

Nitrous oxide (N₂O) synthesis by the freshwater cyanobacterium *Microcystis aeruginosa*

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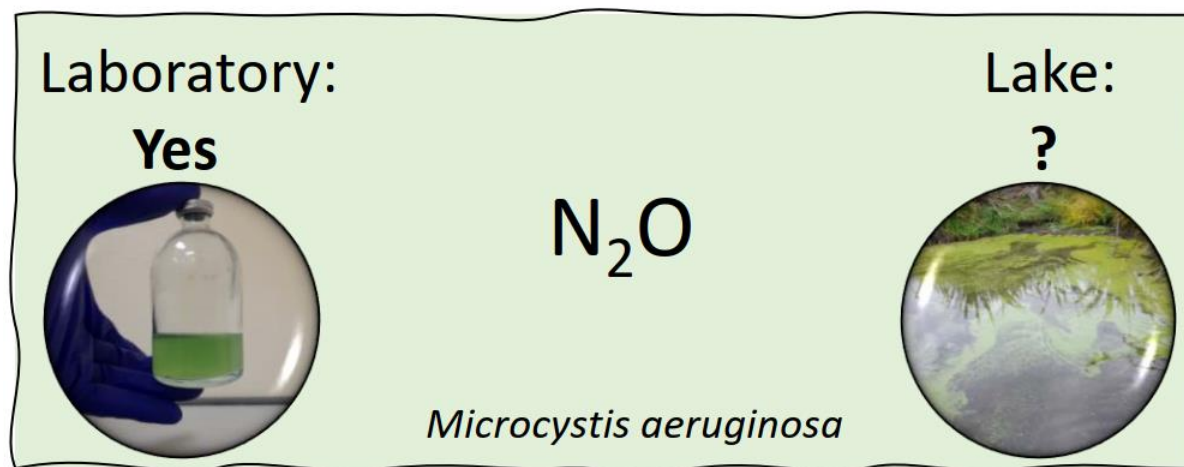
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Abstract. Pure cultures of the freshwater cyanobacterium *Microcystis aeruginosa* synthesized nitrous oxide (N₂O) when supplied with nitrite (NO₂⁻) in darkness (198.9 nmol·g-DW⁻¹·h⁻¹ after 24 hours) and illumination (163.1 nmol·g-DW⁻¹·h⁻¹ after 24 hours) whereas N₂O production was negligible in abiotic controls supplied with NO₂⁻ and in cultures deprived of exogenous nitrogen. N₂O production was also positively correlated to the initial NO₂⁻ and *M. aeruginosa* concentrations, but low to negligible when nitrate (NO₃⁻) and ammonium (NH₄⁺) were supplied as the sole exogenous N source instead of NO₂⁻. A protein database search revealed *M. aeruginosa* possesses protein homologues to eukaryotic microalgae enzymes known to catalyse the successive reduction of NO₂⁻ into nitric oxide (NO) and N₂O. Our laboratory study is the first demonstration that *M. aeruginosa* possesses the ability to synthesize N₂O. As *M. aeruginosa* is a bloom-forming cyanobacterium found globally, further research (including field monitoring) is now needed to establish the significance of N₂O synthesis by *M. aeruginosa* under relevant conditions (especially in terms of N supply). Further work is also needed to confirm the biochemical pathway and potential function of this synthesis.

Graphical Abstract.



1 Introduction

Emissions of the potent ozone depleting greenhouse gas nitrous oxide (N_2O) have been reported from various aquatic ecosystems characterized by a high level of photosynthetic activity and several authors have suggested that N_2O emissions from eutrophic lakes could be globally significant (Delsontro et al., 2018; Plouviez et al., 2019a). Noteworthy, Delsontro et al. (2018) determined that N_2O emissions from lakes and impoundments could be expected to increase as a function of lake size and chlorophyll a (an indicator of the presence of primary producer such as microalgae). Because eutrophication is an increasing global issue (Delsontro et al., 2018; Kapsalis and Kalavrouziotis, 2021; Maure et al., 2021), N_2O emissions from these ecosystems could also be expected to increase. Several species of microalgae and cyanobacteria can indeed synthesize N_2O (Weathers, 1984; Weathers and Niedzielski, 1986; Bauer et al., 2016; Plouviez et al., 2019a) and a biochemical pathway for this synthesis has been established in the model microalga *Chlamydomonas reinhardtii* (Plouviez et al., 2017b; Burlacot et al., 2020). Despite these critical advances, the true global significance of microalgal N_2O synthesis in microalgae-rich eutrophic aquatic bodies remains unknown (Plouviez et al., 2019a; Burlacot et al., 2020; Plouviez and Guieysse, 2020). Microcystis species are cyanobacteria commonly found in eutrophic ecosystems (Xiao et al., 2018; Zhou et al., 2020; Hernandez-Zamora et al., 2021) but the ability of this genus to synthesize N_2O is currently unknown. We, therefore, investigated the ability of N_2O production by the most notorious bloom-forming cyanobacterium reported in freshwaters and model cyanobacterium *M. aeruginosa* (Qian et al., 2010; Kataoka et al., 2020; Zhou et al., 2020) under conditions known to induce or impact N_2O production in microalgae (Guieysse et al., 2013; Alcantara et al., 2015; Bauer et al., 2016; Plouviez et al., 2017b; Burlacot et al., 2020).

