

1 Methane, carbon dioxide and nitrous oxide emissions from two clear- 2 water and two turbid-water urban ponds in Brussels (Belgium)

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7 **Abstract.** Shallow ponds can occur either in a clear-water state dominated by macrophytes or a turbid-water state dominated
8 by phytoplankton, but it is unclear if and how these two states affect the emission to the atmosphere of greenhouse gases
9 (GHGs) such as carbon dioxide (CO₂), methane (CH₄) and nitrous oxide (N₂O). We measured on 46 occasions over 2.5 years
10 (between June 2021 and December 2023) the dissolved concentration of CO₂, CH₄, and N₂O from which the diffusive air-
11 water fluxes were computed, in four urban ponds in the city of Brussels (Belgium): two clear-water macrophyte-dominated
12 ponds (Silex and Tenreuken), and two turbid-water phytoplankton-dominated ponds (Leybeek and Pêcherries). CH₄ ebullitive
13 fluxes were measured with bubble traps in the four ponds during deployments in spring, summer, and fall, totalling 48 days
14 of measurements. To characterize methanogenic pathways (acetoclastic or hydrogenotrophic) and quantify water column
15 methane oxidation (MOX) we measured the ¹³C/¹²C ratio of CH₄ (δ¹³C-CH₄) from gas trapped in the bubble traps, from
16 bubbles deliberately released by the perturbation of the sediments, and in dissolved CH₄ in the water column. Measured
17 ancillary variables include water temperature, oxygen saturation level (%O₂), concentrations of chlorophyll-*a* (Chl-*a*), total
18 suspended matter (TSM), soluble reactive phosphorus (SRP), nitrite (NO₂⁻), nitrate (NO₃⁻) and ammonium (NH₄⁺). The
19 turbid-water and clear-water ponds did not differ significantly in terms of diffusive emissions of CO₂ and N₂O. Clear-water
20 (macrophyte-dominated) ponds exhibited higher values of annual ebullitive CH₄ fluxes compared to turbid-water
21 (phytoplankton-dominated) ponds, most probably in relation to the delivery to sediments of organic matter from
22 macrophytes. At seasonal scale, CH₄ emissions exhibited a temperature dependence in all four ponds, with ebullitive CH₄
23 fluxes having a stronger dependence to temperature than diffusive CH₄ fluxes. The temperature sensitivity of ebullitive CH₄
24 fluxes decreased with increasing water depth. In summer, the δ¹³C-CH₄ values of sediment bubbles indicated that the
25 hydrogenotrophic methanogenesis pathway seemed to dominate in clear-water ponds and acetoclastic methanogenesis
26 pathway seemed to dominate in turbid-water ponds. The δ¹³C-CH₄ values of bubbles traps suggested a seasonal shift from
27 the acetoclastic methanogenesis pathway in spring-summer to the hydrogenotrophic methanogenesis pathway in fall. The
28 δ¹³C-CH₄ of dissolved CH₄ indicated higher rates of MOX in turbid-water ponds compared to clear-water ponds, with an
29 overall positive correlation with TSM and Chl-*a* concentrations. The presence of suspended particles putatively enhanced
30 MOX by reducing light inhibition of MOX and/or by serving as substrate in the water column for fixed methanotrophic
31 bacteria. Total CH₄ emissions (diffusive+ebullitive) in CO₂ equivalents either equalled or exceeded those of CO₂, while N₂O
32 emissions were negligible compared to the other two GHGs. Total annual GHG emissions in CO₂ equivalents from all four
33 ponds increased from 2022 to 2023 due to higher CO₂ diffusive fluxes, likely driven by higher annual precipitation in 2023
34 compared to 2022, possibly in response to the intense El Niño event of 2023.

35 **1. Introduction**

36 Greenhouse gas (GHG) emissions from inland water (rivers, lakes, and reservoirs) to the atmosphere such as carbon dioxide
37 (CO₂), methane (CH₄) and nitrous oxide (N₂O) are quantitatively important for global budgets (Lauerwald et al., 2023).
38 GHG emissions from lakes are lower than from rivers for CO₂ (Raymond et al., 2013) and N₂O (Lauerwald et al., 2019;
39 Maavara et al., 2019). However, emissions of CH₄ from lakes (Rosentreter et al., 2021; Johnson et al., 2022) are significant
40 compared to rivers (Stanley et al., 2016; Rocher-Ros et al., 2023). Emissions of CO₂ and CH₄ from lakes to the atmosphere
41 represent 1.25 to 2.30 Pg CO₂ equivalents (CO₂-eq) annually with a significant proportion from CH₄ emissions, and
42 represent nearly 20% of global CO₂ emissions from fossil fuels (Delsontro et al., 2018). The contribution of CO₂ and CH₄
43 emissions from small lentic water bodies (small lakes and ponds) can be disproportionately high compared to large systems
44 (Holgerson and Raymond, 2016) as small lakes and ponds are the most abundant of all water body types in number
45 (Verpoorter et al., 2014, Cael et al., 2017), and flux intensities (per m²) are usually higher in smaller water bodies. The
46 emissions of GHGs from artificial water bodies such as agricultural reservoirs, urban ponds, and storm-water retention
47 basins could be higher than those from natural systems (Martinez-Cruz et al., 2017; Grinham et al., 2018; Herrero Ortega et
48 al., 2019; Gorsky et al., 2019; Ollivier et al., 2019; Peacock et al., 2019, 2021; Webb et al., 2019; Bauduin et al., 2024).
49 These higher emissions seem to result from higher external inputs of anthropogenic carbon and nitrogen in artificial systems
50 such as rainfall runoff that brings organic matter and dissolved inorganic nitrogen (DIN), but might also reflect other
51 differences compared to natural systems such as in hydrology (Clifford and Heffernan, 2018). Among artificial systems,
52 urban ponds are the subject of a growing body of literature (Singh et al., 2000; Natchimuthu et al., 2014; van Bergen et al.,
53 2019; Audet et al., 2020; Peacock et al., 2021; Goeckner et al., 2022; Ray and Holgerson, 2023; Bauduin et al., 2024). Urban
54 areas can have numerous small artificial water bodies mostly associated to green spaces such as public parks, and their
55 number is increasing due to rapid urbanisation worldwide (Brans et al., 2018; Audet et al., 2020; Gorsky et al., 2024; Rabaey
56 et al., 2024). Urban ponds are generally small, shallow, and usually their catchment consists in majority of impervious
57 surfaces with a smaller contribution from soils (Davidson et al., 2015; Peacock et al., 2021).

58 In shallow ponds and lakes, including urban ponds, aquatic primary production is either dominated by submerged
59 macrophytes or by phytoplankton, corresponding to two alternate states (Scheffer et al., 1993). These two alternative states
60 correspond to clear waters (macrophyte-dominated) or turbid waters (phytoplankton-dominated), during the productive
61 period of the year (spring and summer in mid-latitudes). Submerged macrophytes and phytoplankton regulate CO₂ dynamic
62 directly through photosynthesis that can be more or less balanced by community respiration in the water column. However, it
63 is not clear whether the presence of macrophytes increases or decreases the CO₂ emissions from ponds and lakes. Some
64 studies have shown a decrease of CO₂ emissions with increasing macrophyte density (Kosten et al., 2010; Ojala et al., 2011;
65 Davidson et al., 2015), but other studies showed the opposite pattern (Theus et al., 2023). In phytoplankton-dominated lakes,
66 CO₂ concentrations depend in part on the development stage of the phytoplankton, with the growth and peak phases
67 generally coinciding with lower CO₂ concentrations due to intense photosynthesis (Grasset et al., 2020; Vachon et al., 2020).
68 CH₄ emissions have been reported to increase with the concentration of chlorophyll-*a* (Chl-*a*) in phytoplankton-dominated
69 lakes (DelSontro et al., 2018; Borges et al., 2022). The presence of macrophytes strongly affects CH₄ cycling in freshwaters
70 (Bastviken et al., 2023) and vegetated littoral zones of lakes exhibit higher CH₄ emissions than non-vegetated zones
71 (Desrosiers et al., 2022; Theus et al., 2023). Macrophytes influence organic matter decomposition processes in sediments
72 depending on the quality and quantity of plant matter they release into their environment (Reitsema et al., 2018; Grasset et
73 al., 2019; Harpenslager et al., 2022; Theus et al., 2023). Yet, few studies have consistently compared CH₄ emissions in clear-
74 water and turbid-water ponds (Hilt et al., 2017). A study in Argentina reported higher dissolved CH₄ concentrations in clear-

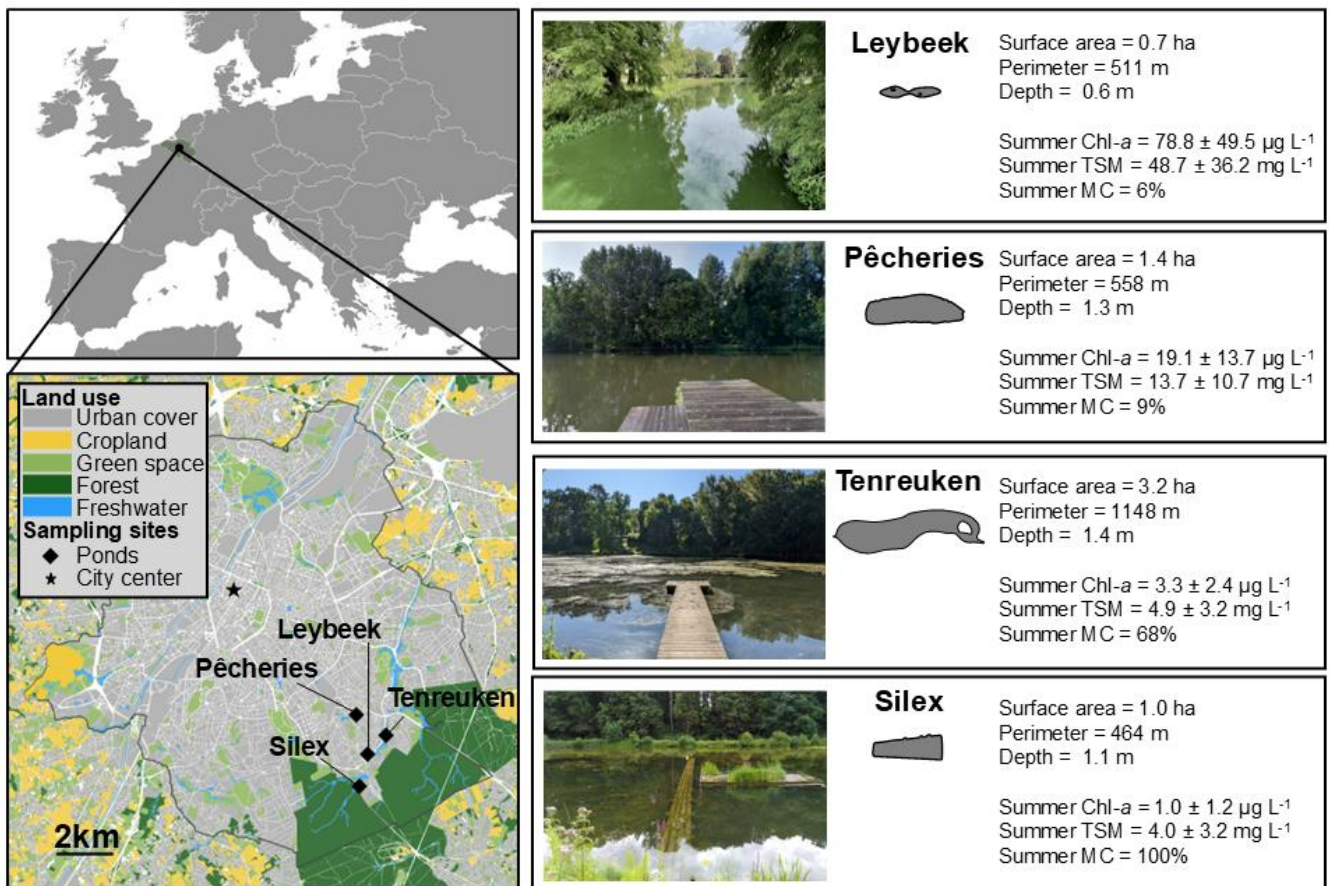
75 water ponds with submerged macrophytes compared to turbid-water phytoplankton-dominated ponds, but no differences in
76 measured CH₄ emissions (Baliña et al., 2023). The production of N₂O predominantly occurs through microbial nitrification
77 and denitrification that depend on DIN and O₂ levels (Codispoti and Christensen, 1985; Mengis et al., 1997). Competition
78 for DIN between primary producers and N₂O-producing microorganisms can impact N₂O production. Additionally, the
79 transfer of labile phytoplankton organic matter to sediments fuels benthic denitrification. Combined, these two processes
80 could explain that some lakes can act as sinks of N₂O under elevated Chl-*a* concentrations (Webb et al., 2019; Borges et al.,
81 2022). The presence of macrophytes also strongly influences nitrogen cycling in sediments of lakes and ponds (Barko et al.,
82 1991; Choudhury et al., 2018; Deng et al., 2020; Dan et al., 2021) and should in theory also affect N₂O emissions, although
83 seldom investigated, and available studies provide contradictory conclusions. N₂O emissions has been showed to follow
84 diurnal cycles of O₂ concentrations in areas dominated by submerged macrophytes in Lake Wuliangshuai (China) (Ni et al.,
85 2022) and the seasonal cycle of aboveground biomass of emerged macrophytes (*Phragmites*) in Baiyangdian Lake (China)
86 (Yang et al., 2012). On the contrary, some studies showed there were no significant differences of N₂O production in
87 sediments of macrophyte-rich (n=10) and macrophyte-free (n=12) lakes in subtropical China (Liu et al., 2018). There have
88 been a very limited number of studies investigating systematically how emissions differ between ponds dominated by
89 phytoplankton and those dominated by macrophytes (Harpenslager et al., 2022; Baliña et al., 2023), and none investigating
90 simultaneously CO₂, CH₄, and N₂O emissions including both diffusive and ebullitive components.

91 The emissions from aquatic systems of CO₂ and N₂O are exclusively through diffusion across the air-water interface
92 (diffusive flux), while CH₄ can be additionally emitted as bubbles released from sediments to the atmosphere (ebullitive
93 flux). At annual scale, ebullitive CH₄ flux usually represents more than half of total (diffusive+ebullitive) CH₄ emissions
94 from shallow lakes (Wik et al., 2013; Deemer and Holgerson, 2021), although the relative contribution of ebullitive and
95 diffusive CH₄ emissions is highly variable seasonally (*e.g.* Wik et al., 2023; Ray and Holgerson, 2023). Ebullitive CH₄
96 fluxes are particularly high in the littoral zone of lakes at depths <5 m (Wik et al., 2013; DelSontro et al., 2016; Borges et al.,
97 2022) and strongly increase in response to temperature (DelSontro et al., 2016; Aben et al., 2017), as well as organic matter
98 availability (DelSontro et al., 2016; 2018). Ebullitive CH₄ fluxes tend to be higher in small and shallow water bodies
99 (Deemer and Holgerson, 2021) but are notoriously variable in time and space, and are difficult to estimate reliably
100 (DelSontro et al., 2011).

101 The two primary metabolic pathways for CH₄ production in sediments by methanogenic archaea are the fermentation of
102 acetate (acetoclastic pathway) and the reduction of carbon dioxide by H₂ (hydrogenotrophic pathway) (Whiticar et al., 1986;
103 Conrad, 1989). CH₄ produced by these two pathways exhibits distinct ¹³C/¹²C ratios (δ¹³C-CH₄) (Whiticar et al., 1986) and
104 can be used to discriminate which pathway is dominant. When CH₄ diffuses from sediments to the water column, it can be
105 oxidized by methanotrophic bacteria who preferentially consume CH₄ with ¹²C over ¹³C, resulting in an increase of δ¹³C-CH₄
106 of the residual CH₄ in the water column (Bastviken et al., 2002). Fractionation models then allow estimating methane
107 oxidation (MOX) from measurements of δ¹³C-CH₄ of dissolved CH₄ in the water column. Bastviken et al. (2008) report that
108 30 to 99% of the CH₄ produced in sediments of freshwater lakes can be removed by MOX that is as a significant CH₄ sink in
109 these water bodies. MOX is known to be inhibited by light (Dumestre et al., 1998) and increases with the presence
110 suspended particles (Abril et al. 2007) so that MOX might vary between clear and turbid waters (Morana et al. 2020).

111 Here, we report a dataset of CO₂, CH₄, and N₂O dissolved concentrations in four shallow and small urban ponds (Leybeek,
112 Pêcherries, Silex, and Tenreuken) in the city of Brussels (Belgium) (Fig. 1), with data collected 46 times at regular intervals
113 (between June 2021 and December 2023) on each pond. The air-water diffusive fluxes of CO₂, CH₄, and N₂O were

114 calculated from dissolved concentrations and the gas transfer velocity, while the ebullitive CH₄ fluxes were measured with
 115 inverted funnels during 8 deployments (totalling 48 days) in the four ponds. The δ¹³C-CH₄ in the sedimentary bubbles and in
 116 the water provides additional information on CH₄ dynamics such as the methanogenesis pathway (acetoclastic or
 117 hydrogenotrophic) and MOX. We test the hypothesis that the two alternative states in shallow lakes (a clear-water state
 118 dominated by macrophytes, or a turbid-water state dominated by phytoplankton) drive differences in the CO₂, CH₄, and N₂O
 119 dissolved concentration and diffusive emissions from the four studied artificial ponds, that have similar depth, surface area,
 120 and catchment urban coverage, and that mainly differ by the phytoplankton-macrophyte dominance. We also test the
 121 hypothesis that the two alternative states in shallow lakes drive differences in the ebullitive CH₄ emissions, water column
 122 MOX, and sedimentary methanogenesis pathway (acetoclastic or hydrogenotrophic) in the four studied ponds. The final
 123 objective of the present work is to determine the relative contribution of CO₂, CH₄, and N₂O to the total GHG emissions in
 124 CO₂-eq and to test the hypothesis that the relative contribution of each GHG differs according to the two alternative states in
 125 shallow lakes.



126

127 **Figure 1: Location of the four sampled ponds in Brussels (Belgium, Europe). Bottom left map shows the metropolitan area of the**
 128 **region of Brussels delineated by the black line and the surrounding region of Flanders in Belgium, showing land cover and**
 129 **sampld urban ponds (black diamonds). The star corresponds to the center of the city (50.8504°N, 4.3487°E). Additional**
 130 **information for each pond is indicated on right panels: shapes of the ponds, surface area (ha), perimeter (m), average depth (m),**
 131 **mean±standard deviation of summer chlorophyll-*a* (Chl-*a*, in µg L⁻¹) and summer total suspended matter (TSM, in mg L⁻¹) from**
 132 **21 June to 21 September in 2021, 2022, 2023, and summer total macrophyte cover (MC, in %) (Table S1).**

133 **2. Material and Methods**

134 **2.1. Field sampling and meteorological data**

135 Sampling was carried out from a pontoon in the four ponds on the same day between 9am and 11am, 46 times on each pond
136 between June 2021 and December 2023 at a frequency ranging from one (winter) to three (summer) times per month at a
137 single fixed station in each of the four ponds. Water was sampled 5cm below the surface with 60ml polypropylene syringes
138 for analysis of dissolved concentrations of CO₂, CH₄, and N₂O. Samples for CH₄ and N₂O were transferred from the syringes
139 with a silicone tube into 60 ml borosilicate serum bottles (Weathon), preserved with 200 µl of a saturated solution of HgCl₂,
140 sealed with a butyl stopper and crimped with aluminium cap, without a headspace, samples were stored at ambient
141 temperature protected from direct light prior to analysis in laboratory. The partial pressure of CO₂ (pCO₂) was measured
142 directly in the field, within 5 minutes of sample collection, with a Li-Cor Li-840 infrared gas analyser (IRGA) based on the
143 headspace technique with 4 polypropylene syringes (Borges et al., 2019). A volume of 30 ml of sample water was
144 equilibrated with 30 ml of atmospheric air within the syringe by shaking vigorously for 5 minutes. The headspace of each
145 syringe was then sequentially injected into the IRGA and a fifth syringe was used to measure atmospheric CO₂. The final
146 pCO₂ value was computed taking into account the partitioning of CO₂ between water and the headspace, as well as
147 equilibrium with HCO₃⁻ (Dickson et al., 2007) using water temperature measured in-situ and after equilibration, and total
148 alkalinity (data not shown). Samples for total alkalinity were conditioned, stored and analysed as described by Borges et al.
149 (2019). The IRGA was calibrated in the laboratory with ultrapure N₂ and a suite of gas standards (Air Liquide Belgium) with
150 CO₂ mixing ratios of 388, 813, 3788 and 8300 ppm. The precision of pCO₂ measurements was ±2.0%. Water temperature,
151 specific conductivity, and oxygen saturation level (%O₂) were measured in-situ with VWR MU 6100H probe 5cm below the
152 surface. A 2 liter polyethylene water container was filled with surface water for conditioning the samples for other variables
153 at the laboratory in Université Libre de Bruxelles.

154 Surveys to identify and quantify visually the relative coverage of emerged and submerged macrophytes were conducted in
155 summer 2023 (Table S1). The resulting list of macrophyte species agreed with past studies in Brussels ponds (Peretyatko et
156 al., 2009).

