 1.92 at 30°C (54, 55). The dissolved concentrations of NO_3^- and NO_2^- were determined colorimetrically using the Griess reaction (56). As the assay measures NO_2^-, vanadium chloride (VCl_3) was used to reduce NO_3^- to NO_2^-. The concentration of NH_4^+ was measured using the salicylate method (57).

The total bacterial population was quantified using TaqMan-based quantitative polymerase chain reactions (qPCRs) targeting the eubacterial 16S rRNA gene (1055f, 5′-ATGGCTGTCGTCAGCT-3′; 1392r, 5′-ACGGGCGGTGTGTAC-3′; and Bac1115_probe, 5′-CAACGAGCGCAACCC-3′) (58). The primers and probe set exclusively targeting the *pmoA* gene (encoding the β subunit of particulate methane monooxygenase) of *M. trichosporium* OB3b (OB3b_pmoAf, 5′-CGCTCGACATGCGGATAT-3′; OB3b_pmoAr, 5′-TTTCCCGATCAGCCTGGT-3′; and OB3b_ pmoA_probe, 5′-AGCCACAGCGCCGGAACCA-3′) were designed *de novo* using Geneious v9.1.7 software (Biomatters, Ltd., Auckland, New Zealand) and used for

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TABLE 1 Primer/probe sets used for qPCR and RT-qPCR analyses

Primer/probe set	Sequence	Target gene	Amplicon length (bp)	Slope	*y* intercept	Amplification efficiency	R^2	Reference
16S rRNA 1055f	5′-ATGGCTGTCGTCAGCT-3′	Bacterial 16S rRNA	338	−3.30	37.7	101.1	0.996	59
16S rRNA 1392r	5′-ACGGGCGGTGTGTAC-3′							
Bac1115_probe	5′-CAACGAGCGCAACCC-3′							
OB3b_pmoAf	5′-CGCTCGACATGCGGATAT-3′	*pmoA*	266	−3.38	36.4	97.8	0.998	This study
OB3b_pmoAr	5′-TTTCCCGATCAGCCTGGT-3′							
OB3b_ pmoA_probe	5′-AGCCACAGCGCCGGAACCA-3′							
mbnAf672	5′-GCTCGTCATACCATTCGGG-3′	*mbnA*	100	−3.32	37.7	100.1	0.999	This study
mbnAr771	5′-GCTTGGCGATACGGATGGTC-3′							
lucf	5′-TACAACACCCCAACATCTTCGA-3′	Luciferase control mRNA	67	−3.37	37.1	98.0	0.999	71
lucr	5′-GGAAGTTCACCGGCGTCAT-3′							

quantification of *M. trichosporium* OB3b cells. qPCR assays were performed with a QuantStudio 3 real-time PCR system (Thermo Fisher Scientific). The calibration curves for the targeted genes were constructed using serial dilutions of PCR2.1 vectors (Thermo Fisher Scientific) carrying the target fragments. For each qPCR quantification, three biological replicates were processed separately. The complete list of the primers and probes used in this study is provided in Table 1.

Monitoring of denitrification and N_2O production in rice paddy soil suspensions amended with methanobactin from *M. trichosporium* OB3b. Methanobactin was isolated from *Methylosinus trichosporium* OB3b cultures and purified as previously described (59). The effects of MB-OB3b on N_2O emissions were examined with suspensions of five rice paddy soils (soils A to E) with various physicochemical properties. Each soil suspension was prepared by suspending 1 g of air-dried rice paddy soil in 50 ml Milli-Q water in 160-ml serum bottles, and the headspace was replaced with N_2 gas after sealing. Methanobactin stock solution was prepared by dissolving 10 μmol (11.5 mg) freeze-dried methanobactin in 10 ml Milli-Q water. The stock solution was equilibrated in an anaerobic chamber filled with 95% N_2 and 5% H_2 (Coy Laboratory Products, Inc., Grass Lake, MI) for 1 h to remove dissolved O_2, and 0.5 ml of the solution was added to the soil suspensions. Potassium nitrate and sodium acetate were then added to final concentrations of 5 and 10 mM, respectively. Soil suspensions were incubated with shaking at 140 rpm at 30°C. The headspace N_2O concentration and the dissolved concentrations of NO_3^- and NO_2^- were monitored until depletion of NO_3^- and NO_2^-. The dissolved concentrations of NH_4^+ were also monitored to ensure the absence of significant dissimilatory nitrate reduction to ammonium (DNRA) activity. Negative controls were prepared identically, but without methanobactin amendment. Abiotic control experiments with sterilized soils with and without methanobactin amendment were performed with 5 mM NO_3^- or NO_2^- added, to confirm that the abiotic contribution to N_2O production was minimal.

