Methane, carbon dioxide, and nitrous oxide emissions from two clear-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 alternative states affect the emission of greenhouse gases (GHGs) 9 such as carbon dioxide (CO₂), methane (CH₄), and nitrous oxide (N₂O) to the atmosphere. We measured the dissolved 10 concentration of CO₂, CH₄, and N₂O from which the diffusive air-water fluxes were computed, in four urban ponds in the 11 city of Brussels (Belgium): two clear-water macrophyte-dominated ponds (Silex and Tenreuken), and two turbid-water 12 phytoplankton-dominated ponds (Leybeek and Pêcheries) on 46 occasions over 2.5 years (between June 2021 and December 13 2023). Ebullitive CH₄ fluxes were measured with bubble traps in the four ponds during deployments in spring, summer, and fall, totalling 48 days of measurements. Measured ancillary variables included water temperature, oxygen saturation level 14 15 (%O₂), concentrations of chlorophyll-a (Chl-a), total suspended matter (TSM), soluble reactive phosphorus (SRP), nitrite 16 (NO₂), nitrate (NO₃), and ammonium (NH₄⁺). The turbid-water and clear-water ponds did not differ significantly in terms of 17 diffusive emissions of CO₂ and N₂O. Clear-water ponds exhibited higher values of ebullitive CH₄ emissions compared to 18 turbid-water ponds, most probably in relation to the delivery of organic matter from macrophytes to sediments, but the 19 diffusive CH₄ emissions were not significantly different between clear- and turbid-water ponds. Across seasons, CH₄ 20 emissions increased with water temperature in all four ponds, with ebullitive CH₄ fluxes having a stronger dependence on water temperature (Q10) than diffusive CH4 fluxes. The temperature sensitivity of ebullitive CH4 fluxes decreased with 21 22 increasing water depth, implying that shallow sediments would respond more strongly to warming (e.g. heat waves). Total 23 annual CH₄ emissions (diffusive+ebullitive) in CO₂ equivalents equalled those of CO₂ in turbid-water ponds and exceeded 24 those of CO₂ in clear-water ponds, while N₂O emissions were negligible compared to the other two GHGs. Total annual 25 GHG emissions in CO₂ equivalents from all four ponds increased from 2022 to 2023 due to higher CO₂ diffusive fluxes, likely driven by higher annual precipitation in 2023 compared to 2022 (leading putatively to higher inputs for organic or 26 27 inorganic carbon from run-off), possibly in response to the intense El Niño event of 2023. The findings of this work suggest that it might be necessary to account for the presence of submerged macrophytes when extrapolating ebullitive CH₄ fluxes in 28 29 ponds at larger scale (regional or global) (particularly if Chl-a is used as a descriptor), although it might be less critical for 30 the extrapolation of diffusive CH₄, CO₂, and N₂O fluxes.

1. Introduction

- 32 Greenhouse gas (GHG) emissions from inland water (rivers, lakes, and reservoirs) to the atmosphere such as carbon dioxide
- 33 (CO₂), methane (CH₄), and nitrous oxide (N₂O) are quantitatively important for global budgets (Lauerwald et al., 2023).
- 34 Global GHG emissions from lakes are lower than from rivers for CO₂ (Raymond et al., 2013) and for N₂O (Lauerwald et al.,
- 35 2019; Maavara et al., 2019). However, reported global emissions of CH₄ from lakes (Rosentreter et al., 2021; Johnson et al.,
- 36 2022) are equivalent or even higher compared to rivers (Stanley et al., 2016; Rocher-Ros et al., 2023). Global emissions of