2 Results and discussion

2.1 N_2O synthesis bioassays

The ability of *M. aeruginosa* to synthesize N_2O was investigated using a protocol successfully used for the microalgae *C. vulgaris* and *C. reinhardtii* (Alcantara et al., 2015; Guieysse et al., 2013; Plouviez et al., 2017b). As can be seen in **Fig. 1**, N_2O production was only recorded in cultures supplied NO_2^- as there was no significant production in the absence of the cyanobacterium (abiotic control) or the absence of NO_2^- (negative control). Further assays showed a positive correlation between biomass concentration and N_2O production (**Fig. 2**), confirming the biological origin of N_2O synthesis.

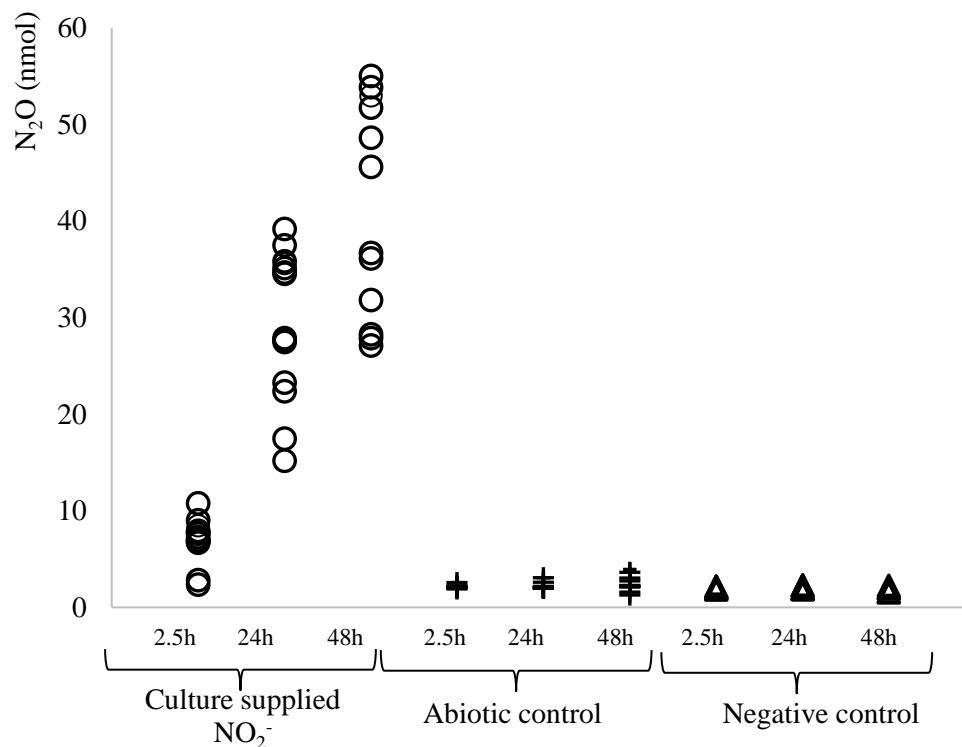


Figure 1. Total N_2O accumulation (nmole) from *M. aeruginosa* supplied with 10 mM NO_2^- under continuous illumination (\circ , $n \geq 12$), abiotic control N-free media with 10 mM NO_2^- (+, $n \geq 10$) and negative control: *M. aeruginosa* cultures incubated in N-free media (Δ , $n \geq 10$).