Monitoring of denitrification and N_2O production in subcultures extracted from a quasi-steady-state chemostat culture of *Methylocystaceae*-dominated soil methanotrophs. A continuous methanotrophic enrichment culture was prepared with soil E. The inoculum was generated by suspending 1 g (wet weight) soil in 50 ml NMS medium in a sealed 160-ml serum vial with air headspace. The vials were fed twice with 41 μmol CH_4, yielding a headspace concentration of ~0.5% (vol/vol) upon each injection. After the initial batch cultivation, 20 ml of the enrichment was transferred to 5 liters NMS medium in a 6-liter fermentor controlled with a BioFlo 120 Bioprocess Control Station (Eppendorf, Hamburg, Germany). The gas stream carrying a relatively low CH_4 concentration (0.5% [vol/vol] in air) was fed continuously at a rate of 145 ml min^{-1} through a gas diffuser to stimulate growth of *Methylocystaceae* methanotrophs, as previous fed-batch and chemostat incubation of rice paddy soils with a continuous supply of 20% (vol/vol) CH_4 resulted in domination by gammaproteobacterial methanotrophs (60). The reactor culture was maintained in the dark at pH 6.8 and 30°C with agitation at 500 rpm. After the initial fed-batch operation, the reactor culture was transitioned to continuous operation with the dilution rate set to 0.014 h^{-1}. After the quasi-steady state was attained, as indicated by constant effluent cell density and CH_4 concentration, 1.0-ml aliquots were collected, and the pellets were stored at −20°C for analyses of the microbial community composition and denitrification functional genes and qPCR quantification of *mbnA* and 16S rRNA genes. Additionally, 0.4-ml aliquots were treated with RNAprotect bacterial reagent (Qiagen, Hilden, Germany) and stored at −80°C for reverse transcription-qPCR (RT-qPCR) analyses of *mbnA* expression.

Triplicate anoxic batch cultures were prepared by distributing 50 ml culture harvested from the running quasi-steady-state reactor to 160-ml serum bottles and flushing the headspace with N_2 gas. A denitrifying enrichment was prepared separately by incubating 1 g (wet weight) of soil E in 100 ml anoxic NMS medium amended with 10 mM sodium acetate. One milliliter of this denitrifier inoculum was added to the methanotroph enrichments, to which 20 mM sodium acetate was added as the electron donor for denitrification. The cultures were amended with or without 2 μM $CuCl_2$, the culture vessels were

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incubated in the dark with shaking at 140 rpm at 30°C, and the changes to the amounts of NO_3^-, NO_2^-, and N_2O-N in the vessels were monitored until no further change was observed.

Denitrification and N_2O production in soil enrichments with altered methanotrophic population composition. Various amounts of preincubated *M. trichosporium* OB3b cells were added to rice paddy soil slurry microcosms before enrichment with CH_4, to artificially vary the proportion of the methanobactin-producing subgroup among the total methanotroph population. The total indigenous gammaproteobacterial methanotroph population in the soil suspension was approximated by multiplying the total eubacterial 16S rRNA copy number per ml suspension (as determined with qPCR) by the proportion of the 16S rRNA gene sequences affiliated with gammaproteobacterial methanotrophs (all affiliated with the *Methylococcaceae* family) in soil B (as computed from the MiSeq amplicon sequencing data [see Table S5 in the supplemental material]). The numbers of *M. trichosporium* OB3b cells in the precultures were estimated with the TaqMan qPCR targeting the duplicate *pmoA* genes of the strain. The *M. trichosporium* OB3b preculture was used to prepare soil suspensions with the augmented *Methylocystaceae* population. The ratios of *M. trichosporium* OB3b *pmoA* copy numbers to the estimated 16S rRNA gene copy numbers of the *Methylococcaceae* methanotrophs were set to 1 and 10 in the 40-ml suspensions. It should be stressed that these ratios cannot be directly translated to cell number ratios, as the numbers of 16S rRNA genes in the completed genomes of gammaproteobacterial methanotrophs deposited in the NCBI GenBank database vary between 1 and 4. Soil suspensions without *M. trichosporium* OB3b augmentation were also prepared. As a negative control, the Δ*mbnAN* mutant strain of *M. trichosporium* OB3b was added in place of the wild-type strain OB3b (61). A copper-replete control was also prepared for the experimental condition with no *M. trichosporium* OB3b augmentation by amending a subset of cultures with 2 μM $CuCl_2$.

The soil suspensions were prepared by adding 0.1 g wet weight of the soil to the autoclaved modified NMS medium bottles prepared as described above (40 ml NMS medium in 250-ml serum bottles). After sealing of the bottles and addition of *M. trichosporium* OB3b or Δ*mbnAN* mutant preculture or $CuCl_2$ to the calculated target concentrations, 20% of the headspace was replaced with CH_4 and sodium acetate was added to a final concentration of 5.0 mM. When the headspace CH_4 concentration decreased to 10%, the headspace was replenished with an 80:20 mixture of air and CH_4 and the cultures were amended with 200 μmol NO_3^- and 400 μmol sodium acetate. At each sampling time point, the headspace N_2O concentration was measured and 0.5 ml of the culture was collected from each bottle for determination of NO_3^-, NO_2^-, and NH_4^+ concentrations. The cell pellets collected immediately after the CH_4 concentration dropped to ~10% were subjected to qPCR targeting strain OB3b *pmoA* genes and eubacterial 16S rRNA genes. The 16S rRNA amplicon sequences of the pellets were also analyzed.