37 CO₂ and CH₄ from lakes to the atmosphere represent 1.25 to 2.30 Pg CO₂ equivalents (CO₂-eq) annually with a significant 38 proportion from CH₄ emissions, and represent nearly 20% of global CO₂ emissions from fossil fuels (Delsontro et al., 2018). The contribution of CO₂ and CH₄ emissions from small lentic water bodies (small lakes and ponds) can be disproportionately 39 40 high compared to large systems (Holgerson and Raymond, 2016) as small lakes and ponds are the most abundant of all water body types in number (Verpoorter et al., 2014; Cael et al., 2017), and fluxes per m² are usually higher from smaller water 41 42 bodies. The emissions of GHGs from artificial water bodies such as agricultural reservoirs, urban ponds, and storm-water 43 retention basins could be higher than those from natural systems (Martinez-Cruz et al., 2017; Grinham et al., 2018; Herrero 44 Ortega et al., 2019; Gorsky et al., 2019; Ollivier et al., 2019; Peacock et al., 2019, 2021; Webb et al., 2019; Bauduin et al., 45 2024). These higher GHG emissions seem to result from higher external inputs of anthropogenic organic carbon and 46 dissolved inorganic nitrogen (DIN) into artificial systems, but might also reflect other differences compared to natural 47 systems such as in hydrology (Clifford and Heffernan, 2018). Among artificial systems, urban ponds are the subject of a 48 growing body of literature on GHG emissions (Singh et al., 2000; Natchimuthu et al., 2014; van Bergen et al., 2019; Audet 49 et al., 2020; Peacock et al., 2021; Goeckner et al., 2022; Ray and Holgerson, 2023; Ray et al. 2023; Bauduin et al., 2024). 50 Urban areas can have numerous small artificial water bodies mostly associated to green spaces such as public parks, and 51 their number is increasing due to rapid urbanisation worldwide (Brans et al., 2018; Audet et al., 2020; Gorsky et al., 2024; 52 Rabaey et al., 2024). Urban ponds are generally small, shallow, and usually their catchment consists in majority of 53 impervious surfaces with a smaller contribution from soils (Davidson et al., 2015; Peacock et al., 2021). In general, the main 54 function of urban ponds is for storm-water management but they provide additional benefits including aesthetic/recreational 55 amenities and habitats for wildlife (e.g. Tixier et al., 2011; Hassall, 2014).

56 Shallow ponds and lakes occur in two alternative states corresponding to systems with either clear waters (macrophyte-57 dominated) or turbid waters (phytoplankton-dominated), during the productive period of the year (spring and summer in 58 mid-latitudes) (Scheffer et al., 1993). Submerged macrophytes and phytoplankton regulate CO₂ dynamics directly through 59 photosynthesis that can be more or less balanced by community respiration in the water column (e.g., Sand-Jensen and 60 Staehr, 2007). However, it is not clear whether the presence of macrophytes increases or decreases the net CO₂ emissions 61 from ponds and lakes. Some studies have shown a decrease of CO₂ emissions with increasing macrophyte density (Kosten et 62 al., 2010; Ojala et al., 2011; Davidson et al., 2015), but other studies showed the opposite pattern (Theus et al., 2023). In 63 phytoplankton-dominated lakes, CO₂ concentrations depend in part on the developmental stage of phytoplankton, with the 64 growth and peak phases generally coinciding with lower CO₂ concentrations due to photosynthesis (Grasset et al., 2020; 65 Vachon et al., 2020).

CH₄ emissions have been reported to increase with the concentration of chlorophyll-a (Chl-a) in phytoplankton-dominated 66 lakes (DelSontro et al., 2018; Borges et al., 2022). The presence of macrophytes strongly affects the production of CH₄ in 67 freshwaters (Bastviken et al., 2023) and vegetated littoral zones of lakes exhibit higher CH₄ emissions than non-vegetated 68 69 zones (Hyvönen et al., 1998; Huttunen et al., 2003; Juutinen et al., 2003; Desrosiers et al., 2022; Theus et al., 2023). 70 Macrophytes influence organic matter decomposition processes in sediments depending on the quality and quantity of plant 71 matter they release into their environment (Reitsema et al., 2018; Grasset et al., 2019; Harpenslager et al., 2022; Theus et al., 72 2023). Yet, few studies have consistently compared CH₄ emissions in clear-water and turbid-water ponds (Hilt et al., 2017). 73 A study in Argentina reported higher dissolved CH₄ concentrations in clear-water ponds with submerged macrophytes 74 compared to turbid-water phytoplankton-dominated ponds, but no differences in measured CH₄ emissions (Baliña et al., 75 2023).