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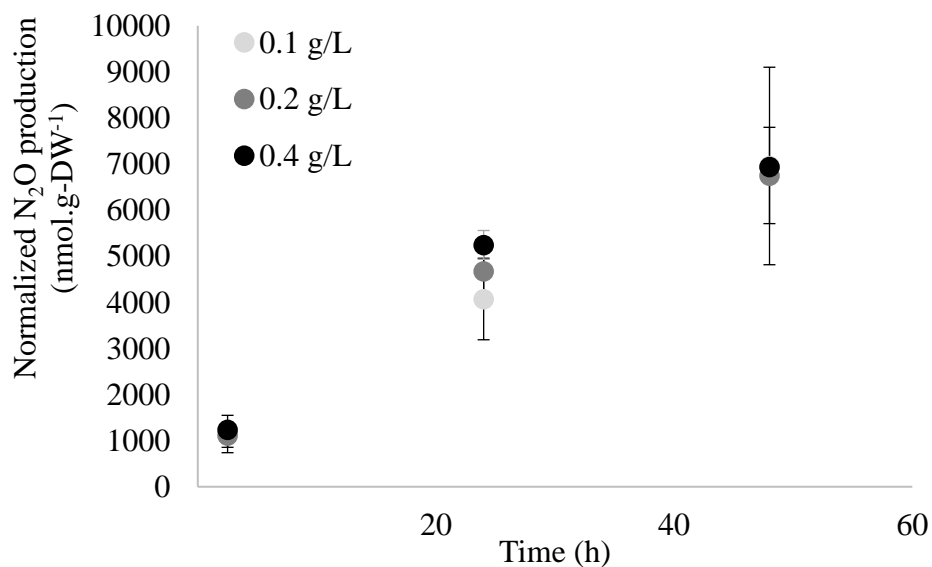


Figure 2. Normalized N₂O production (nmol·g-DW⁻¹) recorded from *M. aeruginosa* cultures with different biomass concentrations (0.1, 0.2 and 0.4 g-DW·L⁻¹; n ≥ 6, n ≥ 12, n = 4, respectively) in sealed flasks supplied light and 10 mM NO₂⁻. N₂O synthesis was statistically different when comparing the rates between 2.5 and 24 h and between 24 and 48 h (p < 0.05, two samples t-test). Specific N₂O production rates (nmol N₂O·g DW⁻¹·h⁻¹) can be found in Table S1.1.

In comparison to cultures supplied with NO₂⁻, low and negligible N₂O synthesis was recorded in cultures supplied with NO₃⁻ and NH₄⁺, respectively (**Table. 1**). This result showed that NO₂⁻ was the substrate to N₂O synthesis, as reported for other microalgae (Weathers, 1984; Weathers and Niedzielski, 1986; Guieysse et al., 2013; Alcantara et al., 2015; Bauer et al., 2016; Plouviez et al., 2017b; Burlacot et al., 2020). The positive correlation between NO₂⁻ concentration and N₂O synthesis also confirmed that NO₂⁻ was the substrate to N₂O synthesis (**Figure S2.1**, Vmax = 185 nmol·g-DW⁻¹·h⁻¹ and Km for NO₂⁻ = 2.22 mM).

Table 1. N₂O emissions in different conditions (n = number of replicates)

Light conditions	N source	N ₂ O production (nmol·g-DW ⁻¹ ·h ⁻¹)	Standard error	n
Light	1 mM NO ₂ ⁻	59.5	13.7	18
	5 mM NO ₂ ⁻	131.5	21.5	16
	10 mM NO ₂ ⁻	163.1	31.5	23
	10 mM NO ₃ ⁻	3.9	1.4	6
	10 mM NH ₄ ⁺	0.07	0.7	4
	-	0.9	0.5	4
Dark	10 mM NO ₂ ⁻	198.9	30.5	5
	-	1.5	1.7	6