Microbial community composition analyses and prediction of genomic potentials for denitrification reactions from 16S rRNA amplicon sequences. Genomic DNA was extracted with the DNeasy PowerSoil kit (Qiagen, Hilden, Germany). The V6 to V8 region of the 16S rRNA gene amplified with the universal primers 926F (5′-AAACTYAAAKGAATTGRCGG-3′)/1392R (5′-ACGGGCGGTGTGTRC-3′) was sequenced at Macrogen, Inc. (Seoul, South Korea), using a MiSeq sequencing platform (Illumina, San Diego, CA) (62). The raw sequence data were processed using the QIIME pipeline v1.9.1 with the parameters set to default values (63). After demultiplexing and quality trimming, the filtered sequences were clustered into operational taxonomic units (OTUs) using the USEARCH algorithm, and the OTUs were assigned a taxonomic classification using the RDP classifier against the Silva SSU database release 132. The raw sequences were deposited into the NCBI SRA database (BioProject accession number PRJNA685001).

Metagenomic analysis of the chemostat enrichment and RT-qPCR targeting *mbnA*. Shotgun metagenomic sequencing of the reactor culture was performed at Macrogen, Inc., using a HiSeq X sequencing platform (Illumina) generating ~5 Gb of paired-end reads with 150-bp read length (BioProject accession number PRJNA684997). The raw short reads were quality screened using Trimmomatic v0.36 with default parameters (64). The processed reads were assembled into contigs using metaSPAdes v3.12.0, and coding sequences were identified using Prodigal v2.6.3 (65, 66). The Hidden Markov model (HMM) algorithms for *nirS*, *nirK*, clade I and clade II *nosZ*, and *recA* were downloaded from the FunGene database (http://fungene.cme.msu.edu/). A previously constructed HMM algorithm for *mbnA* was used for screening of *mbnA* genes (67). The predicted contigs were screened for these targeted genes using *hmmsearch* in HMMER 3.1b2, with the E value set to 10^{-5}. The identified gene sequences were further validated by running blastx against the RefSeq database, with the E value cutoff set to 10^{-3}. For taxonomy assignment, the translated amino acid sequences of the screened functional genes were searched against NCBI's nonredundant protein database (nr) using blastp, with an E value cutoff of 10^{-10}. The quality-trimmed reads were mapped onto the screened functional gene sequences using the bowtie2 v2.3.4.1 software, with the parameters set to default values (68). The coverage of each target gene was calculated using the *genomecov* command of the BEDtools v2.17.0 software and normalized by its length (69).

A degenerate primer set was designed with the sole unique *mbnA* sequence recovered from the shotgun metagenome data and publicly available *Methylocystis mbnA* sequences with >85% translated amino acid sequence identity with this metagenome-derived *mbnA* sequence (Fig. S1), using Geneious v9.1.7 software (Biomatters, Ltd., Auckland, New Zealand). This primer set (672f, 5′-GCTCGTCATA CCATTCGGG-3′; and 771r, 5′-GCTTGGCGATACGGATGGTC-3′) was used for quantification of *mbnA* genes and transcripts in the chemostat culture (Fig. S1). Extraction and purification of RNA were performed using the RNeasy minikit (Qiagen) and RNA MinElute kit (Qiagen), respectively, and reverse transcription was performed using SuperScript III reverse transcriptase (Invitrogen, Carlsbad, CA), according to the established protocols (70). As previously described, a known quantity of luciferase control mRNA (Promega, Madison, WI) was used as the internal standard to correct for RNA loss during the extraction

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and purification procedure. The *mbnA* transcript copy numbers were normalized to *mbnA* copy numbers in the genomic DNA.

Statistical analyses. All experiments were performed in triplicate, and the average values are presented with the error bars representing the standard deviations from triplicate samples. The statistical significance of the data from two different experimental conditions was tested with two-sample *t* tests, and statistical comparisons between two different time points were tested with one-sample *t* tests. The SPSS Statistics 24 software (IBM Corp., NY) was used for statistical analyses.

Data availability. The raw short reads from 16S rRNA amplicon sequencing and shotgun metagenome sequencing were deposited in the NCBI BioProject database under accession numbers PRJNA684997 and PRJNA685001.

SUPPLEMENTAL MATERIAL

Supplemental material is available online only.

SUPPLEMENTAL FILE 1, PDF file, 2.3 MB.

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The authors declare they have no conflicts of interest.

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