157 Three bubble traps were deployed 50 cm apart for measuring ebullitive CH₄ flux. The bubble traps consisted of inverted
158 polypropylene funnels (diameter 23.5 cm) mounted with 60 ml polypropylene syringes, with three way stop valves allowing
159 to collect the gas without contamination from ambient air. The polypropylene funnel was attached with steel rods to a
160 polystyrene float. The volume of gas collected in the funnels was sampled with graduated polypropylene 60 ml syringes
161 every 24 hours. The value of the collected volume of gas was logged, and the gas was transferred immediately after
162 collection to pre-evacuated 12 ml vials (Exetainers, Labco, UK) that were stored at ambient temperature protected from
163 direct light prior to the analysis of CH₄ concentration and δ¹³C-CH₄ in the laboratory. The time-series of measurement were
164 longer at the Silex pond than the other three ponds, because the Silex pond is closed to the public during the week, while the
165 other three ponds are open to the public all the time.

166 In summer 2023, the bubbles present in the sediment were directly collected with bubble traps by physically perturbing the
167 sediment below the traps with a wooden rod. The gas collected in the funnels was stored in pre-evacuated 12 ml vials
168 (Exetainers, Labco, UK) that were stored at ambient temperature protected from direct light prior to the analysis of δ¹³C-CH₄
169 in the laboratory. These samples are referred hereafter to as from “perturbed sediments.” The samples collected in the bubble
170 traps during the ebullition measurements are referred to as from “trapped bubbles.”

171 Air temperature, precipitation, wind speed, and atmospheric pressure, were retrieved from <https://wow.meteo.be/en> for the
172 meteorological station of the Royal Meteorological Institute of St-Lambert (50.8408°N, 4.4234°E) in Brussels, located
173 between 2.5 and 5.0 km from the surveyed ponds. Air temperature, wind speed and atmospheric pressure were averaged over
174 24 h to obtain a daily mean value. Precipitation was integrated each day to obtain cumulated daily rainfall.

175 **2.2. Laboratory analysis**

176 **2.2.1. Chlorophyll-*a*, total suspended matter, and dissolved inorganic nutrients**

177 Water was filtered through Whatman GF/F glass microfiber filters (porosity 0.7 μm) with a diameter of 47 mm for total
178 suspended matter (TSM) and Chl-*a*. Filters for TSM were dried in an oven at 50 °C and filters for Chl-*a* were kept frozen (-
179 20 °C). The weight of each filter was determined before and after filtration of a known volume of water using an Explorer™
180 Pro EP214C analytical microbalance (accuracy ± 0.1 mg) for determination of TSM concentration. Chl-*a* concentration was
181 measured on extracts with 90% acetone by fluorimetry (Kontron model SFM 25) (Yentsch and Menzel, 1963) with a limit of
182 detection of 0.01 $\mu\text{g L}^{-1}$. Filtered water was stored frozen (-20 °C) in 50 ml polypropylene bottles for analysis of dissolved
183 nutrients. Soluble reactive phosphorus (SRP) was determined by the ammonium molybdate, ascorbic acid and potassium
184 antimony tartrate staining method (Koroleff, 1983), with a limit of detection of 0.1 $\mu\text{mol L}^{-1}$. Ammonium (NH_4^+) was
185 determined by the nitroprusside-hypochlorite-phenol staining method (Grasshoff and Johannsen, 1972), with a limit of
186 detection of 0.05 $\mu\text{mol L}^{-1}$. Nitrite (NO_2^-) and nitrate (NO_3^-) were determined before and after reduction of NO_3^- to NO_2^- by a
187 cadmium-copper column, using the Griess acid reagent staining method (Grasshoff and Kremling, 2009), with a detection
188 limit of 0.01 and 0.1 $\mu\text{mol L}^{-1}$, respectively. Concentration of dissolved inorganic nitrogen (DIN) was calculated as the sum
189 NH_4^+ , NO_2^- and NO_3^- concentrations in $\mu\text{mol L}^{-1}$.

190 **2.2.2. CH₄ and N₂O measurements by gas chromatography and $\delta^{13}\text{C}$ -CH₄ by cavity ring-down spectrometry**

191 Measurements of N₂O and CH₄ concentrations dissolved in water and in the gas samples from bubbles were made with the
192 headspace technique (Weiss 1981) with an headspace volume of 20ml of ultra-pure N₂ (Air Liquid Belgium) and a gas
193 chromatograph (GC) (SRI 8610C) with a flame ionisation detector for CH₄ and an electron capture detector for N₂O
194 calibrated with CH₄:N₂O:N₂ gas mixtures (Air Liquide Belgium) with mixing ratios of 1, 10 and 30 ppm for CH₄, and 0.2,
195 2.0 and 6.0 ppm for N₂O. The precision of measurement based on duplicate samples was $\pm 3.9\%$ for CH₄ and $\pm 3.2\%$ for N₂O.

196 The CO₂ concentration is expressed as partial pressure ($p\text{CO}_2$) in parts per million (ppm) and CH₄ as dissolved concentration
197 (nmol L^{-1}), as frequently used in topical literature. CH₄ concentration were systematically and distinctly above saturation
198 level (2-3 nmol L^{-1}) and $p\text{CO}_2$ values were only five times below saturation out of the 187 measurements. The N₂O
199 concentrations fluctuated around atmospheric equilibrium, so data are presented as percent of saturation level (%N₂O, where
200 atmospheric equilibrium corresponds to 100%). The equilibrium with atmosphere for N₂O was calculated from the average
201 air mixing ratios of N₂O provided by the Global Monitoring Division (GMD) of the National Oceanic and Atmospheric
202 Administration (NOAA) Earth System Research Laboratory (ESRL) (Dutton and Hall, 2023), and using the Henry's
203 constant given by Weiss and Price (1980).

204 The $\delta^{13}\text{C}$ -CH₄ was measured in the gas of the headspace (20 ml of synthetic air, Air Liquid Belgium) equilibrated with the
205 water sample (total volume 60 ml) for water samples and directly in the gas stored in Exetainers for samples from the bubble
206 traps. The gas samples were diluted to obtain a final partial pressure of CH₄ in the cavity below 10 ppm (target value of 6
207 ppm) to fall within the operational concentration range of the instrument recommended by the manufacturer, prior to
208 injection into a cavity ring-down spectrometer (G2201i, Isotopic Analyzer, Picarro) with a Small Sample Introduction

209 Module 2 (SSIM, Picarro). Data were corrected with curves of $\delta^{13}\text{C-CH}_4$ as a function of concentration based on two gas
210 standards from Airgas Specialty Gases with certified $\delta^{13}\text{C-CH}_4$ values of $-23.9\pm 0.3\text{‰}$ and $-69.0\pm 0.3\text{‰}$.

211 2.3. Calculations

212 2.3.1. Diffusive GHG emissions

213 The diffusive air-water CO_2 , CH_4 , or N_2O fluxes (F_G) were computed according to:

$$214 F_G = k\Delta[G], \quad (1)$$

215 where k is the gas transfer velocity and $\Delta[G]$ is the air-water gas concentration gradient.

216 The atmospheric pCO_2 was measured in the field with the Li-Cor Li-840. For CH_4 , the global average present day
217 atmospheric mixing ratio of 1.9 ppm was used (Lan et al., 2024). Atmospheric N_2O concentration was calculated from the
218 average air mixing ratios of N_2O provided by the GMD of the NOAA ESRL (Dutton et al., 2017). k was computed from a
219 value normalized to a Schmidt number of 600 (k_{600}) and from the Schmidt number of CO_2 , CH_4 and N_2O in freshwater
220 according to the algorithms as function of water temperature given by Wanninkhof (1992). k_{600} was calculated from the
221 parameterization as a function of wind speed of Cole and Caraco (1998). CH_4 and N_2O emissions were converted into CO_2
222 equivalents ($\text{CO}_2\text{-eq}$) considering a 100-year timeframe, using global warming potential of 32 and 298 for CH_4 and N_2O ,
223 respectively (Myrhe et al., 2013).

224 2.3.2. Ebullitive flux

225 Bubble flux ($\text{ml m}^{-2} \text{d}^{-1}$) measured with the inverted funnels was calculated according to:

$$226 F_{bubble} = \frac{V_g}{A \times \Delta t}, \quad (2)$$

227 where V_g is the volume of gas collected in the funnels (ml), A is the cross-sectional area of the funnel (m^2), Δt is the
228 collection time (d).

229 A multiple linear regression model of F_{bubble} dependent on water temperature and drops of atmospheric pressure was fitted to
230 the data according to:

$$231 \log_{10}(F_{bubble}) = \alpha \times T_w + \beta \times \Delta p, \quad (3)$$

232 where α and β are the slope coefficients of the multiple linear regression model, T_w is the water temperature ($^\circ\text{C}$), and Δp
233 quantifies the drops in atmospheric pressure (atm), calculated according to Zhao et al. (2017):

$$234 \Delta p = -\frac{1}{\Delta t} \int_0^t p - p_0; \quad \forall p < p_0, \quad (4)$$

235 where p is the atmospheric pressure (atm), p_0 a threshold pressure fixed at 1 atm and Δt the time interval between two
236 measurements (d) (Fig. S1).

237 Ebullitive CH_4 fluxes ($\text{mmol m}^{-2} \text{d}^{-1}$) were calculated according to:

$$238 E_{CH_4} = [\text{CH}_4] \times F_{bubble}, \quad (5)$$

239 where $[CH_4]$ is the CH_4 concentration in bubbles ($mmol\ ml^{-1}$).

240 The methane ebullition Q_{10} represents the proportional change in the ebullitive CH_4 flux per $10^\circ C$ change in water
241 temperature (DelSontro et al., 2016) and was computed according to:

$$242 \quad Q_{10} = 10^{10b} , \quad (6)$$

243 where b is the slope of the linear regression between the logarithm of the ebullitive CH_4 flux (E_{CH_4}) and T_w , and c is the y -
244 intercept, according to:

$$245 \quad \log_{10}(E_{CH_4}) = b \times T_w + c , \quad (7)$$

246 The flux of CH_4 from dissolution of rising bubbles was computed using the model of McGinnis et al. (2006) implemented in
247 the SiBu-GUI graphical user interface (Greinert and McGinnis, 2009).

248 **2.3.3. Methane oxidation**

249 The fraction of CH_4 oxidized (FOX) was calculated with a closed-system Rayleigh fractionation model (Liptay et al., 1998)
250 according to:

$$251 \quad \ln(1 - FOX) = \frac{\ln(\delta^{13}C-CH_4_{initial}+1000)-\ln(\delta^{13}C-CH_4 + 1000)}{\alpha - 1} , \quad (8)$$

252 where $\delta^{13}C-CH_4_{initial}$ is the $^{13}C/^{12}C$ ratio of dissolved CH_4 as produced by methanogenesis in sediments, $\delta^{13}C-CH_4$ is the
253 $^{13}C/^{12}C$ ratio of dissolved CH_4 in-situ, and α is the fractionation factor.

254 We used a value of 1.02 for α based on laboratory culture experiments carried out at $26^\circ C$ (Coleman et al., 1981) and field
255 measurements in three Swedish lakes (Bastviken et al., 2002) and one tropical lake (Morana et al., 2015). The α values
256 gathered in the three Swedish lakes were independent of season and temperature according to Bastviken et al. (2002).

257 For $\delta^{13}C-CH_4_{initial}$, we used a value of -69‰ for spring and summer, and -83‰ for fall based on average of measured $\delta^{13}C$ -
258 CH_4 in trapped bubbles (see Sect. 3.5). For winter we used a value of -76‰ corresponding to the average of the fall and
259 spring/summer values.

260 MOX was computed from FOX and the F_G of CH_4 (F_{CH_4}) according to (Bastviken et al., 2002):

$$261 \quad MOX = F_{CH_4} \times \frac{FOX}{1-FOX} , \quad (9)$$

262 **2.4. Statistical analysis**

263 Statistical analysis was conducted with R version 4.4.1. Pearson's linear correlation coefficients and the r^2 coefficient were
264 used to assess relationships between log-transformed variables within each pond and across the dataset, to identify potential
265 pond-specific and overall direct relationships between variables and GHGs. Statistical significance was determined using
266 Fisher's F test and the associated p -value. This approach was also applied to study the relationships between $\delta^{13}C-CH_4$, FOX
267 and MOX with Chl- a and TSM. To assess the impact of Chl- a concentration, macrophyte cover in summer, water depth, and
268 lake surface area on diffusive and ebullitive CH_4 fluxes, the ratio of ebullitive CH_4 to total CH_4 flux, and CO_2 and N_2O
269 fluxes, both linear and quadratic relationships were applied to log-transformed averaged data. This approach allowed for the

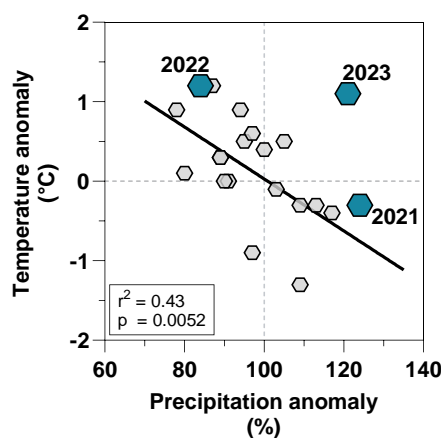
270 observation of trends between explanatory and dependent variables. For N₂O fluxes, additional explanatory variables
271 included NO₂⁻, NO₃⁻, NH₄⁺, and DIN concentrations.

272 A two-way repeated measures analysis of variance (ANOVA) was used to test for differences in categorical variables, with
273 the four seasons and the four ponds serving as independent factors, pond was set as a random effect to account for repeated
274 measurements. A one-way repeated measures ANOVA was used to test for differences in δ¹³C-CH₄ from “perturbed
275 sediments” with the four ponds serving as independent factors. After conducting an ANOVA and establishing significant
276 differences among at least two groups (p<0.05), Tukey's Honestly Significant Difference (HSD) post-hoc test was employed
277 to perform pairwise comparisons across all groups. Statistical outcomes are visually represented on boxplots, where upper-
278 and lower-case letters are used to denote significant differences (p<0.05). Different lower- and upper-case letters indicate
279 significant differences between groups.

280 3. Results and discussion

281 3.1. Seasonal variations of meteorological conditions and GHG concentrations

282 Belgium has a west coast marine climate with mild weather year-round, and evenly distributed abundant rainfall totalling on
283 average 837 mm annually for the reference period 1991-2020. The average annual air temperature was 11°C, with summer
284 average of 17.9 °C and winter average of 4.1 °C for the reference period 1991-2020. During the sampling period, from June
285 2021 to December 2023, water temperature in the surface of the four sampled ponds (Leybeek, Pêcheries, Sillex, and
286 Tenreuken; Fig. 1) tracked closely the air temperature that ranged between -1.5 and 30.0°C following the typical seasonal
287 cycle at mid-latitudes in the Northern Hemisphere (Fig. S2). Years 2022 and 2023 were about 1 °C warmer than the average
288 for the period 1991-2020 (11 °C), while year 2021 was closer to the long-term average (Fig. 2). Year 2022 was warmer and
289 drier than 2021 and 2023 (Fig. 2), with positive temperature anomalies observed evenly throughout the year (9 months out of
290 12) and negative precipitation anomalies in summer, fall and early winter (Fig. S2). Year 2021 had warmer and drier months
291 in June and September, colder and wetter months in July and August, and was overall wetter and colder than 2022 (Fig. 2).
292 Year 2023 was marked by both positive temperature and precipitation anomalies (Fig. S2), resulting in a wetter and warmer
293 year than normal and compared to 2021 and 2022. (Fig. 2). Daily wind speed was generally low (<1 m s⁻¹) except for a
294 windier period in spring 2022 (up to 5.8 m s⁻¹, corresponding to the Eunice storm) and in fall 2023 (up to 9.7 m s⁻¹,
295 corresponding to the Ciarán storm) (Fig. S2).

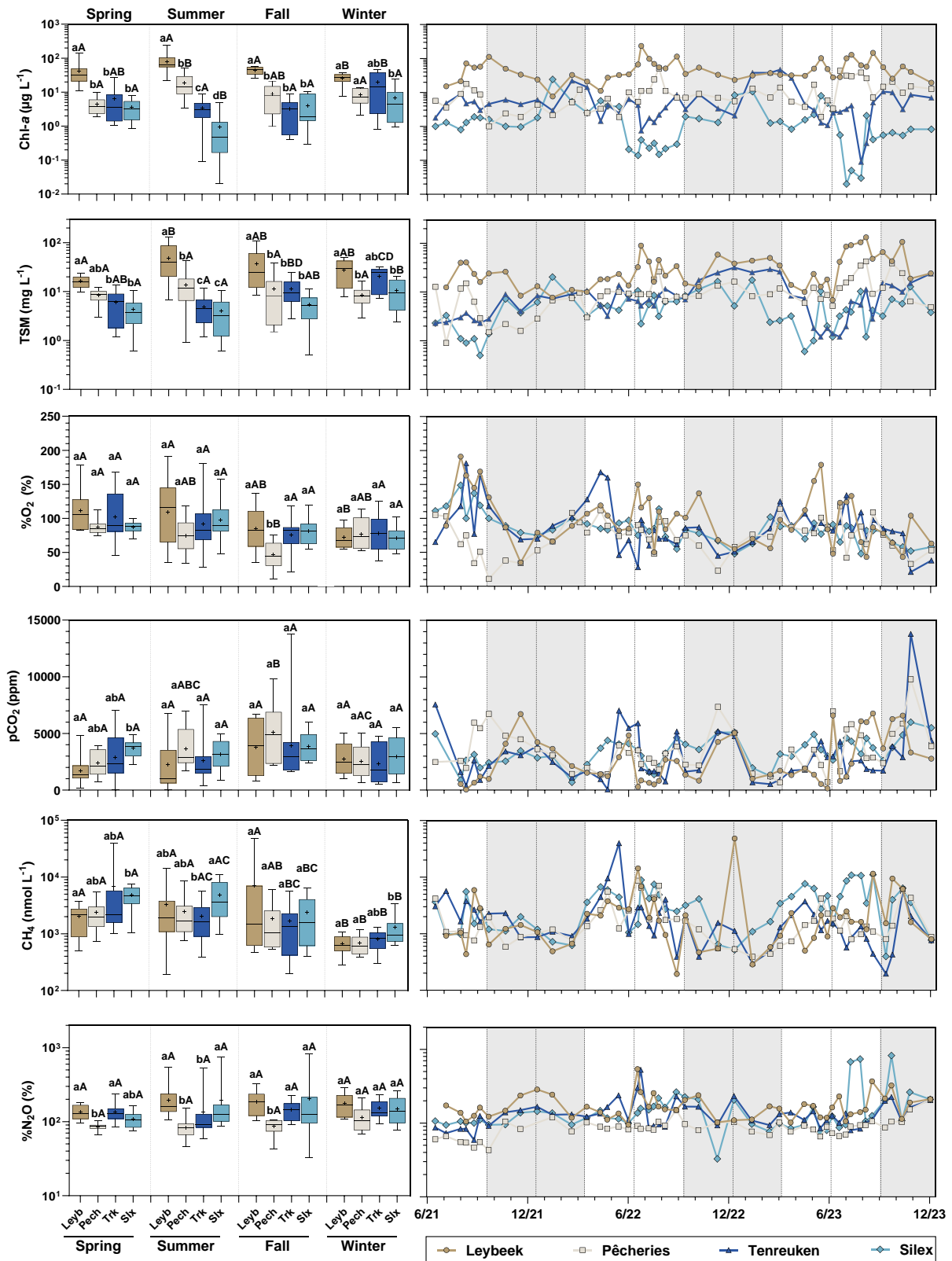


296

297 **Figure 2: Temperature anomaly (difference between the average annual temperature and the normal annual temperature for the**
298 **reference period 1991-2020 (11 °C), in °C) plotted against precipitation anomaly (ratio between annual precipitation and normal**

299 annual precipitation for the reference period 1991-2020 (837 mm), in %) from 2003 to 2023. Each small grey hexagon represents
300 values for years from 2003 to 2020 and larger blue hexagons represent the years of sampling from this study (2021, 2022 and
301 2023). Linear regression for years 2003-2020 is shown by a black solid line ($Y = 3.29 - 0.03 \cdot X$, $n=20$, Table S11). Note the
302 anomalous rainy year in 2023 relative to the pattern as function of temperature for the other years, possibly in response to the
303 strong El Niño event of 2023 (Chen et al., 2024).