M. aeruginosa was able to synthesize N₂O in both darkness and illumination (**Table. 1**), representing 0.07% and 0.06% of the amount of N supplied (g-N-N₂O produced/g-N supplied × 100) respectively. The N₂O produced under illumination was statistically lower than in darkness (p-value < 0.05, two samples t-test, n = 5 replicates from experiments performed on the same day). The negative impact of light was previously observed in *C. vulgaris* and *C. reinhardtii* tested under similar conditions (Guieysse et al., 2013; Alcantara et al., 2015; Plouviez et al., 2017b), although N₂O production was positively correlated with light supply in *C. vulgaris* grown outdoors (Plouviez et al., 2017a). The difference we observed during this study may be explained by light-dependent mechanisms impacting enzymatic activities and consequently intracellular NO₂⁻ accumulation (e.g. the rates of NO₂⁻ reduction into NH₄⁺ and N₂O), as suggested by Plouviez et al, (2017a). However, O₂ production during photosynthesis could also influence N₂O synthesis. Burlacot et al. (2020) indeed reported that one of the

75 enzymes involved in NO reduction to N₂O (Flavodiiron, as discussed in the next section) can also catalyse the reduction of O₂ into H₂O. Because of this dual activity and the reactivity of NO with O₂, N₂O production could be sensitive to O₂. Further research is therefore needed to understand if O₂ influence N₂O production by competitive NO conversion to products such as nitrogen oxides and peroxynitrite, or/and by competitive O₂ reduction into H₂O instead of its reduction to N₂O by the enzymes with nitric reductase ability.

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While small, N₂O synthesis was statistically significant in *M. aeruginosa* fed NO₃⁻ as the sole exogenous N source (p-value < 0.05, two samples t-test when compared with the negative controls). As in *C. vulgaris* and *C. reinhardtii*, the intracellular reduction of NO₃⁻ into NO₂⁻ by the enzyme nitrate reductase (narB) is the first step of NO₃⁻ assimilation in *M. aeruginosa* (Ohashi et al., 2011; Zhou et al., 2020). Hence, intracellular NO₂⁻ production likely generated this substrate for N₂O synthesis during NO₃⁻ exogenous supply but competitive use of NO₂⁻ (for protein synthesis via NH₄⁺ generation) could have competed with N₂O synthesis. Intracellular NO₂⁻ production and accumulation is not expected when cells assimilate NH₄⁺ (Plouviez et al., 2019), explaining the absence of N₂O production in the flasks supplied NH₄⁺ as sole exogenous N source (p-value = 0.91, two samples t-test when compared with the negative controls). In *M. aeruginosa*, NO₃⁻ uptake and the transcriptional regulation of nitrate reductase have been shown to be activated by light, NO₃⁻ and NO₂⁻ (Chen et al., 2009; Ohashi et al., 2011; Chen and Liu, 2015). While the transcriptional and post-translational regulation of nitrate reductase in *M. aeruginosa* still needs to be investigated in relation to N₂O synthesis and varying environmental parameters (e.g. light supply), it is possible that the pattern of N₂O synthesis during outdoor *M. aeruginosa* growth would be similar to that seen in *C. vulgaris*.

2.2 Putative pathways

In the eukaryotic microalga *C. reinhardtii*, cytoplasmic NO₂⁻ is sequentially reduced to nitric oxide (NO) and N₂O. The first step, NO₂⁻ reduction into NO, is catalysed by the dual enzyme nitrate reductase-NO forming nitrite reductase (NR-NoFNiR) or, potentially, the copper containing nitrite reductase (NirK). The second step, NO reduction into N₂O, can then be catalysed by cytochrome P450 (CYP55, Plouviez et al., 2017b; Burlacot et al., 2020), Flavodiirons (FLVs, Burlacot et al., 2020; Bellido-Pedraza et al., 2020), or potentially by the Hybrid Cluster proteins (HCPs, Bellido-Pedraza et al., 2020) involved in nitrogen metabolism (Van Lis et al., 2020). Interestingly, NO₂⁻ reduction into NO by nitrate reductase (narB) has been demonstrated in *M. aeruginosa* (Tang et al., 2011; Song et al., 2017) and here we found that *M. aeruginosa* possesses homologs of the CYP55, FLVs, and HCPs found in *C. reinhardtii* (Table. 2). While the functions of these proteins need to be confirmed, their presence suggests N₂O synthesis in *M. aeruginosa* could involve similar NO₂⁻ and NO reduction pathways to those found in *C. reinhardtii*.