76 The production of N₂O predominantly occurs through microbial nitrification and denitrification that depend on DIN, O₂ levels, and temperature (Codispoti and Christensen, 1985; Mengis et al., 1997; Velthuis and Veraart, 2022). Competition for 77 DIN between primary producers and N₂O-producing microorganisms can impact N₂O production. Additionally, the transfer 78 79 of labile phytoplankton organic matter to sediments fuels benthic denitrification and impacts N₂O fluxes. Eutrophication is 80 assumed to drive high N₂O emissions from lakes and ponds (Audet et al., 2020; Webb et al., 2021; Wang et al., 2021; Xie et 81 al., 2024) but some lakes with elevated Chl-a concentrations can act as sinks of N₂O due to removal of N₂O by denitrification (Webb et al., 2019; Borges et al., 2022; 2023). The presence of macrophytes also strongly influences nitrogen 82 83 cycling in sediments of lakes and ponds (Barko et al., 1991; Choudhury et al., 2018; Deng et al., 2020; Dan et al., 2021) and 84 should in theory also affect N₂O emissions, although seldom investigated, and available studies provide contradictory 85 conclusions. N2O emissions have been shown to follow diurnal cycles of O2 concentrations in areas dominated by submerged macrophytes in Lake Wuliangsuhai (China) (Ni et al., 2022) and the seasonal cycle of aboveground biomass of 86 87 emerged macrophytes (Phragmites) in Baiyangdian Lake (China) (Yang et al., 2012). On the contrary, a study showed there 88 was no significant difference of N₂O production in sediments of macrophyte-rich (n=10) and macrophyte-free (n=12) lakes 89 in subtropical China (Liu et al., 2018). There has been a very limited number of studies systematically investigating how 90 emissions differ between ponds dominated by phytoplankton and those dominated by macrophytes (Baliña et al., 2023), and 91 none investigating simultaneously CO₂, CH₄, and N₂O emissions including both diffusive and ebullitive components.

92 The emissions of CO₂ and N₂O from aquatic systems are exclusively through diffusion across the air-water interface 93 (diffusive flux), while CH₄ can be additionally emitted as bubbles released from sediments to the atmosphere (ebullitive 94 flux). At annual scale, ebullitive CH₄ flux usually represents more than half of total (diffusive+ebullitive) CH₄ emissions 95 from shallow lakes (Wik et al., 2013; Deemer and Holgerson, 2021), although the relative contribution of ebullitive and 96 diffusive CH₄ emissions is highly variable seasonally (e.g. Wik et al., 2013; Ray and Holgerson, 2023; Rabaey and Cotner 97 2024). Ebullitive CH₄ fluxes are particularly high in the littoral zone of lakes at depths <5 m (Wik et al., 2013; DelSontro et 98 al., 2016; Borges et al., 2022) and strongly increase in response to temperature (DelSontro et al., 2016; Aben et al., 2017; 99 Rabaey and Cotner 2024), as well as organic matter availability (DelSontro et al., 2016; 2018). Ebullitive CH₄ fluxes tend to 100 be higher in small and shallow water bodies (Deemer and Holgerson, 2021) but are notoriously variable in time and space, and are difficult to estimate reliably (DelSontro et al., 2011). 101

Here, we report a dataset of CO₂, CH₄, and N₂O dissolved concentrations in four shallow and small urban ponds (Leybeek, Pêcheries, Silex, and Tenreuken) in the city of Brussels (Belgium) (Fig. 1), with data collected 46 times at regular intervals (between June 2021 and December 2023) on each pond. The air-water diffusive fluxes of CO₂, CH₄, and N₂O were calculated from dissolved concentrations and the gas transfer velocity, while the ebullitive CH₄ fluxes were measured with inverted funnels during 8 deployments (totalling 48 days) in the four ponds. The four ponds have similar depth, surface area, and catchment urban coverage, and mainly differ by the phytoplankton-macrophyte dominance, a clear-water state dominated by macrophytes and a turbid-water state dominated by phytoplankton (alternative states) (Fig. 1). We assess whether the differences in terms of (i) CO₂, CH₄, and N₂O dissolved concentration and diffusive emissions; (ii) ebullitive CH₄ emissions; (iii) relative contribution of CO₂, CH₄, and N₂O to the total GHG emissions in CO₂-eq between the four ponds are explained by the two alternative states.

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112 2. Material and Methods

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2.1. Field sampling and meteorological data