304 The four sampled ponds are situated in the periphery of the city of Brussels, with the Silex pond bordered by the Sonian
305 Forest (Fig. 1). The four ponds are relatively small (0.7-3.2 ha) and shallow (60-140 cm) and have not been drained or
306 dredged since at least 2018 (Table S2). The four studied ponds had significantly different Chl-*a* concentration values during
307 summer, with the Leybeek pond having higher Chl-*a* ($78.8 \pm 49.5 \mu\text{g L}^{-1}$), followed by the Pêcherries pond ($19.1 \pm 13.7 \mu\text{g L}^{-1}$),
308 the Tenreuken pond ($3.3 \pm 2.4 \mu\text{g L}^{-1}$), and the Silex pond ($1.0 \pm 1.2 \mu\text{g L}^{-1}$) (Tukey's HSD test $p \leq 0.0001$ for each pair of
309 comparisons, Figs. 1, 3). The Leybeek and Pêcherries ponds with higher summer Chl-*a* concentration had turbid-water
310 (summer TSM = 48.7 ± 36.2 and $13.7 \pm 10.7 \text{ mg L}^{-1}$, respectively), and undetectable submerged macrophyte cover in summer
311 (Fig. 1, Table S1). The Tenreuken and Silex ponds with lower summer Chl-*a* concentrations had clear-water (summer TSM
312 = 4.9 ± 3.2 and $4.0 \pm 3.2 \text{ mg L}^{-1}$, respectively), and a high total macrophyte cover during summer (68 and 100%, respectively,
313 Fig. 1, Table S1). Values of Chl-*a* were higher in summer than in winter in the turbid-water Leybeek and Pêcherries ponds
314 (Tukey's HSD test $p=0.0107$ for the Leybeek pond, $p=0.0211$ for the Pêcherries pond) related to summer algal blooms.
315 Values of Chl-*a* were higher in winter than in summer in the clear-water Tenreuken and Silex ponds (Tukey's HSD
316 test= 0.0296 for the Tenreuken pond, $p=0.0056$ for the Silex pond), probably related to competition for inorganic nutrients
317 from macrophytes, with the Silex pond showing lower summer Chl-*a* (Tukey's HSD test $p<0.0001$), lower summer TSM
318 concentrations (Tukey's HSD test $p<0.0001$) and higher summer total macrophyte cover compared to the Tenreuken pond
319 (Fig. 1).



320

321 **Figure 3: Seasonal variations of Chlorophyll-a (Chl-a, in $\mu\text{g L}^{-1}$), total suspended matter (TSM, in mg L^{-1}), oxygen saturation**
 322 **(%O₂, in %), partial pressure of CO₂ (pCO₂ in ppm), dissolved CH₄ concentration (CH₄, in nmol L^{-1}), and N₂O saturation level**
 323 **(%N₂O, in %) in four urban ponds (Leybeek (Leyb), Pêcherries (Pech), Tenreuken (Trk), and Silex (Slx)) in the city of Brussels**
 324 **(Belgium) from June 2021 to December 2023. Box plots show median (horizontal line), mean (cross), and 25–75% percentiles (box**
 325 **limits). Whiskers extend from minimum to maximum values. White and grey bands in the graphs on the right correspond to the**
 326 **autumn/winter and spring/summer periods, respectively, and dotted vertical bars represent the first days of each season. ANOVA**
 327 **results of the multiple comparison between boxplots are summarized in Tables S4 and S5. Different lower-case letters indicate**
 328 **significant differences between ponds within a season and different upper-case letters indicate significant differences between**
 329 **seasons for a given pond.**

330 The %O₂ values ranged from 11 to 191% (Fig. 3). The highest %O₂ values in the four ponds were observed in spring and
331 summer compared to fall and winter owing to aquatic primary production. In summer, %O₂ was statically higher in the
332 Leybeek pond (109±46 %) characterized by higher Chl-*a* concentration compared to the Pêcherries pond (75±23 %) (Tukey's
333 HSD test p=0.0037). The lowest average %O₂ was observed in fall in the Pêcherries pond (46±22 %) and was statistically
334 lower than in the Leybeek (85±34%, Tukey's HSD test p=0.0302), Tenreuken (76±26 %, Tukey's HSD test p=0.0488), and
335 Silex ponds (81±19 %, Tukey's HSD test p=0.0132).

336 The pCO₂ values ranged from 40 to 13,804 ppm (Fig. 3), within the range of values typically observed in ponds (Holgerson
337 and Raymond, 2016; Peacock et al., 2019; Audet et al., 2020). Undersaturation of CO₂ with respect to atmospheric
338 equilibrium was only observed on five occasions out of the 187 measurements, three times in the turbid-water Leybeek pond
339 in summer (40 ppm on 13 August 2021, 220 ppm on 27 June 2022 and 149 ppm on 13 June 2023), and twice in the clear-
340 water Tenreuken pond in spring and summer (383 ppm on 13 August 2021 and 55 ppm on 2 May 2022). Low values of
341 pCO₂ were generally observed in spring and summer probably due to uptake of CO₂ by primary production from either
342 phytoplankton or submerged macrophytes. High values of pCO₂ were observed in fall in the four ponds and probably reflect
343 the release of CO₂ from degradation of organic matter due to the senescence of phytoplankton or macrophytes (Fig. 3). A
344 general control of pCO₂ by biological activity (primary production and respiration) was confirmed by the strong negative
345 correlation with %O₂ observed in all four ponds (*e.g.* Holgerson, 2015), as well as a positive correlation with DIN observed
346 in three ponds, and a positive correlation with SRP observed in the two clear-water ponds (Table S3; Figs S3, S4, S5, S6). A
347 negative correlation between pCO₂ and Chl-*a* was only observed in the turbid-water Leybeek pond (Table S3; Fig S5),
348 which showed the highest average Chl-*a* concentration, and no correlation was found in clear-water ponds, where aquatic
349 primary production was presumably mainly related to submerged macrophytes (Table S3; Figs S3, S4). In all four ponds,
350 pCO₂ strongly correlated positively to precipitation (Table S3; Figs S3, S4, S5, S6) suggesting a control of external inputs of
351 carbon either as organic carbon sustaining internal degradation of organic matter or as soil CO₂ (*e.g.* Marotta et al., 2011).

352 The CH₄ dissolved concentrations ranged from 194 to 48,380 nmol L⁻¹ (Fig. 3), within the range of values typically observed
353 in ponds (Holgerson and Raymond, 2016; Peacock et al., 2019; Audet et al., 2020). Dissolved CH₄ concentration was
354 positively correlated to water temperature in all four ponds (Table S3; Figs S3,S4,S5,S6), most probably reflecting the
355 increase of sedimentary methanogenesis with temperature (Schulz and Conrad, 1996), with higher dissolved CH₄
356 concentrations observed in spring (3160±5989 nmol L⁻¹) and summer (3979±2993 nmol L⁻¹) than in fall (2645±7315 nmol L⁻¹)
357 and winter (868±601 nmol L⁻¹) (Tukey's HSD test: spring versus fall, p=0.0954; spring versus winter, p<0.0001; summer
358 versus fall, p=0.0154; summer versus winter, p<0.0001). In individual ponds, dissolved CH₄ concentration was negatively
359 correlated to precipitation and DIN in the Pêcherries pond (Table S3; Fig S6), and positively correlated to SRP in the Silex
360 pond (Table S3; Fig S4). These relationships between CH₄ and other variables probably indirectly reflect the seasonal
361 variations of these other variables that showed correlations with temperature, as DIN was negatively correlated to
362 temperature in the Pêcherries pond (r²=0.11, p=0.0028), and SRP was positively correlated to temperature in the Silex pond
363 (r²= 0.10, p=0.0103). Dissolved CH₄ concentration was negatively correlated to Chl-*a* in the Silex pond (Table S3; Fig S4)
364 and to TSM in the Tenreuken pond (Table S3; Fig S3). These relationships probably reflect the negative relationship
365 between Chl-*a* and temperature in the Silex pond (r²=0.13, p=0.0008) and the negative relationship between TSM and
366 temperature in the Tenreuken pond (r²=0.36, p<0.0001) because of the primary production from macrophytes peaks in
367 summer in the two clear-water ponds.

368 The correlations between pCO₂ and precipitation, and between dissolved CH₄ concentration and temperature observed in all
369 four ponds individually were also observed when pooling together the data for all four ponds (“All” in Table S3; Fig S7).
370 The slopes of these correlations were not significantly different between ponds and were not correlated with surface area,
371 depth, or dominance of type of primary producers (phytoplankton or macrophyte) (Table S6). These results suggest that the
372 effect of precipitation on pCO₂ and the impact of temperature on dissolved CH₄ concentration outweigh other factors in
373 explaining seasonal variations.

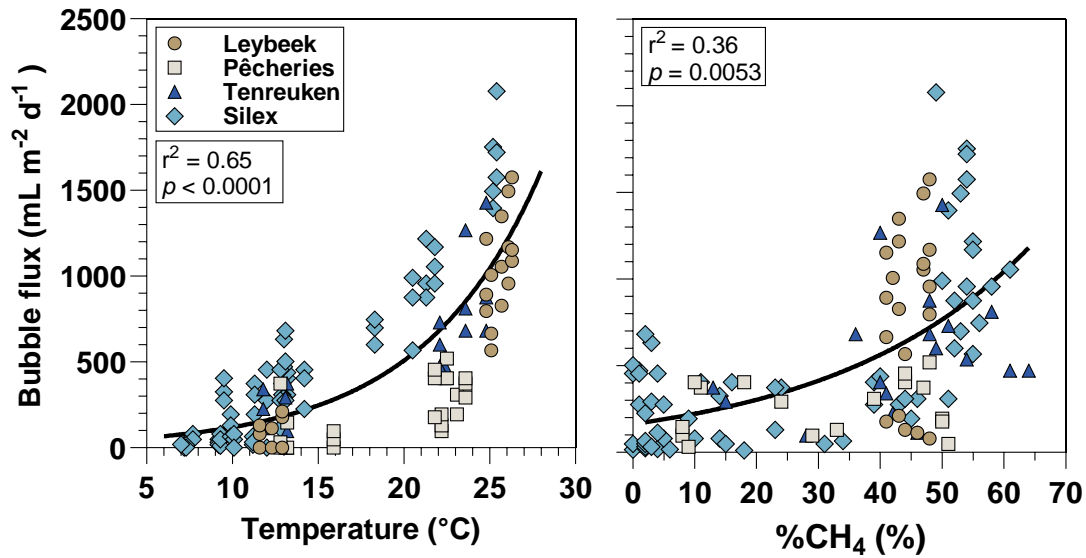
374 The %N₂O values ranged from 32 to 826% (Fig. 3), within the range of values observed in other ponds (Audet et al., 2020;
375 Rabaey and Cotner, 2022). The %N₂O values did not show significant seasonal variations in any of the four sampled ponds
376 (ANOVA F(3,174)=1,127, p=0.4091) (Fig. 3). In individual ponds, %N₂O correlated negatively to temperature in the
377 Tenreuken pond and Chl-*a* in the Silex pond, and positively to SRP in the Silex pond and TSM concentration in the
378 Tenreuken pond (Table S3; Fig S3, S4). We do not have a clear explanation for these correlations that might be spurious.
379 The correlations with Chl-*a* and TSM were surprising since they were observed in the two clear-water ponds and might
380 indirectly reflect seasonal variations (with minimal values of these two quantities in summer). More surprisingly, %N₂O was
381 not correlated with DIN (Table S3; Fig S3, S4, S5, S6) nor with individual forms of DIN (NH₄⁺, NO₂⁻, NO₃⁻) in the four
382 ponds individually or when all the data were pooled together for the individual forms of DIN (Table S3; Fig S7). In a
383 previous study of the variation of GHGs in 22 urban ponds in the city of Brussels sampled only once during each season,
384 %N₂O correlated positively with DIN, NH₄⁺, NO₂⁻, and NO₃⁻. The range of variation of DIN and %N₂O across these 22
385 ponds (2 to 625 μmol L⁻¹ for DIN, and 0 to 10,354% for %N₂O) was wider than the one observed in the present study of only
386 four ponds (1 to 135 μmol L⁻¹ for DIN, and 32 to 826% for %N₂O) (Fig. S8). The four ponds studied here are located at the
387 periphery of the city and most probably receive less atmospheric nitrogen deposition than closer to the city center. A lower
388 atmospheric nitrogen deposition in the periphery than in the city center is consistent with the correlation between %N₂O and
389 atmospheric nitrogen dioxide (NO₂), and the correlation between %N₂O and the distance from the city center (Fig. S8).
390 Atmospheric nitrogen deposition has been shown to enhance denitrification and N₂O production in lakes (McCrackin and
391 Elser, 2010; Palacin-Lizarbe et al., 2020).

392 The relationships between GHG dissolved concentrations and other variables were similar in clear-water macrophyte-
393 dominated ponds and turbid-water phytoplankton-dominated ponds. pCO₂ was positively correlated with precipitation, and
394 dissolved CH₄ concentration was positively correlated with temperature, while no significant correlation was found between
395 %N₂O and other variables in the four ponds taken individually. The negative correlation between pCO₂ and %O₂ reflected
396 the photosynthesis-respiration balance independently from the community driving aquatic primary production (macrophytes
397 in clear-water ponds and phytoplankton in turbid-water ponds).

398 **3.2. Drivers of bubble flux**

399 The bubble flux measured with inverted funnels in the four sampled ponds in the city of Brussels ranged between 0 and 2078
400 ml m⁻² d⁻¹ and strongly increased with water temperature (Fig. 4) and were overall higher in summer (837±434 mL m⁻² d⁻¹)
401 than in spring (198±170 mL m⁻² d⁻¹) and fall (106±63 mL m⁻² d⁻¹) (Tukey’s HSD test p<0.0001 for summer versus spring and
402 summer versus fall). The bubble flux values in the four sampled ponds in the city of Brussels were within the range of values
403 reported in lentic systems of equivalent size by Wik et al. (2013) (0 to 2772 mL m⁻² d⁻¹), Delsontro et al. (2016) (11 to 748
404 mL m⁻² d⁻¹) and Ray and Holgerson (2023) (0 to 2079 mL m⁻² d⁻¹). The mean CH₄ content of the bubbles in the four sampled
405 ponds in the city of Brussels was 31±21%, and comparable to the values obtained by Wik et al. (2013) (35±25%), Delsontro
406 et al. (2016) (58±25%) and Ray and Holgerson (2023) (25±13%) in lentic systems of equivalent size. The CH₄ content of the

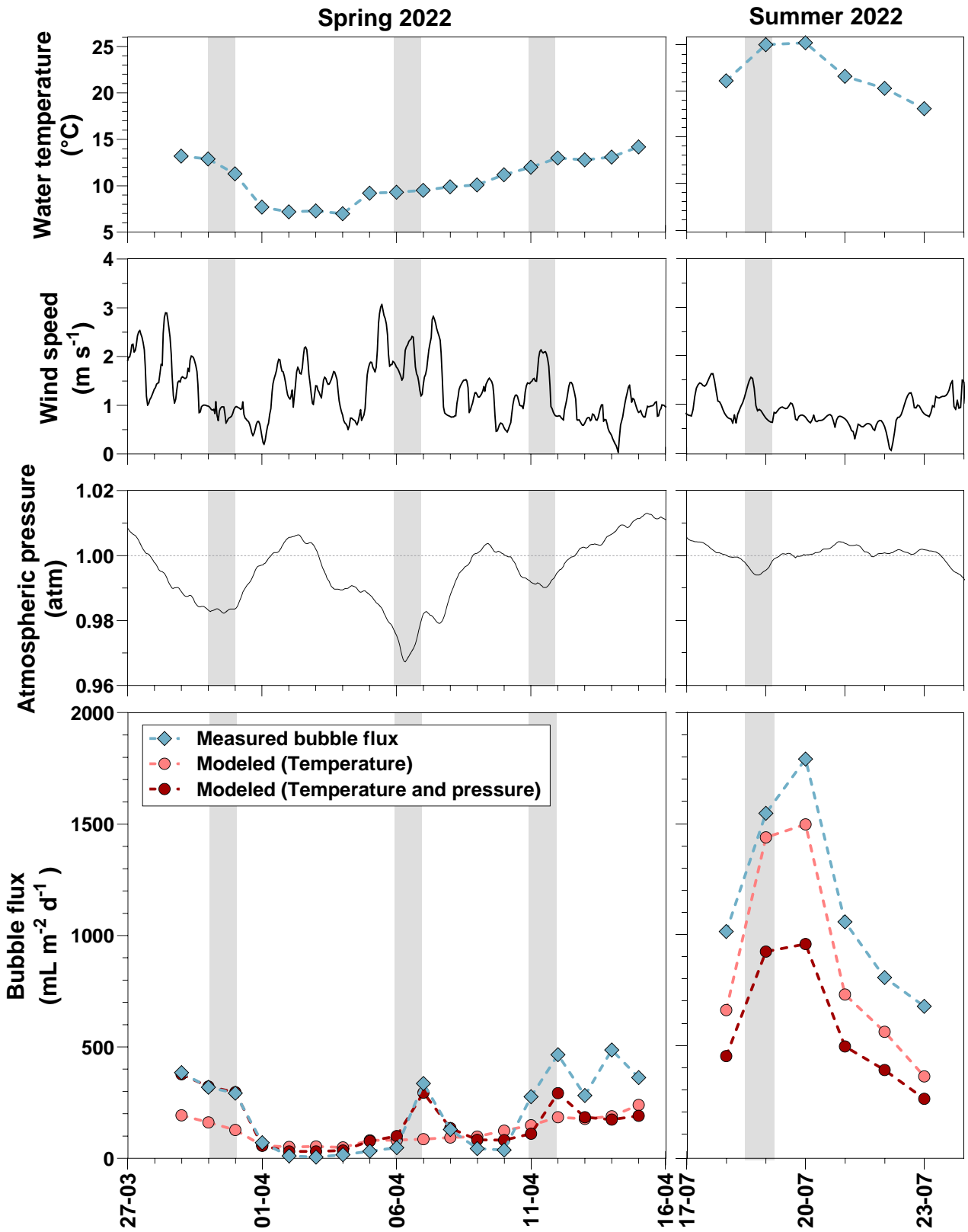
407 bubbles increased with bubble flux (Fig. 4). These patterns between bubble flux and temperature and %CH₄ were most
 408 probably related to the strong dependence of methanogenesis on temperature (Schulz and Conrad, 1996). The increase of
 409 methanogenesis with temperature leads to the build-up of gas bubbles in sediments that are richer in CH₄, and consequently
 410 to higher bubble fluxes with a higher CH₄ content at higher temperatures.



411

412 **Figure 4: Bubble flux (ml m⁻² d⁻¹) as a function of water temperature (°C) and the relative CH₄ content in bubbles (%CH₄, in %)**
 413 **in four urban ponds (Leybeek, Pêcherries, Tenreuken, and Silex) in the city of Brussels (Belgium). Bubbles fluxes were measured**
 414 **with three bubble traps in spring, summer, and fall of 2022 and 2023, totalling 8 days in the Leybeek, Pêcherries, and Tenreuken**
 415 **ponds and 24 days in the Silex pond. Given the shallowness of the sampled systems (<1.5 m, Fig. 1) we assume that sediments**
 416 **experience the same temperature as surface waters. Solid lines represent exponential regression fit of bubble flux as function of**
 417 **temperature ($Y = 28 \cdot e^{0.14 \cdot X}$, n=139), and as function of relative CH₄ content in the bubbles ($Y = 164 \cdot e^{0.03 \cdot X}$, n=123) (Table**
 418 **S11).**

419 Bubbling events are known to also be triggered by a decrease of hydrostatic pressure on the sediments due to water level
 420 fluctuations or changes in atmospheric pressure. Drops in atmospheric pressure have been documented to trigger bubble
 421 fluxes from lake sediments (Tokida et al., 2007; Scandella et al., 2011; Varadharajan and Hemond, 2012; Wik et al., 2013;
 422 Taoka et al., 2020; Zhao et al., 2021). The bubble fluxes were measured during more lengthy series at the Silex pond than
 423 the other three ponds for logistical reasons allowing investigating in more the detail the effects of temperature and
 424 atmospheric pressure variations on bubble fluxes. In spring 2022, the bubble flux at the Silex pond increased during events
 425 of drops in atmospheric pressure (depressions) (Fig. 5). There was no relation between wind speed and peaks of bubble flux
 426 ($r^2 = 0.01$, $p=0.4629$) as shown in Gatun Lake (Keller and Stallard, 1994), suggesting a more important role of changes of
 427 atmospheric pressure than wind speed in the Silex pond in spring 2022. The bubble flux at the Silex pond was higher in
 428 summer (1152 ± 433 mL m⁻² d⁻¹) than during spring (198 ± 170 mL m⁻² d⁻¹) (Tukey's HSD test $p < 0.0001$), and the temporal
 429 changes tracked those of water temperature. The bubble flux was modelled as function of temperature alone or as function of
 430 both temperature and pressure changes (Figs. 5, S9). The inclusion of the term of pressure drops in addition to temperature
 431 improved the performance of the model compared to the original data, for periods of low temperature (<15°C) but not for
 432 warmer periods (>15°C) (Figs. 5, S9) when bubbling fluxes were quantitatively more important. The inclusion of the term of
 433 pressure changes only improved the performance of the model compared to the original data very marginally when
 434 comparing the full temperature range (<15°C and >15°C) (Fig. S9), showing that the intensity of bubble flux was mainly
 435 driven by temperature change at yearly scales, in agreement with previous studies (e.g. Wik et al., 2013; DelSontro et al.,
 436 2016; Aben et al., 2017; Ray and Holgerson, 2023).