105 **Table 2. Summary of Blastp results for proteins potentially involved in N₂O synthesis in *Chlamydomonas reinhardtii*. Accession numbers were retrieved from (Bellido-Pedraza et al., 2020) and used as query sequence for blastp (protein-protein BLAST) protein searches (<https://blast.ncbi.nlm.nih.gov/Blast.cgi>) of *M. aeruginosa* (taxid:1126) protein sequences database.**

Protein	<i>C. reinhardtii</i> accession number	e-value	<i>M. aeruginosa</i> accession number	% Similarity	<i>M. aeruginosa</i> protein
NirK	PNW79625.1	-		-	-
HCP	XP_001694756.1	3e-158	NCR75269.1	45.38	Hydroxylamine reductase
	XP_001694571.1	5e-160	WP_002787796.1	44.79	Hydroxylamine reductase
	XP_001694671.1	2e-157	NCR75269.1	45.03	Hydroxylamine reductase
	XP_001694454.1	2e-159	WP_002787796.1	45.96	Hydroxylamine reductase
CYP55	XP_001700272.1	3e-45	NCR09918.1	29.90	CYP55
FLV	XP_001692916.1	6e-138	WP_193956217.1	43.45	Diflavin flavoprotein
FLV	PNW71243.1	0	WP_110545956.1	52.18	Diflavin flavoprotein

2.3 Metabolic function

110 The metabolic function of N₂O synthesis in eukaryotic microalgae is currently unknown and it has been suggested that NO₂⁻ reduction into N₂O enables cells to expend excess energy or instead, is the fortuitous result of dual enzymatic activity (Guieysse et al., 2013; Plouviez et al., 2017b). The intermediate NO is a ubiquitous signalling molecule in algae (Astier et al., 2021). Interestingly, NO stimulates the production of secondary metabolites (*e.g.* linoleic acid) by *M. aeruginosa* that inhibit the growth of competitors (Song et al., 2017). NO also promotes the growth of this cyanobacterium (Tang et al., 2011). While the link between NO and N₂O still needs to be determined, it is possible that the NO and N₂O biosynthetic pathways is/are involved in cell-to-cell communications in *M. aeruginosa* and more broadly, in microalgae.

2.4 Potential environmental implications

Microalgae species from at least 3 divisions (Bacillariophyta, Chlorophyta, Cyanobacteria) have the ability to synthesize NO (Kim et al., 2008; Kumar et al., 2015; Plouviez et al., 2017b; Tang et al., 2011) and/or N₂O (Weathers, 1984; Weathers and Niedzielski, 1986; Guieysse et al., 2013; Kamp et al., 2013; Plouviez et al., 2017a, b, this study). All these observations suggest that the ability to synthesize N₂O is widely distributed among microalgae. Critically, N₂O emissions from aquatic environments where microalgae abound, such as oceans, lakes and engineered cultivation systems, have been repeatedly reported (Bauer et al., 2016; Plouviez et al., 2019b; Plouviez et al., 2019a; Zhang et al., 2022) even under very low exogenous NO₂⁻ concentrations (Plouviez et al., 2019b). These emissions can be explained by intracellular NO₂⁻ production during reductive nitrate assimilation (Plouviez et al., 2017a, b, 2019b) under conditions when excess NO₂⁻ production (Bristow et al., 2015; French et al., 1983; Mortonson and Brooks, 1980; Schaefer and Hollibaugh, 2018) could support N₂O synthesis.

Based on the data available, DelSontro et al. (2018) and Plouviez and Guieysse, (2020) estimated that global N₂O emissions from eutrophic lakes alone could represent 110 to 450 kt N-N₂O·yr⁻¹, which represent 14-56% of the natural and anthropogenic N₂O emissions reported from inland and coastal waters (Tian et al., 2020). Importantly, DelSontro et al. (2018) predicted that N₂O emissions from lakes and impoundments would increase with lake size and chlorophyll a concentration. The N₂O synthesis rates reported during our study are in the same order of magnitude as the rate previously reported for members of the green microalgae, cyanobacteria, and diatoms (Bauer et al., 2016; Plouviez et al., 2019a). However, we cannot conclude that *M. aeruginosa* (or other species) is or is not a major N₂O producer in lakes and other aquatic environments without evidence from field measurements. Indeed, high NO₂⁻ concentrations are rare in natural and engineered ecosystems environments, which would suggest insignificant microalgal N₂O production in most contexts. Nevertheless, significant N₂O emissions were reported from outdoor cultures of *C. vulgaris* fed NO₃⁻ (Guieysse et al., 2013; Plouviez et al., 2017), despite this alga also producing much more N₂O when fed NO₂⁻ (Guieysse et al., 2013). Plouviez et al. (2017) suggested this was caused by NO₂⁻ intracellular accumulation under varying light, as this condition is known to have different impacts on the rate of NO₃⁻ reduction into NO₂⁻ by NR and the rate of NO₂⁻ reduction into NH₄⁺ by NiR. During our study, N₂O emissions under NO₃⁻ supply were low, but not negligible. Because NR activity is also influenced by light and the availabilities of NO₃⁻ and NO₂⁻ in *M. aeruginosa* (Chen et al., 2009; Ohashi et al., 2011; Chen and Liu, 2015), N₂O synthesis by this microalga could possibly occur in environments where NO₃⁻ is the main nitrogen source.