114 Sampling was carried out at a single fixed station (pontoon) in each of the four ponds, on the same day between 9am and 11am, 46 times on each pond between June 2021 and December 2023 at a frequency ranging from one (winter) to three 115 (summer) times per month. Water was sampled 5 cm below the surface with 60 ml polypropylene syringes for analysis of 116 117 dissolved concentrations of CO₂, CH₄, and N₂O. Samples for CH₄ and N₂O were transferred from the syringes with a silicone 118 tube into 60 ml borosilicate serum bottles (Wheaton), preserved with 200 µl of a saturated solution of HgCl₂, sealed with a 119 butyl stopper and crimped with aluminium cap, without a headspace, and stored at ambient temperature in the dark prior to 120 analysis in the laboratory. The partial pressure of CO₂ (pCO₂) was measured directly in the field, within 5 minutes of sample 121 collection, with a Li-Cor Li-840 infrared gas analyser (IRGA) based on the headspace technique with 4 polypropylene 122 syringes (Borges et al., 2019). A volume of 30 ml of sample water was equilibrated with 30 ml of atmospheric air within the 123 syringe by shaking vigorously for 5 minutes. The headspace of each syringe was then sequentially injected into the IRGA 124 and a fifth syringe was used to measure atmospheric CO₂. The final pCO₂ value was computed taking into account the 125 partitioning of CO₂ between water and the headspace, as well as equilibrium with HCO₃ (Dickson et al., 2007) using water 126 temperature measured in-situ and after equilibration, and total alkalinity (data not shown). Samples for total alkalinity were conditioned, stored, and analysed as described by Borges et al. (2019). The IRGA was calibrated in the laboratory with 127 128 ultrapure N₂ and a suite of gas standards (Air Liquide Belgium) with CO₂ mixing ratios of 388, 813, 3788 and 8300 ppm. 129 The precision of pCO₂ measurements was ±2.0%. Water temperature, specific conductivity, and oxygen saturation level (%O₂) were measured in-situ with VWR MU 6100H probe 5 cm below the surface. A 2 liter polyethylene water container 130 was filled with surface water for conditioning the samples for other variables at the laboratory in Université Libre de 131 132 Bruxelles.

133 Three bubble traps were deployed 50 cm apart for measuring ebullitive CH₄ flux. The bubble traps consisted of inverted polypropylene funnels (diameter 23.5 cm) mounted with 60 ml polypropylene syringes, with three way stop valves allowing 134 135 to collect the gas without contamination from ambient air. The polypropylene funnel was attached with steel rods to a polystyrene float. The volume of gas collected in the funnels was sampled with graduated polypropylene 60 ml syringes 136 137 every 24 hours. The value of the collected volume of gas was logged, and the gas was transferred immediately after collection to pre-evacuated 12 ml vials (Exetainers, Labco, UK) that were stored at ambient temperature in the dark prior to 138 139 the analysis of CH₄ concentration in the laboratory. The time-series of measurements was longer at the Silex pond than the 140 other three ponds.

Surveys to identify and quantify visually the relative coverage of emerged and submerged macrophytes were conducted in summer 2023 (Table S1). The resulting list of macrophyte species agreed with past studies in Brussels' ponds (Peretyatko et al., 2007). Air temperature, precipitation, wind speed, and atmospheric pressure, were retrieved from https://wow.meteo.be/en for the meteorological station of the Royal Meteorological Institute of St-Lambert (50.8408°N, 4.4234°E) in Brussels, located between 2.5 and 5.0 km from the surveyed ponds. Air temperature, wind speed and atmospheric pressure were averaged over 24 h to obtain a daily mean value. Precipitation was integrated each day to obtain cumulated daily value.

2.2. Laboratory analysis

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

150 Water was filtered through Whatman GF/F glass microfiber filters (porosity 0.7 µm) with a diameter of 47 mm for total 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 (-151 20 °C). The weight of each filter was determined before and after filtration of a known volume of water using an ExplorerTM 152 Pro EP214C analytical microbalance (accuracy ±0.1 mg) for determination of TSM concentration. Chl-a concentration was 153 154 measured on extracts with 90% acetone by fluorimetry (Kontron model SFM 25) (Yentsch and Menzel, 1963) with a limit of detection of 0.01 µg L⁻¹. Filtered water was stored frozen (-20 °C) in 50 ml polypropylene bottles for analysis of dissolved 155 nutrients. Soluble reactive phosphorus (SRP) was determined by the ammonium molybdate, ascorbic acid and potassium 156 antimony tartrate staining method (Koroleff, 1983), with a limit of detection of 0.1 µmol L⁻¹. Ammonium (NH₄⁺) was 157 determined by the nitroprusside-hypochlorite-phenol staining method (Grasshoff and Johannsen, 1972), with a limit of 158 159 detection of 0.05 µmol L⁻¹. Nitrite (NO₂) and nitrate (NO₃) were determined before and after reduction of NO₃ to NO₂ by a cadmium-copper column, using the Griess acid reagent staining method (Grasshoff et al., 2009), with a detection limit of 160 0.01 and 0.1 µmol L⁻¹, respectively. Concentration of dissolved inorganic nitrogen (DIN) was calculated as the sum of NH₄⁺, 161 162 NO_2^- and NO_3^- concentrations in μ mol L⁻¹.