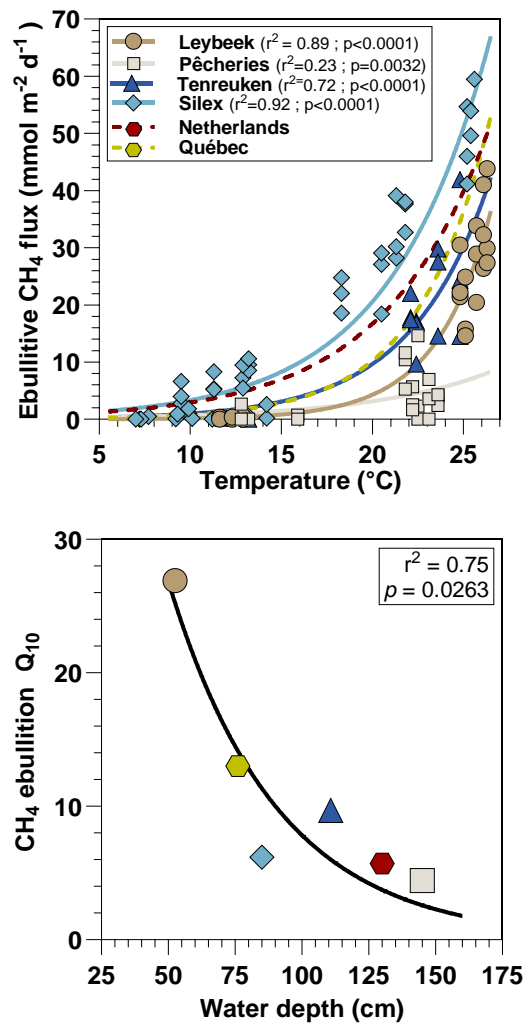


437

438 Figure 5: Water temperature (°C), wind speed (m s⁻¹), atmospheric pressure (atm), and measured and modeled bubble flux (ml m⁻²
 439 d⁻¹) in the Silex pond from the 29 March 2022 to the 15 April 2022 and from the 18 July 2022 to the 23 July 2022. The bubble flux
 440 was modelled from a fit to data based on temperature alone and from both temperature and drops in atmospheric pressure.

441 **3.3. Drivers of methane ebullitive fluxes**

442 Ebullitive CH₄ fluxes in the four ponds ranged between 0 and 59 mmol m⁻² d⁻¹, within the range reported in lentic systems
 443 (e.g. Deemer and Holgerson, 2021) and were positively related to temperature (Fig. 6) as shown previously in other small
 444 lentic systems (e.g. Wik et al., 2013; DelSontro et al., 2016; Aben et al., 2017; Ray and Holgerson, 2023; Rabaey and
 445 Cotner, 2024). The fitted relations between ebullitive CH₄ fluxes and temperature were specific to each pond and
 446 encompassed the fitted relations established in similar systems: four small ponds in Québec (DelSontro et al., 2016) and a
 447 small urban pond in the Netherlands (Aben et al., 2017). The Q₁₀ of CH₄ ebullition values ranged between 4.4 in the deeper
 448 Pêcherries pond and 26.9 in the shallower Leybeek pond, respectively (Table S7). The Q₁₀ of CH₄ ebullition in the four
 449 studied ponds of the city of Brussels, in Québec (DelSontro et al., 2016), and in the Netherlands (Aben et al., 2017) were
 450 negatively related to water depth (Fig. 6). An increase in water temperature leads to a smaller increase in CH₄ ebullitive
 451 fluxes (lower Q₁₀) in deeper ponds as the impact of hydrostatic pressure on sediments is higher in deeper ponds compared to
 452 shallow ponds, restricting bubble formation and release (e.g. DelSontro et al., 2016).

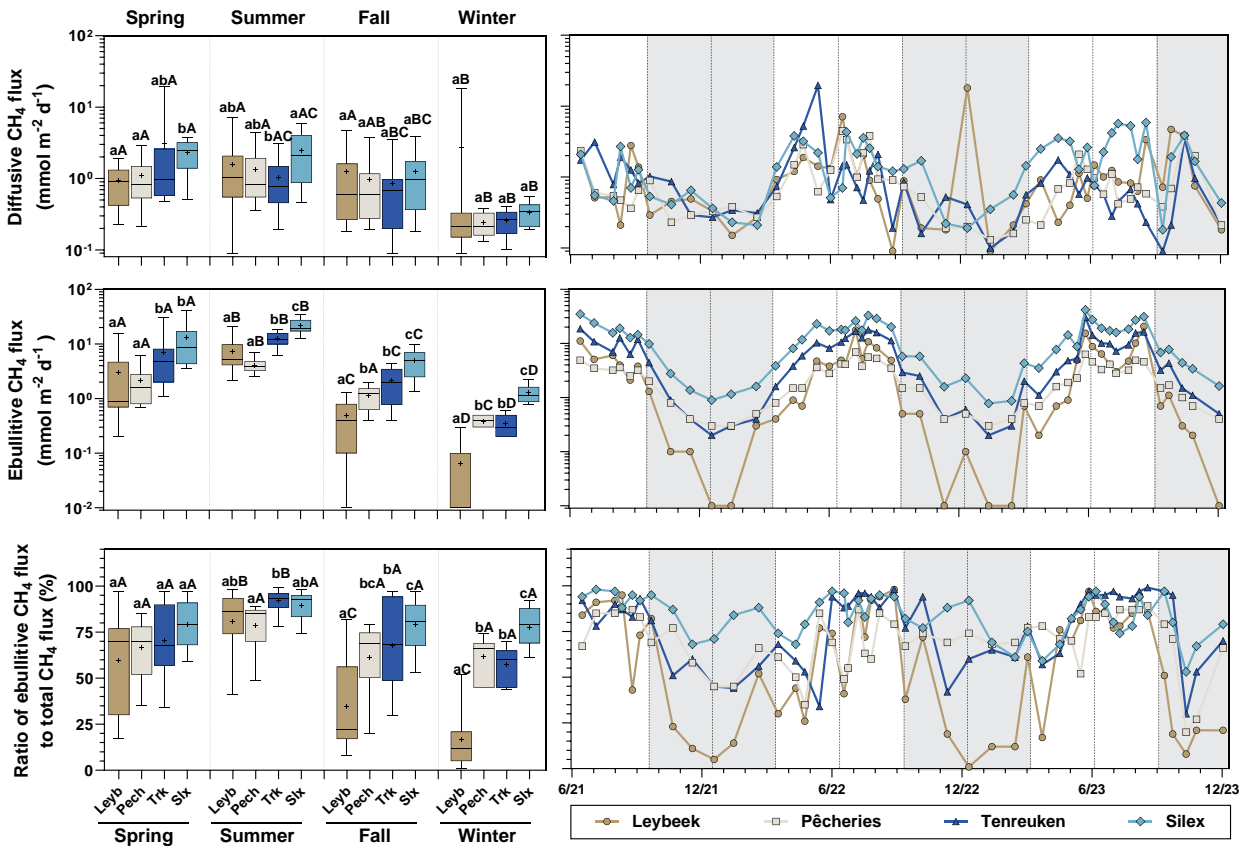


453

454 **Figure 6:** Measured ebullitive CH₄ fluxes (mmol m⁻² d⁻¹) as function of water temperature (°C) in four urban ponds (Leybeek,
 455 Pêcherries, Tenreuken, and Silex) in the city of Brussels (Belgium), in spring, summer, and fall of 2022 and 2023, totalling 8 days in
 456 the Leybeek, Pêcherries, and Tenreuken ponds and 24 days in the Silex pond, with three bubble traps. Solid lines represent
 457 exponential fit for the Leybeek ($Y = 0.01 \cdot e^{0.32 \cdot X}$, n=22), Pêcherries ($Y = 0.16 \cdot e^{0.15 \cdot X}$, n=22), Tenreuken ($Y = 0.10 \cdot e^{0.23 \cdot X}$,
 458 n=19), Silex ($Y = 0.54 \cdot e^{0.18 \cdot X}$, n=72) ponds (Table S7) dashed lines represent exponential fit established in similar systems: four
 459 small ponds in Québec ($Y = 0.06 \cdot e^{0.25 \cdot X}$) (DelSontro et al., 2016) and a small urban pond in the Netherlands ($Y = 0.51 \cdot$
 460 $e^{0.17 \cdot X}$) (Aben et al., 2017). Each exponential curve allows to determine a Q₁₀ of CH₄ ebullition, plotted against water depth; solid
 461 line represents exponential regression fit ($Y = 92 \cdot e^{-0.02 \cdot X}$, n = 6) (Table S11).

462 **3.4. Relative contribution of methane ebullitive and diffusive fluxes**

463 Diffusive CH₄ fluxes computed from dissolved CH₄ concentration and *k* derived from wind speed ranged between 0.1 and
 464 19.7 mmol m⁻² d⁻¹ (Fig. 7) within the range reported in lentic systems (*e.g.* Deemer and Holgerson, 2021). The diffusive CH₄
 465 fluxes tended to be higher in summer and spring than in fall and winter owing to the strong positive dependency between
 466 CH₄ and water temperature (Fig. 3; Table S3). In addition, wind speed only showed small seasonal variations during
 467 sampling (0.6±0.6 m s⁻¹ in spring, 0.3±0.2 m s⁻¹ in summer, 0.7±0.7 m s⁻¹ in fall, and 0.6±0.2 m s⁻¹ in winter) (Fig. 3).
 468 Ebullitive CH₄ fluxes were calculated from the relations with temperature for each pond given in Figure 6 from the
 469 temperature data coincident with the diffusive CH₄ fluxes (Fig. 7). The resulting calculated ebullitive CH₄ fluxes allowed to
 470 compare and integrate seasonally both components of CH₄ emissions to the atmosphere, and to calculate the relative
 471 contribution of ebullition to total (diffusive+ebullitive) CH₄ emissions.



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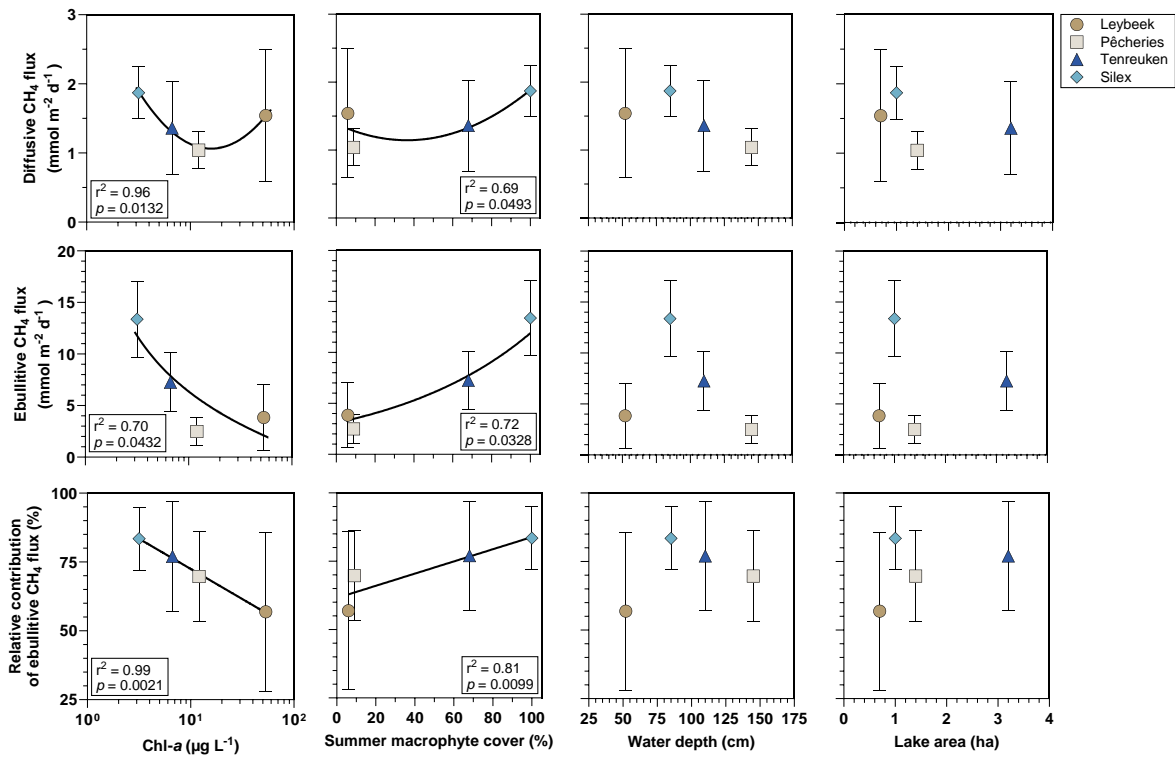
473 **Figure 7: Seasonal variations of diffusive and ebullitive CH₄ fluxes (mmol m⁻² d⁻¹), and the ratio of ebullitive CH₄ flux to total**
 474 **(ebullitive+diffusive) CH₄ flux (%) in four urban ponds (Leybeek (Leyb), Pêcheries (Pech), Tenreuken (Trk), and Silex (Slx)) in the**
 475 **city of Brussels (Belgium) from June 2021 to December 2023. Diffusive fluxes were calculated from CH₄ concentration and gas**
 476 **transfer velocity derived from wind speed. Ebullitive CH₄ fluxes were calculated from the relations with temperature for each**
 477 **pond (Fig. 6; Table S7) from the temperature data coincident with the diffusive CH₄ fluxes. Box plots show median (horizontal**
 478 **line), mean (cross), and 25–75% percentiles (box limits). Whiskers extend from minimum to maximum values. White and grey**
 479 **bands in the graphs on the right correspond to the autumn/winter and spring/summer periods, respectively, and dotted vertical**
 480 **bars represent the first days of each season. ANOVA results of the multiple comparison between boxplots are summarized in**
 481 **Tables S4 and S5. Different lower-case letters indicate significant differences between ponds within a season and different upper-**
 482 **case letters indicate significant differences between seasons for a given pond.**

483 The relative contribution of ebullition to total CH₄ emissions ranged between 1 and 99% in the four sampled ponds in the
 484 city of Brussels (Fig. 7), within the range reported in lentic systems (*e.g.* Deemer and Holgerson, 2021). Owing to the strong
 485 dependency of ebullitive CH₄ fluxes to temperature (Table S7; Fig. 6), the mean relative contribution of ebullition to total
 486 CH₄ emissions for all data pooled together was higher in summer (85±7 %) compared to spring (69±14 %, Tukey's HSD test

487 $p=0.0104$), fall ($61\pm 18\%$, Tukey's HSD test $p<0.0001$), and winter ($53\pm 8\%$, Tukey's HSD test $p<0.0001$) (Fig. 7). This
488 finding is consistent with other studies showing that ebullitive CH_4 fluxes can account for more than half of total CH_4
489 emissions in small and shallow lentic systems (*e.g.* Wik et al., 2013; Deemer and Holgerson, 2021; Ray and Holgerson,
490 2023; Rabaey and Cotner, 2024). The relative contribution of ebullition to total CH_4 emissions was lowest during the other
491 seasons, especially in the Leybeek pond (Fig. 7). Owing to the strong dependency of ebullitive CH_4 fluxes to temperature,
492 the relative contribution of ebullition to total CH_4 emissions was related to temperature in the four ponds (Fig. S10), as
493 previously also shown in Québec ponds (DelSontro et al., 2016).

494 The values of Q_{10} of diffusive CH_4 fluxes were lower than those for ebullitive CH_4 fluxes, less variable (1.2 in the Pêcherries
495 pond to 2.9 in the Silex pond), and less statistically significant (Table S7). Other studies have also reported higher Q_{10} for
496 CH_4 ebullition than for CH_4 diffusion in lentic systems (DelSontro et al., 2016; Xun et al., 2024). The lower dependence to
497 temperature of CH_4 diffusion compared to CH_4 ebullition might be related to a lower relative change of CH_4 concentrations
498 and k to temperature change. CH_4 concentrations in surface water are very strongly affected by MOX (see hereafter). A
499 relative increase of CH_4 production in sediments by methanogenesis will lead to a stronger increase of CH_4 emission by
500 ebullition than by diffusion because of a mitigation by MOX on CH_4 diffusion. Additionally, k depends on wind speed, but
501 the warmer periods of the year (summer) tended to be less windy ($\sim 0.3\text{ m s}^{-1}$) than the other seasons ($>0.6\text{ m s}^{-1}$) also
502 contributing to lower dependence on temperature of CH_4 diffusion compared to ebullition and lower Q_{10} values.

503 The annual averaged diffusive and ebullitive fluxes of CH_4 in the four ponds in the city of Brussels were plotted against Chl-
504 a concentration, total macrophyte cover in summer, water depth, and lake surface area (Fig. 8) that are frequent predictors of
505 variations of CH_4 fluxes among lakes (Holgerson and Raymond, 2016; DelSontro et al., 2018, Deemer and Holgerson, 2021;
506 Casas-Ruiz et al., 2021; Borges et al., 2022). The annual diffusive CH_4 flux was significantly lower in the slightly deeper
507 Pêcherries pond (130 cm depth) than the two slightly shallower ponds (Leybeek (60 cm depth) and Silex (110 cm depth)
508 ponds) (Tukey's HSD test $p=0.0007$ for Pêcherries versus Leybeek, $p<0.0001$ for Pêcherries versus Silex), and the annual
509 ebullitive CH_4 flux was significantly lower in the Pêcherries pond than the Silex pond (Tukey's HSD test $p<0.0001$) but was
510 not significantly different than the Leybeek pond (Tukey's HSD test $p=0.3847$). No other significant differences in annual
511 diffusive and ebullitive CH_4 fluxes related to water depth or surface area were observed. The narrow range of variation of
512 water depth (50 to 150 cm) and surface area (0.7 to 3.2 ha) could explain the lack of a clear decrease of diffusive and
513 ebullitive CH_4 fluxes with increasing depth or surface that are frequent predictors of variations of CH_4 fluxes among ponds
514 (*e.g.* Holgerson, 2015; Holgerson and Raymond, 2016; Ray et al., 2023; Theus et al., 2023) and lakes (*e.g.* Kankaala et al.,
515 2013; DelSontro et al., 2018, Deemer and Holgerson, 2021; Casas-Ruiz et al., 2021; Borges et al., 2022). Correlations
516 between CH_4 fluxes and depth or lake surface area have been shown among lakes across much larger ranges of variation of
517 lake depth (Borges et al., 2022) and surface area (Kankaala et al., 2013; Holgerson and Raymond, 2016; Casas-Ruiz et al.,
518 2021).



519

520 **Figure 8:** Mean diffusive and ebullitive CH₄ fluxes (mmol m⁻² d⁻¹) and mean ratio of ebullitive CH₄ flux to total
 521 (diffusive+ebullitive) CH₄ flux (%) versus chlorophyll-*a* (Chl-*a*, in µg L⁻¹), total macrophyte cover in summer (%), water depth
 522 (cm), and lake surface area (ha) in four ponds (Leybeek, Pêcherries, Tenreuken, and Silex) in the city of Brussels (Belgium) from
 523 June 2021 to December 2023. Error bars indicate the standard deviation. Dashed lines indicate trends in relationship between
 524 variables (Table S11).

525 The annual ebullitive CH₄ fluxes were higher in the two clear-water ponds (7.3±2.9 and 13.4±3.7 m⁻² d⁻¹ in the Tenreuken
 526 and Silex ponds, respectively) than the two turbid-water ponds (3.8±3.2 and 2.5±1.4 m⁻² d⁻¹ in the Leybeek and Pêcherries
 527 ponds, respectively) (Tukey's HSD test p<0.0001 for each comparison between a clear-water pond and a turbid-water pond).
 528 The annual ebullitive CH₄ fluxes were significantly higher in the Silex pond than the Tenreuken pond (Tukey's HSD test
 529 p<0.0001) that showed a higher macrophyte cover during summer (100% in the Silex pond and 68% in the Tenreuken pond)
 530 and were not significantly different in the two turbid-water Leybeek and Pêcherries ponds (Tukey's HSD test p=0.3847) that
 531 showed similar macrophyte cover during summer (6 and 9% in the Leybeek and Pêcherries ponds, respectively) (Fig. 8). The
 532 annual ebullitive CH₄ fluxes were overall positively correlated to macrophyte cover and negatively correlated to Chl-*a* (Fig.
 533 8). The higher ebullitive CH₄ emissions from the clear-water ponds would suggest that the delivery of organic matter to
 534 sediments from macrophytes sustained a larger methane production than from phytoplankton. This finding is consistent with
 535 the notion that vegetated littoral zones of lakes are hot spots of CH₄ production and emission (e.g. Hyvönen et al., 1998;
 536 Huttunen et al., 2003; Juutinen et al., 2003; Desrosiers et al., 2022). In other small lentic systems, the CH₄ dissolved
 537 concentrations and diffusive fluxes have also been shown to correlate positively with macrophyte cover (e.g. Ray et al.,
 538 2023; Theus et al., 2023).

539 The annual diffusive CH₄ flux was higher in the two clear-water ponds (1.4±0.7 and 1.9±0.4 mmol m⁻² d⁻¹ in the Tenreuken
 540 and Silex ponds, respectively) than in the turbid-water Pêcherries pond (1.0±0.3 mmol m⁻² d⁻¹) (Tukey's HSD test p=0.0404
 541 for Tenreuken versus Pêcherries, and p<0.0001 for Silex versus Pêcherries), which was consistent with the pattern of higher
 542 ebullitive CH₄ emissions from clear-water ponds (Fig. 8). In the four sampled urban ponds, annual CH₄ diffusive fluxes were
 543 significantly higher in the pond with the highest total macrophyte cover in the clear-water ponds, and significantly higher in
 544 the pond with highest Chl-*a* concentration in the turbid-water ponds (Fig. 8). An increase in methane production with

545 phytoplankton biomass in turbid-water ponds has also been reported by other studies in lakes (*e.g.* Yan et al., 2019;
546 Bartosiewicz et al., 2021). Since total macrophyte cover and Chl-*a* were anti-correlated, we hypothesize that the variations of
547 CH₄ diffusive fluxes follow a U-shaped relation with either Chl-*a* or macrophyte cover. Higher values of annual CH₄
548 diffusive fluxes occurred at the extreme values of Chl-*a* or of macrophyte cover (minimum or maximum), and lower values
549 occurred at the intermediate values of Chl-*a* or macrophyte cover. The relative contribution of ebullitive CH₄ fluxes to the
550 total flux was higher in the clear-water Silex pond, which had the highest macrophyte cover, compared to the two turbid-
551 water ponds with lower macrophyte cover (Tukey's HSD test $p < 0.0001$ for Silex versus Leybeek, $p = 0.0056$ for Silex versus
552 Pêcherries), and was higher in the clear-water Tenreuken pond than in the turbid-water Leybeek pond (Tukey's HSD test
553 $p < 0.0001$) (Fig. 8). The relative contribution of ebullitive CH₄ fluxes to the total CH₄ flux seems to increase concomitantly
554 with the macrophyte cover (Fig. 8), and was overall strongly positively correlated to macrophyte cover and negatively to
555 Chl-*a* (Fig. 8). These patterns are consistent with the idea of an increase of ebullition relative to diffusive CH₄ emissions in
556 vegetated sediments compared to unvegetated sediments (*e.g.* Desrosiers et al., 2022; Ray et al., 2023; Theus et al., 2023).