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Our findings support past predictions of the global relevance of photosynthetic N₂O emissions from eutrophic aquatic bodies as *Microcystis* is globally found and often the dominant genus in these ecosystems (Qian et al., 2010; Kataoka et al., 2020; Zhou et al., 2020). The work from Weathers and Niedzielski, (1986) and ours suggest that *Nostoc spp.*, *Aphanocapsa* (PCC 6308), *Aphanocapsa* (PCC 6714) and *M. aeruginosa* have the ability to synthesize N₂O. Consequently, other cyanobacteria species may also have this ability. Further research is now needed to quantify N₂O emissions from eutrophic aquatic ecosystems where cyanobacteria abound. This is especially timely considering that the frequency and geographic distribution of harmful algae blooms have increased due to anthropogenic activities (Paerl et al., 2018; Kataoka et al., 2020). In addition, algae blooms can lead to the decrease of O₂ in oceans, coastal waters and lakes (Jenny et al., 2015; Rabalais and Turner, 2019), a condition that can increase the accumulation of NO₂⁻ in aquatic ecosystems (Schaefer and Hollibaugh, 2018; Bristow et al., 2015). Because microalgal N₂O synthesis is rapid and influenced by factors such as the cell biology (Plouviez et al., 2019b) and, as observed during our study, the type and concentration of the nitrogen source microalgae receive, extensive monitoring (i.e. long-term with wide spatial coverage and high sampling frequency) of several types of microalgae-rich environments are required (e.g. hypoxic waters).

3 Conclusions

160 We herein present the first demonstration that *M. aeruginosa* synthesizes N₂O. *Microcystis aeruginosa* synthesized N₂O when supplied with NO₂⁻ in darkness (198.9 nmol·g-DW⁻¹·h⁻¹ after 24 hours) and illumination (163.1 nmol·g-DW⁻¹·h⁻¹ after 24 hours), and this production was positively correlated to the initial NO₂⁻ and *M. aeruginosa* concentrations. A protein database search also revealed *M. aeruginosa* possesses proteins homologues to eukaryotic microalgae known to catalyse the successive reduction of NO₂⁻ into NO and N₂O. Further studies are needed to confirm the genes/proteins involved as a better understanding of the biochemical pathway involved during microalgal N₂O synthesis is critical to efficiently monitor (*i.e.* identify the source) and mitigate N₂O emissions.

Our study is another evidence of the ability of photosynthetic microorganisms, especially cyanobacteria, to synthesize N₂O. Preliminary estimation showed that N₂O emissions from eutrophic lakes alone could represent 110 to 450 kt N-N₂O·yr⁻¹, which represent 14-56% of the natural and anthropogenic N₂O emissions reported from inland and coastal waters. However, how much microalgae contribute to these emissions is currently unknown. As *M. aeruginosa* is globally distributed, further research (including field monitoring with wide spatial coverage, high sampling frequency and water type) is now needed to evaluate the significance of N₂O synthesis by these cyanobacteria under relevant conditions (especially in terms of N supply).

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4 Appendix: Materials and Methods

4.1 Strain and culture maintenance

Microcystis aeruginosa UTEX 2385 was obtained from the culture collection of the University of Texas at Austin (<https://utex.org/>). Pure cultures were maintained on 100 mL low-phosphate minimal media (Plouviez et al., 2021) incubated at 25°C (INFORS HT Multitron) under continuous illumination (20 μmol·cm⁻²·s⁻¹) and agitation (150 rotation per minutes, rpm). Cultures thus incubated for more than a week were supplied with 100 μL of a solution of KH₂PO₄/K₂HPO₄ (0.4 M/0.6 M) to prevent P limitation. The purity of the cultures was verified via sequencing (S3).