163 2.2.2. CH₄ and N₂O measurements by gas chromatography

- Measurements of N_2O and CH_4 concentrations dissolved in water were made with the headspace technique (Weiss, 1981) with an headspace volume of 20 ml of ultra-pure N_2 (Air Liquid Belgium) and a gas chromatograph (GC) (SRI 8610C) with a flame ionisation detector for CH_4 and an electron capture detector for N_2O calibrated with $CH_4:N_2O:N_2$ gas mixtures (Air Liquide Belgium) with mixing ratios of 1, 10 and 30 ppm for CH_4 , and 0.2, 2.0 and 6.0 ppm for N_2O . The precision of measurement based on duplicate samples was $\pm 3.9\%$ for CH_4 and $\pm 3.2\%$ for N_2O . Measurements of CH_4 concentration in the gas samples from bubble traps were also made by GC with the same set-up and calibration as for the determination of the dissolved concentrations in water samples.
- The CO₂ concentration is expressed as partial pressure (pCO₂) in parts per million (ppm) and CH₄ as dissolved concentration 171 (nmol L-1), as frequently used in topical literature. CH₄ concentration were systematically and distinctly above saturation 172 level (2-3 nmol L⁻¹) and pCO₂ values were below saturation only five times out of the 187 measurements. The N₂O 173 concentrations fluctuated around atmospheric equilibrium, so data are presented as percent of saturation level (%N₂O, where 174 175 atmospheric equilibrium corresponds to 100%). The equilibrium with atmosphere for N₂O was calculated from the average air mixing ratios of N₂O provided by the Global Monitoring Division (GMD) of the National Oceanic and Atmospheric 176 177 Administration (NOAA) Earth System Research Laboratory (ESRL) (Dutton and Hall, 2023), and using the Henry's 178 constant given by Weiss and Price (1980).

179 **2.3. Calculations**

180 2.3.1. Diffusive GHG emissions

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

$$182 F_G = k \times \Delta[G], (1)$$

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

- 184 The atmospheric pCO₂ was measured in the field with the Li-Cor Li-840. For CH₄, the global average present day
- 185 atmospheric mixing ratio of 1.9 ppm was used (Lan et al., 2024). k was computed from a value normalized to a Schmidt
- number of 600 (k_{600}) and from the Schmidt number of CO₂, CH₄ and N₂O in freshwater according to the algorithms as
- 187 function of water temperature given by Wanninkhof (1992). k₆₀₀ was calculated from the parameterization as a function of
- wind speed of Cole and Caraco (1998). CH₄ and N₂O emissions were converted into CO₂ equivalents (CO₂-eq) considering a
- 189 100-year timeframe, using global warming potentials of 32 and 298 for CH_4 and N_2O , respectively (Myrhe et al., 2013).

190 **2.3.2. Ebullitive flux**

191 Bubble flux (ml m⁻² d⁻¹) was calculated according to:

$$F_{bubble} = \frac{V_g}{A \times \Delta t} \,, \tag{2}$$

- where V_q is the volume of gas collected in the inverted funnel (ml), A is the cross-sectional surface area of the funnel (m²),
- 194 and Δt is the collection time (d).
- 195 A multiple linear regression model of F_{bubble} dependent on water temperature (T_w in °C) and drops of atmospheric pressure
- 196 (Δp in atm) was fitted to the data according to:

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

- where γ is the y-intercept, α and β are the slope coefficients of the multiple linear regression model.
- 199 Δp was calculated according to Zhao et al. (2017):

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

- 201 where p is the atmospheric pressure (atm), p_0 a threshold value fixed at 1 atm, and Δt the time interval between two
- 202 measurements (d) (Fig. S1).
- 203 A linear regression model of F_{bubble} dependent on T_w alone was fitted to the data according to:

$$\log_{10}(F_{bubble}) = \alpha' \times T_w + \gamma', \tag{5}$$

- where α' is the slope coefficient and γ' is the y-intercept.
- To evaluate the relative importance of T_w and Δp combined and T_w alone in driving F_{bubble} , the values modelled based on
- Equations (3) and (5) were compared to the observations in the Silex pond alone or in all four ponds together, for three T_w
- 208 ranges (T_w < 15°C, T_w > 15°C, and the full T_w range).
- 209 Ebullitive CH₄ fluxes (E_{CH4} in mmol m⁻² d⁻¹) were calculated according to:

$$210 \quad E_{CH4} = [CH_4] \times F_{bubble} \,, \tag{6}$$

where $[CH_4]$ is the measured CH₄ concentration in bubbles (mmol ml⁻¹).