557 The annual diffusive and ebullitive fluxes in the four ponds in the city of Brussels were within the range of values for ponds
558 of similar surface area (0.4 to 4.0 ha) compiled by Deemer and Holgerson (2021) (Fig. S11). The linear regression of
559 ebullitive CH₄ fluxes as a function of diffusive CH₄ fluxes allows comparing the data of ebullitive CH₄ fluxes from the four
560 Brussels ponds "normalized" to the diffusive CH₄ fluxes. The ebullitive CH₄ fluxes from the two turbid-water ponds
561 (Pêcherries and Leybeek) were very close to the linear regression showing they were characterized by ebullitive CH₄ fluxes
562 equivalent to those in the ponds compiled by Deemer and Holgerson (2021) when normalized by the diffusive fluxes. The
563 ebullitive CH₄ fluxes from the two clear-water ponds (Tenreuken and Silex) were above the linear regression showing they
564 were characterized by ebullitive CH₄ fluxes above those in the ponds compiled by Deemer and Holgerson (2021) when
565 normalized by the diffusive fluxes. We hypothesize the relatively higher ebullitive fluxes in the two clear-water ponds were
566 related to enhancement of ebullition from organic matter subsidized by macrophytes. This hypothesis is consistent with the
567 two clear-water ponds in Brussels having higher ebullitive fluxes than in the ponds compiled by Deemer and Holgerson
568 (2021) at equivalent Chl-*a* values (Fig. S11). The observed high ebullitive fluxes in the clear-water ponds would suggest that
569 Chl-*a* concentration alone fails to predict ebullitive fluxes in macrophyte-dominated clear-water ponds. Consequently, global
570 scaling of CH₄ fluxes in lentic systems using Chl-*a* as a predictor as used in lakes (*e.g.* DelSontro et al., 2018) might under-
571 estimate ebullitive CH₄ emissions due to a misrepresentation of macrophyte-dominated clear-water ponds.

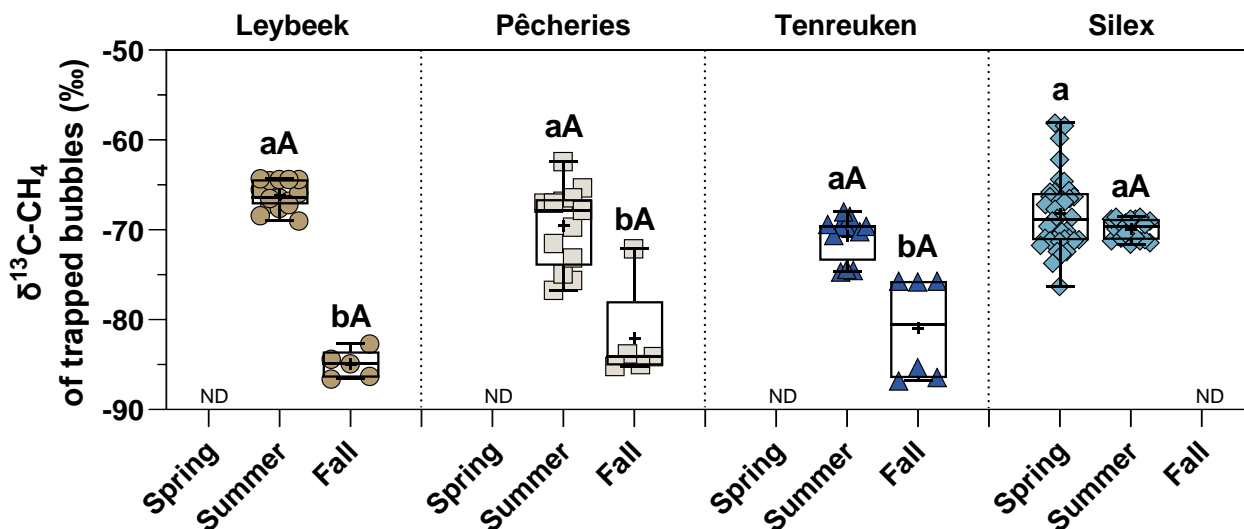
572 The annual averaged diffusive fluxes of CO₂ (F_{CO_2}) and N₂O (F_{N_2O}) in the four ponds in the city of Brussels were also plotted
573 against Chl-*a* concentration, total macrophyte cover in summer, water depth, and lake surface area, as well as DIN for N₂O
574 fluxes (Figs. S12, S13, S14). Annual F_{CO_2} did not show significant differences between the four studied ponds (Tukey's HSD
575 test: $p > 0.05$ for each comparison), and F_{CO_2} did not significantly correlate to the other variables (Chl-*a* concentration, total
576 macrophyte cover, water depth, and lake surface area). This might be surprising since other studies have reported lower CO₂
577 fluxes in more productive lentic systems (*e.g.* Sand-Jensen and Staehr 2007; Borges et al. 2022). We hypothesize that given
578 that the four systems were either phytoplankton-dominated or macrophyte-dominated, in both cases, the ponds had an
579 important submerged productivity resulting in a relatively invariant F_{CO_2} as function of either Chl-*a* or macrophyte cover.
580 Annual mean F_{CO_2} was also uncorrelated to water depth and lake area (Fig. S12). This might have resulted from the relative
581 similarity of depth and surface area of the four studied ponds, as it is well established that CO₂ emissions strongly increase
582 with decreasing size of ponds (Holgerson and Raymond, 2016). Annual F_{N_2O} was not significantly different between clear-
583 water and turbid-water ponds. F_{N_2O} was significantly lower in the slightly deeper Pêcherries pond than the two slightly
584 shallower Leybeek and Silex ponds (Fig. S13) (Tukey's HSD test $p = 0.0012$ for Pêcherries vs. Leybeek, and $p = 0.0052$ for

585 Pêcherries vs. Silex), and F_{N_2O} showed a significant negative relationship with water depth (Fig. S13). We hypothesize that
 586 this might reflect a larger dilution of N_2O diffusing from sediments in the deeper systems. F_{N_2O} did not correlate to DIN,
 587 NH_4^+ , NO_2^- , and NO_3^- (Fig. S14). We hypothesize that this reflects the rather narrow range of annual DIN average values in
 588 the four studied ponds (~24 to ~29 $\mu\text{mol L}^{-1}$), as DIN, NH_4^+ , NO_2^- , and NO_3^- were not statistically different between ponds
 589 (Tukey's HSD test $p > 0.05$ for every comparison).

590 3.5. Methanogenesis pathway inferred from $\delta^{13}\text{C-CH}_4$ in bubbles

591 $\delta^{13}\text{C-CH}_4$ was measured in bubbles trapped during the ebullition flux measurements and in bubbles collected by perturbing
 592 the sediments. The variations of $\delta^{13}\text{C-CH}_4$ suggest that there could have been variations of the relative importance of
 593 hydrogenotrophic versus acetoclastic pathways of methanogenesis among different ponds but also seasonally.
 594 Methanogenesis by the hydrogenotrophic pathway produces CH_4 with more negative $\delta^{13}\text{C-CH}_4$ values (-100‰ to -60‰)
 595 compared to the acetoclastic pathway (-65‰ to -50‰) (Whiticar et al., 1986). Yet, it remains unclear which environmental
 596 factors determine the relative importance of hydrogenotrophic and acetoclastic methanogenesis pathways (Conrad et al.,
 597 2011).

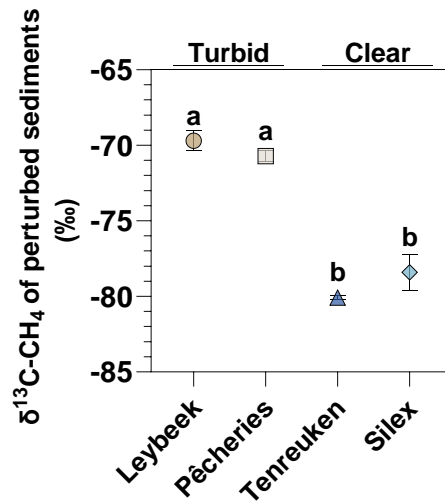
598 The $\delta^{13}\text{C-CH}_4$ values in the trapped bubbles for the all dataset were statistically more negative in fall (-83.2±5.2 ‰) than
 599 summer (-69.5±3.2 ‰) and spring (-68.2±4.4 ‰) (Fig. 9; Table S8) (Tukey's HSD test $p < 0.0001$ for fall versus summer, and
 600 fall versus spring), suggesting a dominance of hydrogenotrophic methanogenesis in fall compared to spring and summer
 601 when acetoclastic methanogenesis seemed dominant. Hydrogenotrophic methanogenesis occurs at higher temperatures than
 602 acetoclastic methanogenesis (Schulz and Conrad, 1996; Schulz et al., 1997), however, temperature in fall (11.9±3.7 °C) was
 603 lower than in summer (21.1±1.9 °C) (Tukey's HSD test $p < 0.0001$). A shift from acetoclastic methanogenesis to
 604 hydrogenotrophic methanogenesis has been documented in response to the increase of NH_4^+ concentration (Ni et al., 2022;
 605 Wang et al., 2022) and the decrease of pH (Kotsyurbenko et al., 2007) expected in response to an increase of CO_2 . An
 606 increase of NH_4^+ and decrease of pH in pore waters in fall compared to summer and spring would be consistent with the
 607 sustained benthic organic matter degradation leading to a gradual change of pore water chemistry from spring to fall.



608
 609 **Figure 9:** $^{12}\text{C}/^{13}\text{C}$ ratio of CH_4 ($\delta^{13}\text{C-CH}_4$, in ‰) in bubbles collected during ebullitive flux measurements (“trapped bubbles”) in
 610 four urban ponds (Leybeek, Pêcherries, Tenreuken, and Silex) in the city of Brussels (Belgium), measured in spring, summer, and
 611 fall in 2022 and 2023 (September 2023 and October 2023 in the Leybeek pond; July 2023 and October 2023 in the Pêcherries pond;
 612 August 2023 and October 2023 in the Tenreuken pond; April 2022 and July 2022 in the Silex pond). Box plots show median
 613 (horizontal line), mean (cross), and 25–75% percentiles (box limits). Whiskers extend from minimum to maximum values. ND = no

614 data. ANOVA results of the multiple comparison between boxplots are summarized in Table S8. Different lower-case letters
615 indicate significant differences between seasons for a given pond and upper-case letters indicate significant differences between
616 ponds within a season.

617 In summer 2023, a survey of all four ponds was made to simultaneously sample bubbles by perturbation of the sediment for
618 the determination of the $\delta^{13}\text{C-CH}_4$ in the released bubbles. The $\delta^{13}\text{C-CH}_4$ values of perturbed sediments were more negative
619 in the clear-water macrophyte-dominated ponds (-80.1 ± 0.1 ‰ and -78.4 ± 1.2 ‰ in the Tenreuken and Silex ponds,
620 respectively) than in the turbid-water phytoplankton-dominated ponds (-69.7 ± 0.7 ‰ and -70.7 ± 0.4 ‰ in the Leybeek and
621 Pêcherries ponds, respectively) (Tukey's HSD test $p < 0.0001$ for each comparison between a clear pond and a turbid pond)
622 (Fig. 10). This pattern of $\delta^{13}\text{C-CH}_4$ of perturbed sediments could suggest a higher contribution of the hydrogenotrophic
623 methanogenesis pathway compared to the acetoclastic pathway in the clear-water ponds where organic matter for
624 methanogenesis was assumed to be mainly related to macrophytes rather than phytoplankton. Based on gene expression
625 during incubations, Wang et al. (2023) suggested that acetoclastic methanogenesis pathway was stimulated by macrophyte
626 organic carbon compared to phytoplankton organic matter in lakes Chaohu and Taihu in China. The distribution of $\delta^{13}\text{C-CH}_4$
627 data in the four urban ponds of the city of Brussels suggests the opposite pattern, with macrophyte organic carbon
628 stimulating the hydrogenotrophic methanogenesis pathway. This pattern seems consistent with the more refractory nature of
629 macrophyte organic carbon compared to the more labile nature of phytoplankton organic carbon. Organic matter from
630 macrophytes has a large share of molecules difficult to degrade such as cellulose unlike organic matter from phytoplankton
631 that is rich in polysaccharides and proteins (West et al., 2015; Berberich et al., 2020). In presence of more refractory organic
632 matter, a partial fermentation would favour the production of H_2 over acetate which would favour hydrogenotrophic
633 methanogenesis over acetoclastic methanogenesis (Liu et al., 2017).



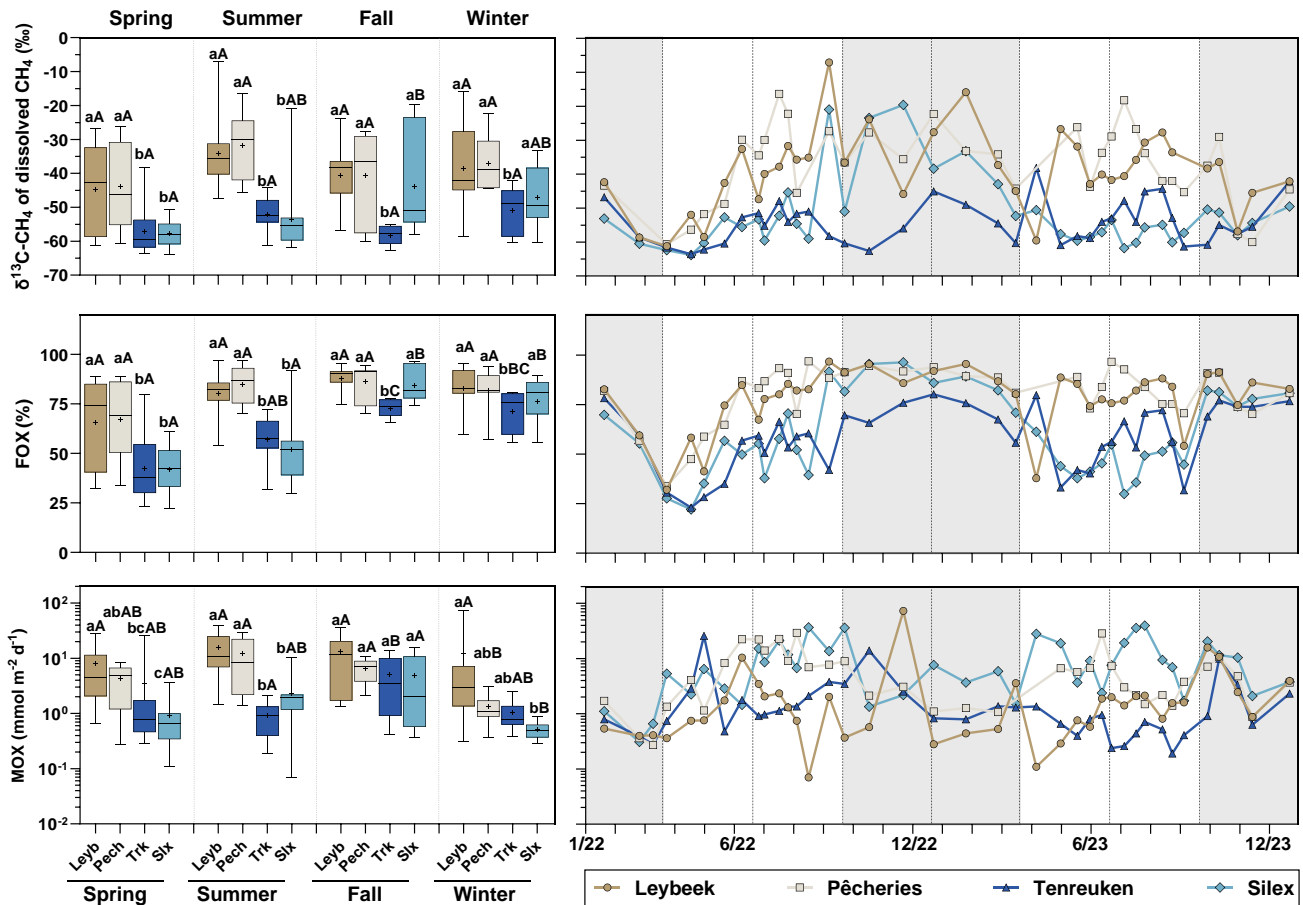
634

635 **Figure 10: Mean \pm standard deviation $^{13}\text{C}/^{12}\text{C}$ ratio of CH_4 ($\delta^{13}\text{C-CH}_4$, in ‰) in bubbles released from sediments after physical**
636 **perturbation (“perturbed sediments”) in four ponds (Leybeek, Pêcherries, Tenreuken, and Silex) in the city of Brussels (Belgium)**
637 **in summer 2023 (4th September 2023). Error bars indicate standard deviation on the mean. ANOVA results of the multiple**
638 **comparison between boxplots are summarized in Table S9. Different lower-case letters indicate significant differences between**
639 **ponds.**

640 3.6. Methane oxidation

641 The $\delta^{13}\text{C-CH}_4$ of dissolved CH_4 in surface waters in the four sampled ponds in the city of Brussels ranged between -16 and -
642 64 ‰ (Fig. 11). The $\delta^{13}\text{C-CH}_4$ of dissolved CH_4 in surface waters were generally higher than in sediments based on trapped
643 bubbles during the ebullition measurements (-55 to -87 ‰; Fig. 9). The ^{13}C enriched values of dissolved CH_4 in surface
644 waters samples probably resulted from MOX. FOX in surface waters in the four sampled ponds in the city of Brussels

645 ranged between 22 and 97%. MOX in surface waters in the four sampled ponds in the city of Brussels ranged between 0.1
 646 and 73.0 $\text{mmol m}^{-2} \text{d}^{-1}$ (Fig. 11).