4.2 Cultivation and Bioassays

M. aeruginosa was cultivated on 400 mL low-phosphorus minimal media in 500 mL Duran bottles for 5 days. These cultures were incubated under fluorescent tubes (F15W/GRO sylvania gro-lux) providing illumination at 20 μmol·cm⁻²·s⁻¹ at the culture surface. Mixing was provided by bubbling filtered (0.22μm) air at 1.5 L·min⁻¹. On the day of the experiment, 15 mL aliquots were withdrawn from the cultures to measure the cell dry weight (DW) according to (Bechet et al., 2015). Then, 100-400 mL aliquots were centrifuged at 4400 rpm for 3 min. The supernatants were discarded, and the pellets were re-suspended in N-free

medium to a final concentration of 0.2 g-DW L⁻¹ as previously described (Guieysse et al., 2013). Twenty-five mL aliquots of these suspensions were transferred into 120 mL serum flasks supplied with 1 mL of NaNO₂, NaNO₃ or NH₄Cl stock solutions (250 mM) to reach a final concentration of 10 mM. Sterile abiotic controls were not inoculated but were supplied with 10 mM nitrite (NO₂⁻) while negative controls were *M. aeruginosa* cultures incubated in N-free media. The flasks were immediately sealed with rubber septa and aluminium caps and incubated at 25°C under continuous agitation (150 rpm) under either constant illumination (20 µmol·cm⁻²·s⁻¹) or darkness. A similar protocol was used to evaluate the impact of the initial cell (0.1 – 0.4 g-DW·L⁻¹), NO₂⁻ (1 – 10 mM) or nitrate (NO₃⁻, 10 mM) or ammonium (NH₄⁺, 10 mM) concentrations on N₂O synthesis. Unless otherwise stated, each condition was tested in triplicate flasks and repeated at least twice. All glassware and media were autoclaved prior to the experiments. An additional experiment confirmed the purity of the *M. aeruginosa* stock cultures and the cultures used during the bioassays (S3).

4.3 Analysis

Gas samples (5 mL) were withdrawn from the flask headspace using a syringe equipped with a needle. The headspace N₂O concentration in those samples was then quantified using gas chromatography (Shimadzu GC-2010, Shimadzu, Japan). Total N₂O was calculated as the sum of gaseous N₂O and dissolved N₂O as described by Guieysse et al. (2013). Briefly, Assuming the gas and the liquid phase N₂O concentrations were at equilibrium at the time of sampling, the total amount of N₂O produced in the flask was calculated by summing up the amounts of N₂O present in the gas and liquid phases. The amount of dissolved N₂O in the liquid phase was calculated using Henry's law at 25°C (Eq. 1):

$$n_{N_2O_{total}}^t = x_{N_2O}^t \cdot P^t \cdot \left(\frac{V_g}{R \cdot T} + H_{N_2O} \cdot V_l \right) \quad (1)$$

Where $n_{N_2O_{total}}^t$ is the total amount of N₂O produced in the Duran bottle at time t (moles N₂O); $x_{N_2O}^t$ is the molar fraction of N₂O in the gas phase at time t (mol N₂O·mol gas⁻¹); P^t is the pressure in the gas headspace at time t (typically 101325 Pa unless otherwise stated); V_g is the volume of gas in the flask (mL); R is the ideal gas constant (8.314 J·mol⁻¹·K⁻¹); T is the temperature inside the bottle (298.15 K); H_{N₂O} is the Henry law constant of N₂O at T (2.5·10⁻⁷ mol·L⁻¹·Pa⁻¹); and V_l is the volume of liquid in the serum flask (mL).

Supplementary information

Specific N₂O production rates are presented in Table S1.1. The enzymatic kinetic of N₂O synthesis in *M. aeruginosa* is shown in Figure S2.1. The purity of *M. aeruginosa* cultures was confirmed by PCR and sequencing: The methodology used and the results obtained are presented in the supplement S3.

Authors contribution

F.F. performed the investigation, data visualization and curation, and contributed to the writing - review & editing of the manuscript. M.P. was involved with the writing - original draft and contributed to conceptualization, methodology, and data curation and visualization with B.J. B.J. and J.P. were involved with the writing - review & editing of the manuscript before submission. Finally, B.J., J.P. and M.P. were all involved with the funding acquisition.

Competing interests

The authors declare that they have no conflict of interest.

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