- 212 The CH₄ content in bubbles expressed in % of total gas (%CH₄) was fitted with a linear regression model dependent on T_W
- in the Silex pond. The correlation of $%CH_4$ with F_{bubble} was tested on the merged data of all the four ponds.
- 214 A linear regression model of E_{CH4} dependent on T_w was fitted to the data according to:

$$215 \quad \log_{10}(E_{CH4}) = b \times T_w + c \,, \tag{7}$$

- 216 where b is the slope and c is the y-intercept of the linear regression.
- Equation (7) is used to predict E_{CH4} in each pond from measured time-series of T_W allowing matching to each diffusive CH_4
- 218 flux estimate derived from Equation (1).
- The ratio of ebullitive CH_4 flux to total (diffusive + ebullitive) CH_4 flux ($\frac{Ebul}{Tot}$) was fitted as a function of T_W according to
- 220 DelSontro et al. (2016):

$$221 \quad \frac{Ebul}{Tot} = \frac{1}{1 + f \times e^{g \cdot Tw}} \tag{8}$$

- The methane ebullition Q_{10} represents the change in E_{CH4} per 10° C change in T_{W} and was computed according to DelSontro
- 223 et al. (2016):

$$224 Q_{10} = 10^{10b} (9)$$

- 225 where b is given by Equation (7)
- The Q_{10} of diffusive CH₄ fluxes was also computed from equation (9) but using the b value derived by replacing E_{CH4} by the
- 227 diffusive CH₄ flux in equation (7).

228 **2.4. Statistical analysis**

- 229 Generalized linear mixed models (GLMMs) were used (1) to find differences in variables among ponds and (2) to relate
- 230 GHG variables to their putative controls across all ponds and within individual ponds. GLMMs were computed with the
- 231 lme4 package (Bates et al., 2015) in R version 4.4.1 (R Core Team, 2021).
- 232 GLMMs allowed to compare pCO₂, dissolved CH₄ concentration, %N₂O, F_{bubble}, %CH₄ in bubbles, E_{CH4}, and diffusive CH₄
- 233 fluxes among the four ponds, using "sampling date" as a random effect and post-hoc tests were performed using estimated
- 234 marginal means (*emmeans* package in R) to assess pairwise differences between ponds. This analysis aimed at investigating
- 235 if patterns in GHG concentrations and emissions differed among ponds, in particular with regards to clear-water and turbid-
- 236 water states. Data were also compared among the four ponds separated by seasons, but GLMMs did not converge due to
- 237 insufficient number of data points. Comparisons on log₁₀-transformed data were then made using repeated measures analysis
- 238 of variance (ANOVA) (ezANOVA package in R) with Tukey's honestly significant difference (HSD) post-hoc tests. This
- analysis aimed at investigating if patterns in data shown by the data analysis with the full data-set (four seasons merged)
- 240 were also observed when analyzing the data separated by seasons.
- 241 For the data-sets covering the whole sampling period (four seasons merged), GLMMs were constructed for pCO₂, dissolved
- 242 CH₄ concentration, $\%N_2O$, F_{bubble} , $\%CH_4$ in bubbles, E_{CH4} , and diffusive CH₄ fluxes that included T_W , precipitation, $\%O_2$,

243 Chl-a, TSM, DIN, SRP as fixed effects, and "pond" and "sampling date" as a random effect to account for repeated measurements with the *lme4* package in R. This analysis aimed at investigating (1) the impact of photosynthesis-respiration 244 on CO₂ concentrations and emissions based the relationships with Chl-a, DIN, SRP; (2) the impact of the response of 245 methanogenesis to warming on CH₄ concentrations and diffusive and ebullitive emissions based on the relationships to T_W; 246 247 (3) the impact on DIN availability and T_W on N₂O concentrations and diffusive emissions. A linear regression model was 248 used to assess the relationship between Chl-a, TSM, %O₂, SRP, and DIN versus T_W. This analysis aimed at investigating if some of the patterns between GHG variables versus Chl-a, TSM, %O2, SRP, and DIN might in fact have reflected indirectly 249 250 a relation between these variables and T_W. The relationships between the annual means of CH₄, CO₂ and N₂O fluxes and the 251 annual means of a subset of variables (Chl-a, macrophyte cover, pond surface area and depth) were assessed with linear or 252 quadratic regressions. The modelled F_{bubble} in the Silex pond and in the four ponds were compared to measured F_{bubble} values 253 with a linear regression in order to evaluate the model performance. A correlation analysis was used to assess the 254 relationship between % CH₄ and F_{bubble} , as well as between anomalies in annual air temperature and annual precipitation.