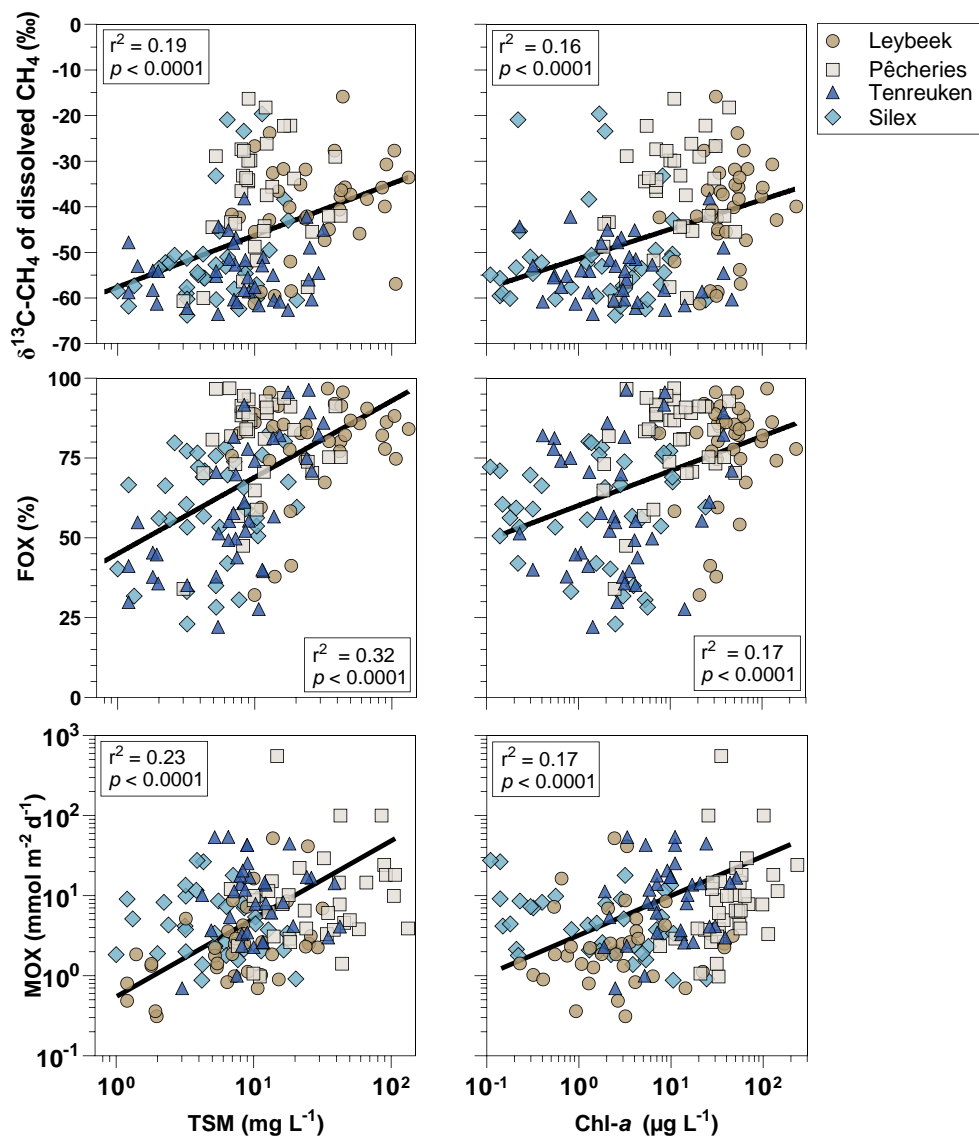
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648 **Figure 11: Seasonal variations of $^{13}\text{C}/^{12}\text{C}$ ratio of dissolved CH_4 in surface waters ($\delta^{13}\text{C}\text{-CH}_4$ of dissolved CH_4 , in ‰), fraction of**
 649 **CH_4 removed by methane oxidation (FOX, in %), and methane oxidation (MOX, in $\text{mmol m}^{-2} \text{d}^{-1}$) in four urban ponds (Leybeek**
 650 **(Leyb), Pêcherries (Pech), Tenreuken (Trk), and Silex (Slx)) in the city of Brussels (Belgium) from January 2022 to December 2023.**
 651 **Box plots show median (horizontal line), mean (cross), and 25–75% percentiles (box limits). Whiskers extend from minimum to**
 652 **maximum values. White and grey bands in the graphs on the right correspond to the fall/winter and spring/summer periods, and**
 653 **dotted vertical bars represent the first days of each season. ANOVA results of the multiple comparison between boxplots are**
 654 **summarized in Tables S4 and S5. Different lower-case letters indicate significant differences between ponds within a season and**
 655 **different upper-case letters indicate significant differences between seasons for a given pond.**

656 FOX and MOX followed the same seasonal variations as $\delta^{13}\text{C}\text{-CH}_4$ of dissolved CH_4 since both quantities were derived from
 657 isotopic models that include $\delta^{13}\text{C}\text{-CH}_4$ of dissolved CH_4 . $\delta^{13}\text{C}\text{-CH}_4$ of dissolved CH_4 , FOX, and MOX showed no significant
 658 differences between seasons in the two turbid-water ponds except in the Pêcherries pond where MOX was lower in winter
 659 ($1.3 \pm 0.86 \text{ mmol m}^{-2} \text{d}^{-1}$) than in summer ($12.3 \pm 10.5 \text{ mmol m}^{-2} \text{d}^{-1}$, Tukey's HSD test $p=0.0010$) and fall ($6.5 \pm 3.0 \text{ mmol m}^{-2}$
 660 d^{-1} , Tukey's HSD test $p=0.0254$) (Fig. 11). In the clear-water Silex pond, FOX was lower in spring ($42 \pm 12 \%$) and summer
 661 ($52 \pm 16 \%$) than in fall ($84 \pm 9 \%$) and winter ($76 \pm 12 \%$) (Tukey's HSD test $p < 0.0001$ for spring or summer versus fall or
 662 winter). In the clear-water Tenreuken pond, FOX was higher in fall ($73 \pm 5 \%$) than in spring ($42 \pm 17 \%$, Tukey's HSD test
 663 $p < 0.0001$) and summer ($57 \pm 11 \%$, Tukey's HSD test $p=0.0324$), and higher in winter ($71 \pm 10 \%$) than in spring ($42 \pm 17 \%$,
 664 Tukey's HSD test $p < 0.0001$). $\delta^{13}\text{C}\text{-CH}_4$ of dissolved CH_4 and FOX were statistically higher in the turbid-water ponds
 665 (Leybeek and Pêcherries) than in the clear-water ponds (Tenreuken and Silex) during spring and summer (Fig. 11) and than in
 666 the Tenreuken pond during fall and winter (Fig. 11; Tables S4 and S5). These seasonal differences led to an annual MOX
 667 that was statistically higher in the turbid-water ponds (10.8 and $7.2 \text{ mmol m}^{-2} \text{d}^{-1}$ in the Leybeek and Pêcherries ponds,

668 respectively) than the clear-water ponds (2.4 and 4.4 mmol m⁻² d⁻¹ in the Tenreuken and Silex ponds, respectively) (Tukey's
 669 HSD test p=<0.0001 for each turbid-water pond versus each clear-water pond). TSM and Chl-*a* concentrations were higher
 670 in the turbid-water ponds than in the clear-water ponds, particularly during productive phytoplanktonic periods of spring and
 671 summer (Fig. 3), when the highest difference of δ¹³C-CH₄ of dissolved CH₄, FOX, and MOX were observed between the
 672 turbid-water and the clear-water ponds (Fig. 11).

673 δ¹³C-CH₄ of dissolved CH₄, FOX, and MOX positively correlated to TSM and Chl-*a* concentrations (Fig. 12). These patterns
 674 could reflect the increase of micro-organisms including methanotrophs fixed on particles leading to an increase of MOX in
 675 parallel to an increase of TSM concentration (Abril et al., 2007). Fixed micro-organisms can grow on inorganic particles and
 676 aggregates of organic matter (Kirchman and Mitchell, 1982), but also on aggregates of living cyanobacteria (Li et al., 2021).
 677 An increase of particles in the water column increases light attenuation in the water column which would alleviate the
 678 inhibition of MOX by light (Dumestre et al., 1999; Murase and Sugimoto 2005; Morana et al., 2020), also possibly
 679 contributing to a positive relation between MOX and TSM and Chl-*a*, along the turbidity gradient. Both processes could co-
 680 occur contributing to the observed positive patterns between MOX and TSM and Chl-*a* concentrations.



681

682 **Figure 12:** ¹²C/¹³C ratio of CH₄ in surface waters (δ¹³C-CH₄, in ‰), fraction of CH₄ removed by methane oxidation (FOX, in %),
 683 and methane oxidation flux (MOX, in mmol m⁻² d⁻¹) versus total suspended matter concentration (TSM, in mg L⁻¹) and chlorophyll-*a*

684 concentration ($\text{Chl-}a$, in $\mu\text{g L}^{-1}$) in four urban ponds (Leybeek, Pêcherries, Tenreuken, and Silex) in the city of Brussels (Belgium)
685 from January 2022 to December 2023. Linear regression shown as black solid line (Table S11).

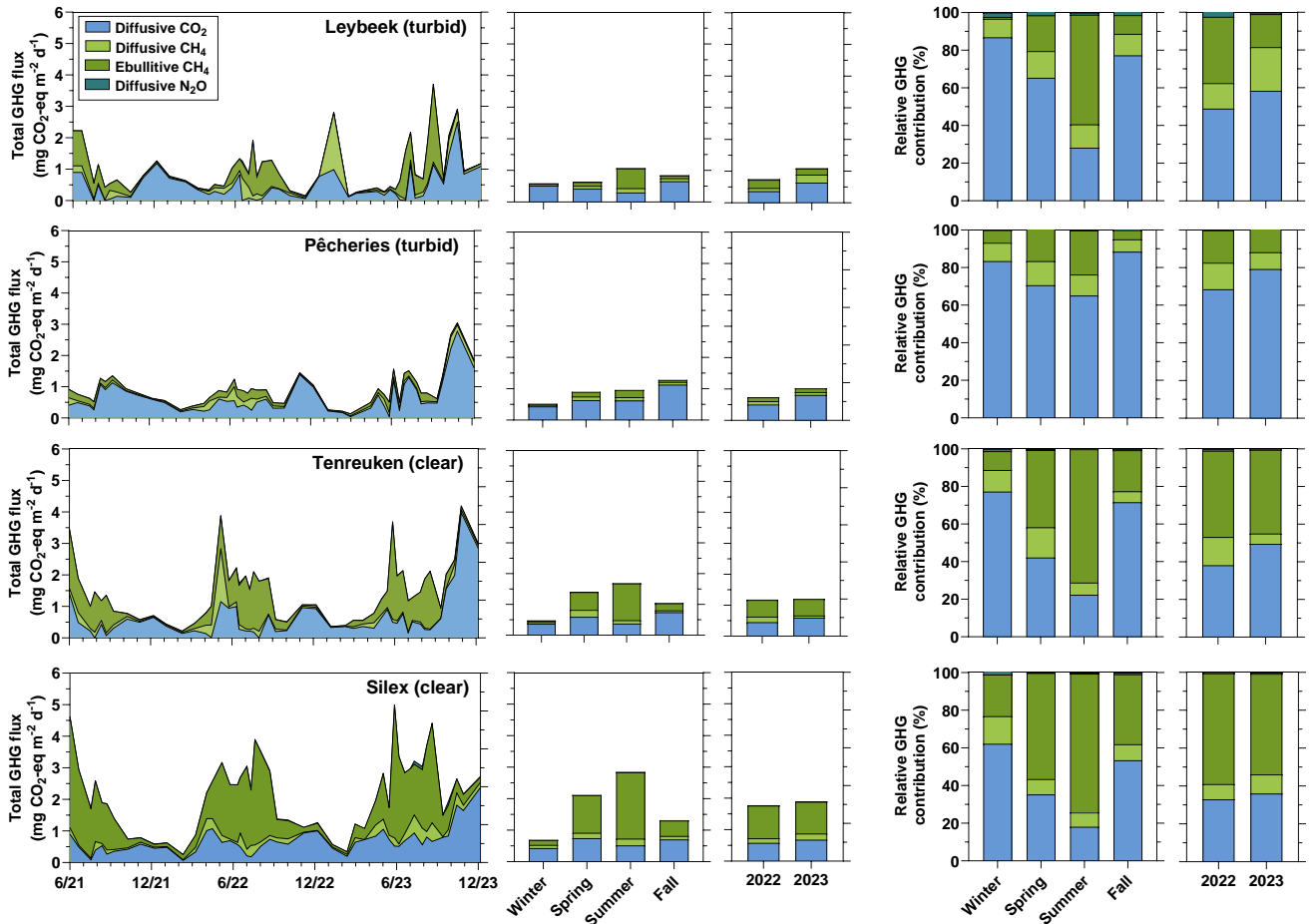
686 Figure S15 compares the main fluxes of dissolved CH_4 in the water column: MOX, diffusive CH_4 emissions, bubble
687 dissolution that were derived from measurements, and the sedimentary diffusive CH_4 flux that was computed as a closing
688 term (assuming a steady state) for comparative purposes. The dissolution of bubbles was a significantly smaller input term of
689 dissolved CH_4 compared to the diffusive sedimentary flux that represented $88\pm 18\%$ of the total input of CH_4 to the water
690 column (Tukey's HSD test $p < 0.0001$ in each pond). The low contribution of dissolution of bubbles resulted from the
691 shallowness of the studied ponds because bubble dissolution depends on the time spent by the bubble in the water column
692 during ascent, which is directly proportional to depth (McGinnis et al., 2006). MOX was a larger sink of dissolved CH_4 than
693 the diffusive CH_4 emission to the atmosphere in the four ponds, representing $80\pm 19\%$ and $80\pm 14\%$ of the total dissolved
694 CH_4 removal in the turbid-water Leybeek and Pêcherries ponds respectively (Tukey's HSD test $p < 0.0001$ for the two ponds),
695 and $59\pm 21\%$ and $51\pm 27\%$ in the clear-water Tenreuken and Silex ponds respectively (Tukey's HSD test $p = 0.3429$ for the
696 Tenreuken pond, and $p = 0.7634$ for the Silex pond). For all four ponds, MOX accounted for $66\pm 26\%$ of the total CH_4
697 dissolved removal from the water column, in agreement with other studies in lentic systems (Kankaala et al., 2006;
698 Bastviken et al., 2008; Morana et al., 2020; Reis et al., 2022).

699 3.7. Relative contribution of CO_2 , CH_4 and N_2O emissions

700 The emissions in CO_2 -eq for the 3 GHGs averaged per season for both 2022 and 2023 peaked seasonally in summer with 2.9
701 and 1.7 $\text{mg CO}_2\text{-eq m}^{-2} \text{d}^{-1}$ in the Silex and the Tenreuken ponds, respectively, and 1.1 $\text{mg CO}_2\text{-eq m}^{-2} \text{d}^{-1}$ in the Leybeek
702 pond (Fig. 13). The GHG fluxes peaked in fall in the Pêcherries pond, with 1.3 $\text{mg CO}_2\text{-eq m}^{-2} \text{d}^{-1}$. The higher value of the
703 total GHG emissions in fall compared to other seasons in the Pêcherries pond was due to an increase of CO_2 emissions in fall
704 that surpassed the peak of CH_4 emissions in summer. The GHG fluxes were the lowest in winter with 1.3 and 0.9 $\text{mg CO}_2\text{-eq}$
705 $\text{m}^{-2} \text{d}^{-1}$ in the Silex and the Tenreuken ponds, respectively, and 0.8 and 0.6 $\text{mg CO}_2\text{-eq m}^{-2} \text{d}^{-1}$ in the Pêcherries and the
706 Leybeek ponds, respectively. The relative contribution of ebullitive CH_4 fluxes peaked in summer in all four ponds, 73.8%
707 and 70.9% in the Silex and the Tenreuken ponds, respectively, and 23.6% and 58.3% in the Pêcherries and the Leybeek
708 ponds, respectively. The relative contribution of ebullitive CH_4 fluxes was lowest in winter with 22.1% and 10.0% in the
709 Silex and the Tenreuken ponds, respectively, and 6.7% and 1.0% in the Pêcherries and the Leybeek ponds, respectively.

710 The annual emissions in CO_2 -eq of the three GHGs (CO_2 , CH_4 , and N_2O) in 2022 and 2023 were higher in the two clear-
711 water ponds (1.3 ± 0.5 and 1.8 ± 0.9 $\text{mg CO}_2\text{-eq m}^{-2} \text{d}^{-1}$ in the Tenreuken and Silex ponds, respectively) than in the two turbid-
712 water ponds (1.0 ± 0.2 and 0.9 ± 0.5 $\text{mg CO}_2\text{-eq m}^{-2} \text{d}^{-1}$ in the Leybeek and Pêcherries ponds, respectively) (Fig. 13) (Tukey's
713 HSD test $p < 0.0001$ for Silex versus Pêcherries, $p < 0.0001$ for Silex versus Leybeek, $p = 0.0107$ for Tenreuken versus
714 Pêcherries, and $p = 0.0467$ for Tenreuken versus Leybeek) due to higher total CH_4 emissions (diffusive+ebullitive) in clear-
715 water ponds (0.7 ± 0.4 and 1.2 ± 0.5 $\text{mg CO}_2\text{-eq m}^{-2} \text{d}^{-1}$ in the Tenreuken and Silex ponds, respectively) than in turbid-water
716 ponds (0.2 ± 0.2 and 0.4 ± 0.3 $\text{mg CO}_2\text{-eq m}^{-2} \text{d}^{-1}$ in the Leybeek and Pêcherries ponds, respectively) (Tukey's HSD test
717 $p < 0.0001$ for Silex versus Pêcherries, $p < 0.0001$ for Silex versus Leybeek, $p = 0.0005$ for Tenreuken versus Pêcherries, and
718 $p = 0.0164$ for Tenreuken versus Leybeek), as there were no significant differences between the four ponds for CO_2 emissions
719 in 2022 and 2023 (Tukey's HSD test $p > 0.05$ for each comparison). N_2O emissions were significantly lower in the Pêcherries
720 pond than the Leybeek and Silex ponds (Tukey's HSD test $p = 0.0012$ for Pêcherries versus Leybeek, and $p = 0.0052$ for
721 Pêcherries versus Silex). The contribution of N_2O to the total GHG emissions was marginal and did not affect the differences
722 in total GHG fluxes between ponds, with the highest contribution observed in the Leybeek pond, with a contribution of
723 1.7%.

724 The majority of GHG emissions in CO₂-eq was related to CO₂ and CH₄ (diffusive+ebullitive) in the four ponds. In turbid-
 725 water ponds CO₂ represented the largest fraction of GHG emissions (68.5% (2022) and 79.3% (2023) in the Pêcheries pond,
 726 and 49.0% (2022) and 58.3% (2023) in the Leybeek pond). In clear-water ponds CH₄ represented the largest fraction of
 727 GHG emissions (66.5% (2022) and 63.3% (2023) in the Silex pond, and 60.8% (2022) and 50.0% (2023) in the Tenreuken
 728 pond). The higher annual GHG emissions in CO₂-eq from the two clear-water ponds than the turbid-water ponds were
 729 related to the higher contribution of ebullitive CH₄ fluxes.



730

731 **Figure 13: Temporal evolution and relative contribution of emissions to the atmosphere of CO₂ (diffusive), CH₄ (diffusive and**
 732 **ebullitive), and N₂O (diffusive) expressed in CO₂ equivalents (in mg CO₂-eq m⁻² d⁻¹), in four urban ponds (Leybeek, Pêcheries,**
 733 **Tenreuken, and Silex) in the city of Brussels (Belgium) from June 2021 to December 2023. Averages per season include data from**
 734 **2021, 2022, and 2023. Year 2023 had a higher annual precipitation (1011 mm) than year 2022 (701 mm).**

735 The annual GHG fluxes increased from 2022 to 2023 due to an increase in relative contribution of CO₂ diffusive emissions
 736 in all four ponds. Diffusive CO₂ emissions averaged annually in all four ponds 0.5 mg CO₂ m⁻² d⁻¹ in 2022 and 0.7 mg CO₂
 737 m⁻² d⁻¹ in 2023. Diffusive CO₂ emissions were two times higher in summer 2023 than in summer 2022, and 2.5 times higher
 738 in fall 2023 than in fall 2022, for similar values between 2023 and 2022 in spring and winter (1.1 higher and 1.1 lower,
 739 respectively). Air temperatures were similar in both years (annual average of 12.2°C in 2022 and 12.1°C in 2023) with
 740 winter, spring and summer marginally colder in 2023 than in 2022 (-0.5, -1.1°C and -0.4°C, respectively), and fall
 741 marginally warmer in 2023 than 2022 (+0.6°C). Spring and summer were rainier in 2023 than 2022 (2.2 and 2.5 and times,
 742 respectively) but fall and winter precipitations were relatively similar in both years (1.4 times wetter and 1.2 times drier in
 743 2023 than 2022, respectively). Higher precipitations are likely to increase the inputs of organic and inorganic carbon from
 744 soils to ponds by ground-waters, soil-waters, and surface runoff, as previously shown in other lentic systems (*e.g.* Marotta et

745 al., 2011; Holgerson, 2015). The highest seasonal increase of diffusive CO₂ emissions was observed in fall 2023. While this
746 hypothesis is only based on the comparison of two years, the increase of the relative contribution of CO₂ diffusive emissions
747 was observed in all four ponds which suggests a common uniform driver that would be consistent with a large variation
748 weather such as annual precipitation. The El Niño event in 2023 induced low-level cyclonic wind anomalies and higher
749 precipitation over Western Europe, including Belgium (Chen et al., 2024).

750 **4. Conclusions**

751 We found very marked differences in CH₄ dynamics between the two clear-water macrophyte-dominated ponds (Tenreuken
752 and Silex) and the two turbid-water phytoplankton-dominated ponds (Pêcherries and Leybeek) of the city of Brussels. MOX
753 was more important in the two turbid-water ponds compared to the clear-water ponds. MOX correlated to TSM and Chl-*a*
754 concentrations possibly owing to a higher abundance of methanotrophs in the water column fixed to particles and/or an
755 attenuation of light limitation of MOX. Ebullitive CH₄ emissions were higher in the two clear-water ponds than the two
756 turbid-water ponds, possibly related to high availability of macrophyte organic matter. The annual diffusive N₂O and CO₂
757 fluxes in 2022-2023 were not statistically different in the two clear-water ponds (Tenreuken and Silex) and in the two turbid-
758 water ponds (Pêcherries and Leybeek). Other studies have found no difference in N₂O sedimentary production in lakes with
759 high and low density of submerged macrophytes. We hypothesize that in human impacted system such as the urban ponds in
760 the city of Brussels, the strong range of variations of DIN was the main driver of N₂O levels and over-rides other possible
761 drivers such as presence or absence of macrophytes. Such a hypothesis was consistent with an overall positive relation
762 between %N₂O and DIN in the urban ponds of the city of Brussels irrespective of presence or absence of macrophytes
763 (Bauduin et al., 2024; this study). We hypothesize that CO₂ fluxes were relatively invariant among the four sampled ponds
764 because of they were of similar size and depth, and that they were all relatively productive irrespective of whether from
765 phytoplankton or submerged macrophytes.

766 The total (diffusive and ebullitive) CH₄ emissions represented 57.7±28.9 % (ranging seasonally from 4.9 to 99.9%) of total
767 GHG emissions in CO₂-eq in the two clear-water ponds compared to 41.0±28.7 % (ranging seasonally from 2.8 to 99.9%) in
768 the two turbid-water ponds. CO₂ represented nearly all the remainder of total GHG emissions in CO₂-eq, and N₂O
769 represented a very marginal fraction (0.8±1.6 %, ranging from 0.0% to 14.9%, with the maximum coinciding with minimal
770 total CO₂-eq GHG flux in the Leybeek pond). The seasonal variations of GHG emissions were dominated by CH₄ ebullitive
771 seasonal variations that peaked in summer (both quantitatively and relatively), as CH₄ ebullition was strongly related to
772 temperature. The pCO₂ values in the four sampled ponds increased with precipitation at seasonal scale, probably in relation
773 to higher inputs of organic and inorganic carbon by surface runoff. Years 2022 and 2023 were abnormally dry and wet,
774 respectively, and consequently, the GHG emissions were higher in 2023 mainly due to an increase in the relative
775 contribution of CO₂ emissions, probably in response to a strong El Niño event. This would suggest that variations of
776 precipitation also affected year-to-year variations of CO₂ emissions in addition to partly regulating seasonal variations of
777 CO₂ emissions from the four studied ponds.