The linear regression model and the Pearson correlation coefficient (r) were computed in R using the functions lm and cor(method = "pearson"), respectively. Statistical significance was set at p < 0.05 for all analyses. Significant differences between groups presented in boxplots are indicated by lower-case letters in the Figures.

3. Results

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3.1. Seasonal variations of meteorological conditions and GHG concentrations

The city of Brussels experiences a temperate climate with mild weather year-round, and evenly distributed abundant precipitation totalling on average 837 mm annually for the reference period 1991-2020. The average annual air temperature was 11 °C, with summer average of 17.9 °C and winter average of 4.1 °C for the reference period 1991-2020. During the sampling period, from June 2021 to December 2023, T_w in the surface of the four sampled ponds (Leybeek, Pêcheries, Silex, and Tenreuken; Fig. 1) tracked closely the air temperature that ranged between -1.5 and 30.0°C following the typical seasonal cycle at mid-latitudes in the Northern Hemisphere (Fig. S2). Years 2022 and 2023 were about 1 °C warmer than the average for the period 1991-2020 (11 °C), while year 2021 was closer to the long-term average (Fig. 2). Year 2022 was warmer and drier than 2021 and 2023 (Fig. 2), with positive air temperature anomalies observed evenly throughout the year (9 months out of 12) and negative precipitation anomalies in summer, fall, and early winter (Fig. S2). Year 2021 had warmer and drier months in June and September, colder and wetter months in July and August, and was overall wetter and colder than 2022 (Fig. 2). Year 2023 was marked by both positive air temperature and precipitation anomalies (Fig. S2), resulting in a wetter and warmer year than normal and compared to 2021 and 2022 (Fig. 2). Daily wind speed was generally low (<1 m s⁻¹) except for a 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, corresponding to the Ciarán storm) (Fig. S2).

The four sampled ponds are situated in the periphery of the city of Brussels, with the Silex pond being bordered by the Sonian Forest (Fig. 1). The four ponds are relatively small (0.7-3.2 ha) and shallow (0.6-1.4 m) and have not been drained or dredged since at least 2018 (Table S2). The four studied ponds had significantly different Chl-*a* concentration values during summer, with the Leybeek pond having higher Chl-*a* (78.8±49.5 μg L⁻¹), followed by the Pêcheries pond (19.1±13.7 μg L⁻¹), the Tenreuken pond (3.3±2.4 μg L⁻¹), and the Silex pond (1.0±1.2 μg L⁻¹) (Figs. 1, 3, Table S3). The Leybeek and Pêcheries ponds with higher summer Chl-*a* concentration had turbid-water (summer TSM = 48.7±36.2 and 13.7±10.7 mg L⁻¹, respectively), and undetectable submerged macrophyte cover in summer (Fig. 1, Table S1). The Tenreuken and Silex ponds