778 **Data availability.** Timestamped and georeferenced data-set is available at [10.5281/zenodo.11103556](https://zenodo.org/doi/10.5281/zenodo.11103556).

779 **Author contributions.** AVB and NG conceived the study; TB collected field samples; TB and AVB made the laboratory
780 analysis; TB and AVB jointly interpreted data and drafted the manuscript with substantial inputs from NG.

781 **Competing interests.** The authors declare that they have no conflict of interest.

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791 **References**

792 Aben, R. C. H., Barros, N., Van Donk, E., Frenken, T., Hilt, S., Kazanjian, G., Lamers, L. P. M., Peeters, E. T. H. M., Roelofs, J.G.M., de
793 Senerpont Domis, L. S., Stephan, S., Velthuis, M., Van de Waal, D., Wik, M., Thornton, B., Wilkinson, J., Delsontro, T., and Kosten, S.:
794 Cross continental increase in methane ebullition under climate change. *Nature communications*, 8(1), 1682.
795 <https://doi.org/10.1038/s41467-017-01535-y>, 2017.

796 Abril, G., Commarieu, M. V., and Guérin, F.: Enhanced methane oxidation in an estuarine turbidity maximum. *Limnology and*
797 *oceanography*, 52(1), 470-475. <https://doi.org/10.4319/lo.2007.52.1.0470>, 2007.

798 Audet, J., Carstensen, M.V., Hoffmann, C.C., Lavaux, L., Thieme, K., and Davidson, T.A.: Greenhouse gas emissions from urban ponds
799 in Denmark. *Inland Waters* 10 (3), 373–385. <https://doi.org/10.1080/20442041.2020.1730680>, 2020.

800 Baliña, S., Sanchez, M. L., Izaguirre, I., and del Giorgio, P. A.: Shallow lakes under alternative states differ in the dominant greenhouse
801 gas emission pathways. *Limnology and Oceanography*, 68(1), 1-13. <https://doi.org/10.1002/lno.12243>, 2023.

802 Barko, J. W., Gunnison, D., and Carpenter, S. R.: Sediment interactions with submersed macrophyte growth and community dynamics.
803 *Aquatic botany*, 41(1-3), 41-65. [https://doi.org/10.1016/0304-3770\(91\)90038-7](https://doi.org/10.1016/0304-3770(91)90038-7), 1991.

804 Bartosiewicz, M., Maranger, R., Przytulska, A., and Laurion, I.: Effects of phytoplankton blooms on fluxes and emissions of greenhouse
805 gases in a eutrophic lake. *Water Research*, 196, 116985. <https://doi.org/10.1016/j.watres.2021.116985>, 2021.

806 Bastviken D., Ejlertsson J. and Tranvik L.: Measurement of methane oxidation in lakes: A comparison of methods. *Environmental*
807 *Science & Technology*, 36, 3354-3361. <https://doi.org/10.1021/es010311p>, 2002.

808 Bastviken, D., Cole, J. J., Pace, M. L., and Van de Bogert, M. C.: Fates of methane from different lake habitats: Connecting whole-lake
809 budgets and CH₄ emissions. *Journal of Geophysical Research: Biogeosciences*, 113(G2). <https://doi.org/10.1029/2007JG000608>, 2008.

810 Bastviken, D., Cole, J., Pace, M., and Tranvik, L.: Methane emissions from lakes: Dependence of lake characteristics, two regional
811 assessments, and a global estimate. *Global biogeochemical cycles*, 18(4). <https://doi.org/10.1029/2004GB002238>, 2004.

812 Bastviken, D., Treat, C.C., Pangala, S.R., Gauci, V., Enrich-Prast, A., Karlson, M., Gålfalk, M., Romano, M.B., and Sawakuchi, H.O.: The
813 importance of plants for methane emission at the ecosystem scale. *Aquat Bot* 184, 103596. <https://doi.org/10.1016/j.aquabot.2022.103596>,
814 2023.

815 Bauduin, T., Gypens, N., and Borges, A. V.: Seasonal and spatial variations of greenhouse gas (CO₂, CH₄ and N₂O) emissions from
816 urban ponds in Brussels. *Water Research*, 121257. <https://doi.org/10.1016/j.watres.2024.121257>, 2024.

817 Berberich, M. E., Beaulieu, J. J., Hamilton, T. L., Waldo, S., and Buffam, I.: Spatial variability of sediment methane production and
818 methanogen communities within a eutrophic reservoir: importance of organic matter source and quantity. *Limnol. Oceanogr.* 65, 1–23.
819 <https://doi.org/10.1002/lno.11392>, 2020.

820 Borges, A.V., Darchambeau, F., Lambert, T., Morana, C., Allen, G.H., Tambwe, E., and Bouillon, S.: Variations in dissolved greenhouse
821 gases (CO₂, CH₄, N₂O) in the Congo River network overwhelmingly driven by fluvial-wetland connectivity. *Biogeosciences* 16 (19),
822 3801–3834. <https://doi.org/10.5194/bg-16-3801-2019>, 2019.

- 823 Borges, A.V., Deirmendjian, L., Bouillon, S., Okello, W., Lambert, T., Roland, F.A.E., Razanamahandry, V.F., Voarintsoa, N.R.G.,
824 Darchambeau, F., Kimirei, I.A., Descy, J., Allen, G.H., and Morana, C.: Greenhouse gas emissions from African lakes are no longer a
825 blind spot. *Sci. Adv.* 8 (25), eabi8716. <https://doi.org/10.1126/sciadv.abi8716>, 2022.
- 826 Brans, K.I., Engelen, J.M., Souffreau, C., and De Meester, L.: Urban hot-tubs: local urbanization has profound effects on average and
827 extreme temperatures in ponds. *Landsc. Urban Plan.* 176, 22–29. <https://doi.org/10.1016/j.lurbplan.2018.05.001>, 2018.
- 828 Cael, B. B., Heathcote, A. J., and Seekell, D. A.: The volume and mean depth of Earth's lakes. *Geophysical Research Letters*, 44(1), 209-
829 218. <https://doi.org/10.1002/2016GL071378>, 2017.
- 830 Casas-Ruiz, J.P., Jakobsson, J., and del Giorgio, P.A.: The role of lake morphometry in modulating surface water carbon concentrations in
831 boreal lakes. *Environ. Res. Lett.* 16 (7), 074037 <https://doi.org/10.1088/1748-9326/ac0be3>, 2021.
- 832 Chen, B., Zhang, L., and Wang, C.: Distinct impacts of the central and eastern Atlantic Niño on the European climate. *Geophysical
833 Research Letters*, 51(2), e2023GL107012. <https://doi.org/10.1029/2023GL107012>, 2024.
- 834 Choudhury, M. I., McKie, B. G., Hallin, S., and Ecke, F.: Mixtures of macrophyte growth forms promote nitrogen cycling in wetlands.
835 *Science of the Total Environment*, 635, 1436-1443. <https://doi.org/10.1016/j.scitotenv.2018.04.193>, 2018.
- 836 Clifford, C.C., and Heffernan, J.B.: Artificial aquatic ecosystems. *Water* 10 (8), 1096. <https://doi.org/10.3390/w10081096>, 2018.
- 837 Codispoti, L.A., and Christensen, J.P.: Nitrification, denitrification and nitrous oxide cycling in the eastern tropical South Pacific Ocean.
838 *Mar. Chem.* 16 (4), 277–300. [https://doi.org/10.1016/0304-4203\(85\)90051-9](https://doi.org/10.1016/0304-4203(85)90051-9), 1985.
- 839 Cole, J.J., and Caraco, N.F.: Atmospheric exchange of carbon dioxide in a low-wind oligotrophic lake measured by the addition of SF6.
840 *Limnol. Oceanogr.* 43 (4), 647–656. <https://doi.org/10.4319/lo.1998.43.4.0647>, 2018.
- 841 Coleman, D. D., Risatti, J. B., and Schoell, M.: Fractionation of carbon and hydrogen isotopes by methane oxidizing bacteria. *Geochimica
842 Cosmochimica Acta*, 45, 1033–1037. [https://doi.org/10.1016/0016-7037\(81\)90129-0](https://doi.org/10.1016/0016-7037(81)90129-0), 1981.
- 843 Conrad, R., Noll, M., Claus, P., Klose, M., Bastos, W. R., and Enrich-Prast, A.: Stable carbon isotope discrimination and microbiology of
844 methane formation in tropical anoxic lake sediments. *Biogeosciences*, 8(3), 795-814. <https://doi.org/10.5194/bg-8-795-2011>, 2011.
- 845 Conrad, R.: Control of Methane Production in Terrestrial Ecosystems, In: Andreae, M.O. and Schimel, D.S., Ed., *Exchange of Trace Gases
846 between Terrestrial Ecosystems and the Atmosphere*, John Wiley, New York, 39-58, 1989.
- 847 Conrad, R.: Quantification of methanogenic pathways using stable carbon isotopic signatures: a review and a proposal. *Organic
848 geochemistry*, 36(5), 739-752. <https://doi.org/10.1016/j.orggeochem.2004.09.006>, 2005
- 849 Dan, Z., Chuan, W., Qiaohong, Z., and Xingzhong, Y.: Sediments nitrogen cycling influenced by submerged macrophytes growing in
850 winter. *Water Science and Technology*, 83(7), 1728-1738. <https://doi.org/10.2166/wst.2021.081>, 2021.
- 851 Davidson, T.A., Audet, J., Svenning, J.C., Lauridsen, T.L., Søndergaard, M., Landkildehus, F., and Jeppesen, E.: Eutrophication effects on
852 greenhouse gas fluxes from shallow-lake mesocosms override those of climate warming. *Glob. Chang. Biol.* 21 (12), 4449–4463.
853 <https://doi.org/10.1111/gcb.13062>, 2015.
- 854 Deemer, B. R., and Holgerson, M. A.: Drivers of methane flux differ between lakes and reservoirs, complicating global upscaling efforts.
855 *Journal of Geophysical Research: Biogeosciences*, 126(4) <https://doi.org/10.1029/2019JG005600>, 2021.
- 856 DelSontro, T., Beaulieu, J. J., and Downing, J. A.: Greenhouse gas emissions from lakes and impoundments: Upscaling in the face of
857 global change. *Limnology and Oceanography Letters*, 3(3), 64-75. <https://doi.org/10.1002/lol2.10073>, 2018.
- 858 DelSontro, T., L. Boutet, A. St-Pierre, P.A. del Giorgio, and Y.T.: Prairie, Methane ebullition and diffusion from northern ponds and lakes
859 regulated by the interaction between temperature and system productivity, *Limnol. Oceanogr.* 61(S1), S62-S77
860 <https://doi.org/10.1002/lno.10335>, 2016.
- 861 DelSontro, T., Kunz, M. J., Kempter, T., Wüest, A., Wehrli, B., and Senn, D. B.: Spatial Heterogeneity of Methane Ebullition in a Large
862 Tropical Reservoir, *Environmental Science & Technology* 45 (23), 9866-9873, <https://doi.org/10.1021/es2005545>, 2011.
- 863 Deng, Hg., Zhang, J., Wu, Jj., Yao, X., and Yang, L.-W.: Biological denitrification in a macrophytic lake: implications for macrophytes-
864 dominated lake management in the north of China. *Environ Sci Pollut Res* 27, 42460–42471. <https://doi.org/10.1007/s11356-020-10230-3>
865 , 2020.
- 866 Desrosiers, K., DelSontro, T., and del Giorgio, P. A.: Disproportionate Contribution of Vegetated Habitats to the CH4 and CO2 Budgets of
867 a Boreal Lake. *Ecosystems*, 1-20. <https://doi.org/10.1007/s10021-021-00730-9>, 2022.

- 868 Downing, J. A.: Emerging global role of small lakes and ponds: little things mean a lot. *Limnetica*, 29(1), 0009-24.
869 <https://doi.org/10.23818/limn.29.02>, 2009.
- 870 Dickson, A.G.; Sabine, C.L. and Christian, J.R.: Guide to best practices for ocean CO₂ measurement. Sidney, British Columbia, North
871 Pacific Marine Science Organization, 191pp. (PICES Special Publication 3; IOCCP Report 8). <https://doi.org/10.25607/OBP-1342>, 2007.
- 872 Dumestre, J. F., Guézennec, J., Galy-Lacaux, C., Delmas, R., Richard, S., and Labroue, L.: Influence of light intensity on methanotrophic
873 bacterial activity in Petit Saut Reservoir, French Guiana. *Applied and environmental microbiology*, 65(2), 534-539.,
874 <https://doi.org/10.1128/aem.65.2.534-539.1999>, 1999.
- 875 Dutton, G., Elkins II, J., Hall, B., NOAA ESRL, Earth System Research Laboratory Halocarbons and Other Atmospheric Trace Gases
876 Chromatograph for Atmospheric Trace Species (CATS) Measurements. NOAA National Centers for Environmental Information.
877 <https://doi.org/10.7289/V5X0659V>. Version 1. [Database: atmospheric nitrous oxide N₂O] [2024-03-27], 2017.
- 878 Goeckner, A. H., Lusk, M. G., Reisinger, A. J., Hosen, J. D., and Smoak, J. M.: Florida's urban stormwater ponds are net sources of
879 carbon to the atmosphere despite increased carbon burial over time. *Communications earth & environment*, 3(1), 53,
880 <https://doi.org/10.1038/s43247-022-00384-y> 2022.
- 881 Gorsky, A.L., Racanelli, G.A., Belvin, A.C., and Chambers, R.M.: Greenhouse gas flux from stormwater ponds in southeastern Virginia
882 (USA). *Anthropocene* 28, 100218. <https://doi.org/10.1016/j.ancene.2019.100218>, 2019.
- 883 Gorsky, A. L., Dugan, H. A., Wilkinson, G. M., and Stanley, E. H.: Under-ice oxygen depletion and greenhouse gas supersaturation in
884 north temperate urban ponds. *Journal of Geophysical Research: Biogeosciences*, 129(6), <https://doi.org/10.1029/2024JG008120>, 2024.
- 885 Grasset, C., Abril, G., Mendonça, R., Roland, F., and Sobek, S.: The transformation of macrophyte-derived organic matter to methane
886 relates to plant water and nutrient contents. *Limnology and Oceanography*, 64(4), 1737-1749, <https://doi.org/10.1002/lno.11148>, 2019.
- 887 Grasset, C., Sobek, S., Scharnweber, K., Moras, S., Villwock, H., Andersson, S., Hiller, C., Nydahl, A.C., Chaguaceda, F., Colom, W., and
888 Tranvik, L.J.: The CO₂-equivalent balance of freshwater ecosystems is non-linearly related to productivity. *Glob. Chang. Biol.* 26 (10),
889 5705–5715. <https://doi.org/10.1111/gcb.15284>, 2020.
- 890 Grasshoff, K., and Johannsen, H.: A new sensitive and direct method for the automatic determination of ammonia in sea water. *ICES J.*
891 *Mar. Sci.* 34 (3), 516–521. <https://doi.org/10.1093/icesjms/34.3.516>, 1972.
- 892 Grasshoff, K., Kremling, K., and Ehrhardt, M.: *Methods of Seawater Analysis: Determination of Nitrite*. John Wiley & Sons, 2009.
- 893 Greinert J., and D.F. McGinnis: Single bubble dissolution model – The graphical user interface SiBu-GUI, *Environmental Modelling &*
894 *Software*, 24, 1012-1013, <https://doi.org/10.1016/j.envsoft.2008.12.011>, 2009.
- 895 Grinham, A., Albert, S., Deering, N., Dunbabin, M., Bastviken, D., Sherman, B., Lovelock, C.E., and Evans, C.D.: The importance of
896 small artificial water bodies as sources of methane emissions in Queensland, Australia. *Hydrol. Earth Syst. Sci.* 22 (10), 5281–5298.
897 <https://doi.org/10.5194/hess-22-5281-2018>, 2018.
- 898 Harpenslager, S. F., Thiemer, K., Levertz, C., Misteli, B., Sebola, K. M., Schneider, S. C., Hilt, S., and Köhler, J.: Short-term effects of
899 macrophyte removal on emission of CO₂ and CH₄ in shallow lakes. *Aquatic Botany*, 182, 103555.
900 <https://doi.org/10.1016/j.aquabot.2022.103555>, 2022.
- 901 Herrero Ortega, S., Romero Gonz´alez-Quijano, C., Casper, P., Singer, G.A., and Gessner, M.O.: Methane emissions from contrasting
902 urban freshwaters: rates, drivers, and a whole-city footprint. *Glob. Chang. Biol.* 25 (12), 4234–4243. <https://doi.org/10.1111/gcb.14799>,
903 2019.
- 904 Hilt, S., Brothers, S., Jeppesen, E., Veraart, A. J., and Kosten, S.: Translating regime shifts in shallow lakes into changes in ecosystem
905 functions and services. *BioScience*, 67(10), 928-936. <https://doi.org/10.1093/biosci/bix106>, 2017.
- 906 Holgerson, M., and Raymond, P.: Large contribution to inland water CO₂ and CH₄ emissions from very small ponds. *Nat. Geosci.* 9, 222–
907 226. <https://doi.org/10.1038/ngeo2654>, 2016.
- 908 Holgerson, MA.: Drivers of carbon dioxide and methane supersaturation in small, temporary ponds, *Biogeochemistry* 124:305–318.
909 <https://doi.org/10.1007/s10533-015-0099-y>, 2015.
- 910 Huttunen, J. T., Alm, J., Liikanen, A., Juutinen, S., Larmola, T., Hammar, T., Silvola, T., and Martikainen, P. J.: Fluxes of methane,
911 carbon dioxide and nitrous oxide in boreal lakes and potential anthropogenic effects on the aquatic greenhouse gas emissions.
912 *Chemosphere*, 52(3), 609-621. [https://doi.org/10.1016/S0045-6535\(03\)00243-1](https://doi.org/10.1016/S0045-6535(03)00243-1), 2003.

- 913 Hyvönen, T., Ojala, A., Kankaala, P., & Martikainen, P. J.: Methane release from stands of water horsetail (*Equisetum fluviatile*) in a
914 boreal lake, *Freshwat. Biol.*, 40, 275–284. <https://doi.org/10.1046/j.1365-2427.1998.00351.x>, 1998.
- 915 Johnson, M.S., Matthews, E., Du, J., Genovese, V., and Bastviken, D.: Methane Emission from Global Lakes: New Spatiotemporal Data
916 and Observation-Driven Modeling of Methane Dynamics Indicates Lower Emissions. *Journal of Geophysical Research: Biogeosciences*,
917 127(7). <https://doi.org/10.1029/2022JG006793>, 2022.
- 918 Juutinen, S., Alm, J., Larmola, T., Huttunen, J. T., Morero, M., Martikainen, P. J., and Silvola, J.: Major implication of the littoral zone for
919 methane release from boreal lakes, *Global Biogeochem. Cycles*, 17(4), 1117, <https://doi.org/10.1029/2003GB002105>, 2003.
- 920 Kankaala, P., Huotari, J., Peltomaa, E., Saloranta, T., and Ojala, A.: Methanotrophic activity in relation to methane efflux and total
921 heterotrophic bacterial production in a stratified, humic, boreal lake. *Limnology and Oceanography*, 51(2), 1195-1204.
922 <https://doi.org/10.4319/lo.2006.51.2.1195>, 2006.
- 923 Kankaala, P., Huotari, J., Tulonen, T., & Ojala, A.: A Lake-size dependent physical forcing drives carbon dioxide and methane effluxes
924 from lakes in a boreal landscape. *Limnol Oceanogr* 58:1915–1930. <https://doi.org/10.4319/lo.2013.58.6.1915>, 2013.
- 925 Keller, M., and R. F. Stallard: Methane emission by bubbling from Gatun Lake, Panama, *J. Geophys. Res.*, 99(D4), 8307–8319,
926 doi:[10.1029/92JD02170](https://doi.org/10.1029/92JD02170), 1994.
- 927 Kelly, C. A., and Chynoweth, D. P.: The contributions of temperature and of the input of organic matter in controlling rates of sediment
928 methanogenesis 1. *Limnology and Oceanography*, 26(5), 891-897. <https://doi.org/10.4319/lo.1981.26.5.0891>, 1981.
- 929 Kirchman D., and Mitchell, R.: Contribution of Particle-Bound Bacteria to Total Microheterotrophic Activity in Five Ponds and Two
930 Marshes, *Applied And Environmental Microbiology*, 43, 200-209, <https://doi.org/10.1128/aem.43.1.200-209.1982>, 1982.
- 931 Koroleff, J.: Determination of total phosphorus by alkaline persulphate oxidation. *Methods of Seawater Analysis*. Verlag Chemie,
932 Wienheim, pp. 136–138, 1983.
- 933 Kotsyurbenko, O. R., Friedrich, M. W., Simankova, M. V., Nozhevnikova, A. N., Golyshin, P. N., Timmis, K. N., and Conrad, R.: Shift
934 from acetoclastic to H₂-dependent methanogenesis in a West Siberian peat bog at low pH values and isolation of an acidophilic
935 *Methanobacterium* strain. *Applied and Environmental Microbiology*, 73(7), 2344-2348. <https://doi.org/10.1128/AEM.02413-06>, 2007.
- 936 Lan, X., K.W. Thoning, and E.J. Dlugokencky: Trends in globally-averaged CH₄, N₂O, and SF₆ determined from NOAA Global
937 Monitoring Laboratory measurements [data set]. Version 2024-08, <https://doi.org/10.15138/P8XG-AA10>, 2024.
- 938 Lauerwald, R., Regnier, P., Figueiredo, V., Enrich-Prast, A., Bastviken, D., Lehner, B., Maavara, T., and Raymond, P.: Natural Lakes Are
939 a Minor Global Source of N₂O to the Atmosphere. *Global Biogeochemical Cycles*, 33(12), 1564–1581.
940 <https://doi.org/10.1029/2019GB006261>, 2019.
- 941 Lauerwald, R., Allen, G. H., Deemer, B. R., Liu, S., Maavara, T., Raymond, P., Alcott, L., Bastviken, D., Hastie, A., Holgerson, M.A.,
942 Johnson, M. S., Lehner, B., Lin, P., Marzadri, A., Ran, L., Tian, H., Yang, X., Yao, Y., and Regnier, P.: Inland water greenhouse gas
943 budgets for RECCAP2: 2. Regionalization and homogenization of estimates. *Global Biogeochemical Cycles*, 37,
944 e2022GB007658. <https://doi.org/10.1029/2022GB007658>, 2023.
- 945 Li, C., Hambright, K. D., Bowen, H. G., Trammell, M. A., Grossart, H. P., Burford, M. A., Hamilton, D.P., Jiang, H., Latour, D., Meyer,
946 E. I., Padišák, J., Zamor, R. M. and Krumholz, L. R.: Global co-occurrence of methanogenic archaea and methanotrophic bacteria in
947 *Microcystis* aggregates, *Environmental Microbiology*, 23(11)<https://doi.org/10.1111/1462-2920.15691>, 2021.
- 948 Liptay, K., Chanton, J., Czepiel, P., and Mosher, B.: Use of stable isotopes to determine methane oxidation in landfill cover soils. *Journal*
949 *of Geophysical Research: Atmospheres*, 103(D7), 8243-8250. <https://doi.org/10.1029/97JD02630>, 1998.
- 950 Liu, Y., Conrad, R., Yao, T., Gleixner, G., and Claus, P.: Change of methane production pathway with sediment depth in a lake on the
951 Tibetan plateau. *Palaeogeography, Palaeoclimatology, Palaeoecology*, 474, 279-286. <https://doi.org/10.1016/j.palaeo.2016.06.021>, 2017.
- 952 Maavara, T., Lauerwald, R., Laruelle, G. G., Akbarzadeh, Z., Bouskill, N. J., Van Cappellen, P., and Regnier, P.: Nitrous oxide emissions
953 from inland waters: Are IPCC estimates too high? *Global Change Biology*, 25(2), 473–488. <https://doi.org/10.1111/gcb.145042>, 2019.
- 954 Marotta, H., Duarte, C. M., Pinho, L., and Enrich-Prast, A.: Rainfall leads to increased pCO₂ in Brazilian coastal lakes. *Biogeosciences*,
955 7(5), 1607-1614. <https://doi.org/10.5194/bg-7-1607-2010>, 2010.
- 956 Martinez-Cruz, K., Gonzalez-Valencia, R., Sepulveda-Jauregui, A., Plascencia- Hernandez, F., Belmonte-Izquierdo, Y., and Thalasso, F.:
957 Methane emission from aquatic ecosystems of Mexico City. *Aquat. Sci.* 79, 159–169. <https://doi.org/10.1007/s00027-016-0487-y>, 2017.

- 958 McCrackin, M.L., and Elser, J. J.: Atmospheric nitrogen deposition influences denitrification and nitrous oxide production in lakes,
959 *Ecology*, 91(2):528-39. <https://doi.org/10.1890/08-2210.1>, 2010.
- 960 McGinnis, D.F., Greinert, J., Artemov, Y., Beaubien, S.E., and Wüest, A.: The fate of rising methane bubbles in stratified waters: what
961 fraction reaches the atmosphere? *Journal of Geophysical Research* 111, C09007. <https://doi.org/10.1029/2005JC003183>, 2006.
- 962 Mengis, M., Gächter, R., and Wehrl, B.: Sources and sinks of nitrous oxide (N₂O) in deep lakes. *Biogeochemistry*, 38, 281-301.
963 <https://doi.org/10.1023/A:1005814020322>, 1997.
- 964 Morana, C., Bouillon, S., Nolla-Ardèvol, V., Roland, F. A., Okello, W., Descy, J. P., Nankabirwa, A., Nabafu, E., Springael, D., and
965 Borges, A. V.: Methane paradox in tropical lakes? Sedimentary fluxes rather than pelagic production in oxic conditions sustain
966 methanotrophy and emissions to the atmosphere, *Biogeosciences*, 17, 5209-5221, <https://doi.org/10.5194/bg-17-5209-2020>, 2020.
- 967 Morana C., Borges A.V., Roland F.A.E., Darchambeau F., Descy J.-P. and Bouillon S.: Methanotrophy within the water column of a large
968 meromictic tropical lake (Lake Kivu, East Africa). *Biogeosciences*, 12, 2077-2088. <https://doi.org/10.5194/bg-12-2077-2015>, 2015.
- 969 Murase, J., and Sugimoto, A.: Inhibitory effect of light on methane oxidation in the pelagic water column of a mesotrophic lake (Lake
970 Biwa, Japan). *Limnology and oceanography*, 50(4), 1339-1343. <https://doi.org/10.4319/lo.2005.50.4.1339>, 2005.
- 971 Natchimuthu, S., Panneer Selvam, B., and Bastviken, D.: Influence of weather variables on methane and carbon dioxide flux from a
972 shallow pond. *Biogeochemistry* 119, 403–413. <https://doi.org/10.1007/s10533-014-9976-z>, 2014.
- 973 Ni, M., Liang, X., Hou, L., Li, W., and He, C.: Submerged macrophytes regulate diurnal nitrous oxide emissions from a shallow eutrophic
974 lake: A case study of Lake Wuliangshuai in the temperate arid region of China. *Science of The Total Environment*, 811, 152451.
975 <https://doi.org/10.1016/j.scitotenv.2021.152451>, 2022a.
- 976 Ni, R., Xu, C., Shi, X., Yang, S., Li, L., Peng, X., and Song, L.: Acetoclastic methanogenesis pathway stability despite the high microbial
977 taxonomic variability in the transition from acidogenesis to methanogenesis during food waste anaerobic digestion. *Journal of Cleaner
978 Production*, 372, 133758. <https://doi.org/10.1016/j.jclepro.2022.133758>, 2022b.
- 979 Ollivier, Q.R., Maher, D.T., Pitfield, C., and Macreadie, P.I.: Punching above their weight: large release of greenhouse gases from small
980 agricultural dams. *Glob. Chang. Biol.* 25 (2), 721–732. <https://doi.org/10.1111/gcb.14477>, 2019.
- 981 Palacin-Lizarbe, C., Camarero, L., Hallin, S., Jones, C. M., and Catalan, J.: Denitrification rates in lake sediments of mountains affected by
982 high atmospheric nitrogen deposition. *Sci Rep* 10, 3003. <https://doi.org/10.1038/s41598-020-59759-w>, 2020.
- 983 Peacock, M., Audet, J., Bastviken, D., Cook, S., Evans, C.D., Grinham, A., Holgerson, M. A., Högbom, L., Pickard, A.E., Zieliński, P.,
984 and Futter, M.N.: Small artificial waterbodies are widespread and persistent emitters of methane and carbon dioxide. *Glob. Chang. Biol.*
985 27 (20), 5109–5123. <https://doi.org/10.1111/gcb.15762>, 2021.
- 986 Peacock, M., Audet, J., Jordan, S., Smeds, J., and Wallin, M.B.: Greenhouse gas emissions from urban ponds are driven by nutrient status
987 and hydrology. *Ecosphere* 10 (3), e02643. <https://doi.org/10.1002/ecs2.2643>, 2019.
- 988 Peretyatko, A., Symoens, J. J., and Triest, L.: Impact of macrophytes on phytoplankton in eutrophic peri-urban ponds, implications for
989 pond management and restoration. *Belgian Journal of Botany*, 83-99. <https://doi.org/10.2307/20794626>, 2007.
- 990 Rabaey, J. and Cotner, J.: Pond greenhouse gas emissions controlled by duckweed coverage. *Front. Environ. Sci.* 10, 889289
991 <https://doi.org/10.3389/fenvs2022.889289>, 2022.
- 992 Rabaey, J. S., and Cotner, J. B.: The influence of mixing on seasonal carbon dioxide and methane fluxes in ponds. *Biogeochemistry*, 1-18,
993 <https://doi.org/10.1007/s10533-024-01167-7>, 2024.
- 994 Ray, N. E., & Holgerson, M. A.: High Intra-Seasonal Variability in Greenhouse Gas Emissions from Temperate Constructed Ponds.
995 *Geophysical Research Letters*, 50(18), e2023GL104235, <https://doi.org/10.1029/2023GL104235>, 2023
- 996 Ray, N. E., Holgerson, M. A., Andersen, M. R., Bikše, J., Bortolotti, L. E., Futter, M., Kokorite, I., Law, A., McDonald, C., Mesman, J.P.,
997 Peacock, M., Richardson, D.C., Arsenault, J., Bansal, S., Cawley, K., Kuhn, M., Shahabinia, A.R., and Smufer, F.: Spatial and temporal
998 variability in summertime dissolved carbon dioxide and methane in temperate ponds and shallow lakes. *Limnology and Oceanography*,
999 68(7), 1530-1545. <https://doi.org/10.1002/lno.12362>, 2023.
- 1000 Raymond, P. A., Hartmann, J., Lauerwald, R., Sobek, S., McDonald, C., Hoover, M., Butman, D., Striegl, R., Mayorga, E., Humborg, C.,
1001 Kortelainen, P., Dürr, H., Meybeck, M., Ciais, P., and Guth, P.: Global carbon dioxide emissions from inland waters. *Nature*, 503(7476),
1002 355–359. <https://doi.org/10.1038/nature12760>, 2013.

- 1003 Reis, P.C.J., Thottathil, S.D. and Prairie, Y.T. The role of methanotrophy in the microbial carbon metabolism of temperate lakes. *Nat*
1004 *Commun* 13, 43. <https://doi.org/10.1038/s41467-021-27718-2>, 2022.
- 1005 Reitsema, R. E., Meire, P., and Schoelynck, J.: The future of freshwater macrophytes in a changing world: dissolved organic carbon
1006 quantity and quality and its interactions with macrophytes. *Frontiers in Plant Science*, 9, 301954. <https://doi.org/10.3389/fpls.2018.00629>,
1007 2018.
- 1008 Rocher-Ros, G., Stanley, E. H., Loken, L. C., Casson, N. J., Raymond, P. A., Liu, S., Amatulli, G., and Sponseller, R. A.: Global methane
1009 emissions from rivers and streams. *Nature* 621:530–535. <https://doi.org/10.1038/s41586-023-06344-6>, 2023.
- 1010 Rosentreter, J. A., Borges, A. V., Deemer, B. R., Holgerson, M. A., Liu, S., Song, C., Melack, J., Raymond, P. A., Duarte, C. M., Allen, G.
1011 H., Olefeldt, D., Poulter, B., Battin, T. I., and Eyre, B. D.: Half of global methane emissions come from highly variable aquatic ecosystem
1012 sources. *Nature Geoscience*, 14(4), 225–230. <https://doi.org/10.1038/s41561-021-00715-2>, 2021.
- 1013 Sand-Jensen, K., & Staehr, P. A.: Scaling of pelagic metabolism to size, trophy and forest cover in small Danish lakes. *Ecosystems*, 10,
1014 128-142. <https://doi.org/10.1007/s10021-006-9001-z>, 2007.
- 1015 Scandella, B. P., Varadharajan, C., Hemond, H. F., Ruppel, C., and Juanes, R.: A conduit dilation model of methane venting from lake
1016 sediments. *Geophysical Research Letters*, 38(6). <https://doi.org/10.1029/2011GL046768>, 2011.
- 1017 Scheffer, M., Hosper, S. H., Meijer, M. L., Moss, B., and Jeppesen, E. (1993). Alternative equilibria in shallow lakes. *Trends in ecology &*
1018 *evolution*, 8(8), 275-279. [https://doi.org/10.1016/0169-5347\(93\)90254-M](https://doi.org/10.1016/0169-5347(93)90254-M)
- 1019 Schulz S., Matsuyama, H., and Conrad, R.: Temperature dependence of methane production from different precursors in a profundal
1020 sediment (Lake Constance) *FEMS Microbiology Ecology*, 22, 207-213; <https://doi.org/10.1111/j.1574-6941.1997.tb00372.x>, 1997.
- 1021 Schulz S., and Conrad, R.: Influence of temperature on pathways to methane production in the permanently cold profundal sediment of
1022 Lake Constance. *FEMS Microbiology Ecology* 201-14; <https://doi.org/10.1111/j.1574-6941.1996.tb00299.x>, 1996.
- 1023 Singh, S.N., Kulshreshtha, K., and Agnihotri, S.: Seasonal dynamics of methane emission from wetlands. *Chemosphere Glob. Chang. Sci.*
1024 2 (1), 39–46. [https://doi.org/10.1016/S1465-9972\(99\)00046-X](https://doi.org/10.1016/S1465-9972(99)00046-X), 2000.
- 1025 Stanley, E. H., Casson, N. J., Christel, S. T., Crawford, J. T., Loken, L. C., and Oliver, S. K.: The ecology of methane in streams and
1026 rivers: patterns, controls, and global significance. *Ecological Monographs*, 86(2), 146–171. <https://doi.org/10.1890/15-1027>, 2016.
- 1027 Taoka, T., Iwata, H., Hirata, R., Takahashi, Y., Miyabara, Y., and Itoh, M.: Environmental controls of diffusive and ebullitive methane
1028 emissions at a subdaily time scale in the littoral zone of a midlatitude shallow lake. *Journal of Geophysical Research: Biogeosciences*, 125,
1029 e2020JG005753. <https://doi.org/10.1029/2020JG005753>, 2020.
- 1030 Theus, M. E., Ray, N. E., Bansal, S., and Holgerson, M. A.: Submersed macrophyte density regulates aquatic greenhouse gas emissions.
1031 *Journal of Geophysical Research: Biogeosciences*, 128(10), <https://doi.org/10.1029/2023JG007758>, 2023.
- 1032 Tokida, T., Miyazaki, T., Mizoguchi, M., Nagata, O., Takakai, F., Kagemoto, A., and Hatano, R.: Falling atmospheric pressure as a trigger
1033 for methane ebullition from peatland. *Global Biogeochemical Cycles*, 21(2). <https://doi.org/10.1029/2006GB002790>, 2007.
- 1034 Vachon, D., Langenegger, T., Donis, D., Beaubien, S. E., and McGinnis, D. F.: Methane emission offsets carbon dioxide uptake in a small
1035 productive lake. *Limnology and Oceanography Letters*, 5(6), 384-392, <https://doi.org/10.1002/lol2.10161>, 2020.
- 1036 van Bergen, T.J.H.M., Barros, N., Mendonça, R., Aben, R.C.H., Althuisen, I.H.J., Huszar, V., Lamers, L.P.M., Lürling, M., Roland, F.,
1037 and Kosten, S.: Seasonal and diel variation in greenhouse gas emissions from an urban pond and its major drivers. *Limnol. Oceanogr.* 64
1038 (5), 2129–2139. <https://doi.org/10.1002/lno.11173>, 2019.
- 1039 Varadharajan, C., and Hemond, H. F.: Time-series analysis of high-resolution ebullition fluxes from a stratified, freshwater lake. *Journal*
1040 *of Geophysical Research: Biogeosciences*, 117(G2). <https://doi.org/10.1029/2011JG001866>, 2012.
- 1041 Verpoorter, C., Kutser, T., Seekell, D. A., and Tranvik, L. J.: A global inventory of lakes based on high-resolution satellite imagery.
1042 *Geophysical Research Letters*, 41(18), 6396-6402. <https://doi.org/10.1002/2014GL060641>, 2014.
- 1043 Wang, T., Zhumabieke, M., Zhang, N., Liu, C., Zhong, J., Liao, Q., and Zhang, L.: Variable promotion of algae and macrophyte organic
1044 matter on methanogenesis in anaerobic lake sediment. *Environmental Research*, 237, 116922.
1045 <https://doi.org/10.1016/j.envres.2023.116922>, 2023.

- 1046 Wang, Z., Wang, S., Hu, Y., Du, B., Meng, J., Wu, G., Liu, H., and Zhan, X.: Distinguishing responses of acetoclastic and
1047 hydrogenotrophic methanogens to ammonia stress in mesophilic mixed cultures. *Water Research*, 224, 119029.
1048 <https://doi.org/10.1016/j.watres.2022.119029>, 2022.
- 1049 Wanninkhof, R.: Relationship between gas exchange and wind speed over the ocean. *J. Geophys. Res.* 97, 7373–7381.
1050 <https://doi.org/10.1029/92JC00188>, 1992.
- 1051 Webb, J.R., Leavitt, P.R., Simpson, G.L., Baulch, H.M., Haig, H.A., Hodder, K.R., and Finlay, K.: Regulation of carbon dioxide and
1052 methane in small agricultural reservoirs: optimizing potential for greenhouse gas uptake. *Biogeosciences* 16 (21), 4211–4227.
1053 <https://doi.org/10.5194/bg-16-4211-2019>, 2019.
- 1054 Weiss, R. F., Price, B. A.: Nitrous oxide solubility in water and seawater. *Marine chemistry*, 8(4), 347-359., [doi.org/10.1016/0304-](https://doi.org/10.1016/0304-4203(80)90024-9)
1055 [4203\(80\)90024-9](https://doi.org/10.1016/0304-4203(80)90024-9), 1980.
- 1056 Weiss, R. F.: Determinations of carbon dioxide and methane by dual catalyst flame ionization chromatography and nitrous oxide by
1057 electron capture chromatography. *J. Chromatogr. Sci.* 19, 611–616. <https://doi.org/10.1093/chromsci/19.12.611>, 1981.
- 1058 West, W. E., Coloso, J. J., and Jones, S. E.: Effects of algal and terrestrial carbon on methane production rates and methanogen community
1059 structure in a temperate lake sediment. *Freshw. Biol.* 57, 949–955. <https://doi.org/10.1111/j.1365-2427.2012.02755.x>, 2012.
- 1060 Whiticar, M. J., Faber, E., and Schoell, M.: Biogenic methane formation in marine and freshwater environments: CO₂ reduction vs. acetate
1061 fermentation—isotope evidence. *Geochimica et Cosmochimica Acta*, 50(5), 693-709. [https://doi.org/10.1016/0016-7037\(86\)90346-7](https://doi.org/10.1016/0016-7037(86)90346-7), 1986.
- 1062 Wik, M., Crill, P. M., Varner, R. K., and Bastviken, D.: Multiyear measurements of ebullitive methane flux from three subarctic lakes. *J.*
1063 *Geophys. Res. Biogeosciences* 118:791 1307–1321. <https://doi.org/10.1002/jgrg.20103>, 2013.
- 1064 Wik, M., Thornton, B. F., Bastviken, D., MacIntyre, S., Varner, R. K., and Crill, P. M.: Energy input is primary controller of methane
1065 bubbling in subarctic lakes. *Geophysical Research Letters*, 41(2), 555-560. <https://doi.org/10.1002/2013GL058510>, 2014.
- 1066 Xun, F., Feng, M., Ma, S., Chen, H., Zhang, W., Mao, Z., Zhou, Y., Xiao, Q, Wu, Q. L., and Xing, P.: Methane ebullition fluxes and
1067 temperature sensitivity in a shallow lake. *Science of The Total Environment*, 912, 169589. <https://doi.org/10.1016/j.scitotenv.2023.169589>,
1068 2024.
- 1069 Yan, X., Xu, X., Ji, M., Zhang, Z., Wang, M., Wu, S., Wang, G., Zhang, C., and Liu, H.: Cyanobacteria blooms: A neglected facilitator of
1070 CH₄ production in eutrophic lakes. *Science of the total environment*, 651, 466-474. <https://doi.org/10.1016/j.scitotenv.2018.09.197>, 2019.
- 1071 Yang, Z., Zhao, Y., and Xia, X.: Nitrous oxide emissions from *Phragmites australis*-dominated zones in a shallow lake. *Environmental*
1072 *pollution*, 166, 116-124. <https://doi.org/10.1016/j.envpol.2012.03.006>, 2012.
- 1073 Yentsch, C.S., and Menzel, D.W.: A method for the determination of phytoplankton chlorophyll and phaeophytin by fluorescence. In:
1074 *Deep Sea Research and Oceanographic Abstracts*, 10. Elsevier, pp. 221–231. [https://doi.org/10.1016/0011-7471\(63\)90358-9](https://doi.org/10.1016/0011-7471(63)90358-9), 1963.
- 1075 Zhao, K., Tedford, E. W., Zare, M., and Lawrence, G. A.: Impact of atmospheric pressure variations on methane ebullition and lake
1076 turbidity during ice-cover. *Limnology and Oceanography Letters*, 6(5), 253-261. <https://doi.org/10.1002/lol2.10201>, 2021.