- with lower summer Chl-a concentrations had clear-water (summer TSM = 4.9 ± 3.2 and 4.0 ± 3.2 mg L⁻¹, respectively), and a
- 282 high total macrophyte cover during summer (68 and 100%, respectively, Fig. 1, Table S1). Seasonally, the highest values of
- 283 Chl-a were observed in summer in the turbid-water Leybeek and Pêcheries ponds, related to algal blooms. Conversely,
- 284 lowest values of Chl-a were observed in summer in the clear-water Tenreuken and Silex ponds (Figs. 1, 3), probably related
- 285 to competition for dissolved inorganic nutrients with macrophytes.
- 286 The %O₂ values ranged from 11 to 191% (Fig. 3). The highest %O₂ values in the four ponds were observed in spring and
- 287 summer compared to fall and winter owing to aquatic primary production. In summer, %O₂ was significantly higher in the
- Leybeek pond (109±46%) characterized by higher Chl-a concentration compared to the Pêcheries pond (75±23%, p=0.0212,
- Table S3). The lowest average %O₂ was observed in fall in the Pêcheries pond (46±22%) and was significantly lower than in
- 290 the Leybeek (85±34%, p=0.0146, Table S3) and Silex ponds (81±19%, p=0.0130, Table S3).
- 291 The pCO₂ values ranged from 40 to 13,804 ppm (Fig. 3). Undersaturation of CO₂ with respect to atmospheric equilibrium
- 292 (~410 ppm) was only observed on five occasions out of the 187 measurements, three times in the turbid-water Leybeek pond
- 293 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-
- 294 water Tenreuken pond in spring and summer (383 ppm on 13 August 2021 and 55 ppm on 2 May 2022). Low values of
- 295 pCO₂ were generally observed in spring and summer and high values of pCO₂ were observed in fall in the four ponds (Fig.
- 296 3). In summer, pCO₂ was lower in the Leybeek pond (2187±2012 ppm) than in the Pêcheries (3427±1672 ppm, p=0.0015,
- 297 Table S3), and the Silex (3222±1175 ppm, p=0.0002, Table S3) ponds. When data were pooled, pCO₂ was negatively
- 298 influenced by %O₂, and positively by DIN, SRP, and precipitation (Table S4). In individual ponds, pCO₂ was negatively
- 299 influenced by %O₂ and positively by precipitation in the four ponds, positively by DIN in the Leybeek pond, by DIN and
- 300 SRP in the Tenreuken pond, and negatively by Chl-a in the Silex pond (Table S5).
- 301 The CH₄ dissolved concentrations ranged from 194 to 48,380 nmol L⁻¹ (Fig. 3) and was always above saturation (~2 nmol L⁻¹
- 302 ¹). High values of CH₄ dissolved concentrations were generally observed in spring and summer and low values of CH₄
- 303 dissolved concentrations were generally observed in winter in the four ponds (Fig. 3). In summer, CH₄ dissolved
- 304 concentration was higher in the Silex pond (4,898±3,384 nmol L⁻¹) than in the Pêcheries (2,518±2,105 nmol L⁻¹, p=0.0385,
- Table S3) and the Tenreuken (2,189±1,365 nmol L⁻¹, p=0.0055, Table S3) ponds. When data were pooled, dissolved CH₄
- 306 concentration was influenced positively by T_W (Table S4). In individual ponds, CH₄ dissolved concentration was also
- 307 influenced positively by T_W in each of the four ponds (Table S5). Additionally, CH₄ dissolved concentration was positively
- 308 influenced by precipitation in the Leybeek pond and by SRP in the Silex pond, and negatively by DIN in the Pêcheries pond
- 309 and by Chl-a in the Tenreuken and the Silex ponds. (Table S5). These relationships between CH₄ and other variables (SRP,
- 310 DIN, Chl-a) probably indirectly reflected the seasonal variations of these other variables that were also influenced by T_w.
- 311 Indeed, DIN was negatively influenced T_W in the Pêcheries pond; Chl-a was negatively influenced by T_W in the Tenreuken
- 312 and the Silex ponds; SRP was positively influenced by T_W in the Silex pond (Table S6).
- 313 The %N₂O values ranged from 32 to 826% (Fig. 3). Undersaturation of N₂O with respect to atmospheric equilibrium was
- 314 observed 66 times out of the 187 measurements. Low values of %N₂O were generally observed in spring and summer and
- 315 high values of %N₂O were generally observed in fall and winter in the four ponds (Fig. 3). During spring, %N₂O was lower
- 316 in the Pêcheries pond (90±11%) than the Leybeek (138±30%, p=0.0043, Table S3) and the Tenreuken (138±41, p=0.0057,
- 317 Table S3) ponds. During summer, $\%N_2O$ was lower in the Pêcheries pond (78±17%) than the Leybeek (191±104%,
- 318 p<0.0001, Table S3) and the Silex ($126\pm49\%$, p=0.001, Table S3) pond, and lower in the Tenreuken pond ($133\pm106\%$) than
- 319 the Leybeek pond (p=0.0219, Table S3). During fall, %N₂O was lower in the Pêcheries pond (103±33%) than the Leybeek

- 320 pond (190 \pm 70%, p=0.0174, Table S3). For the all sampling period, %N₂O was lower in the Pêcheries pond (94 \pm 28%) than
- 321 the Leybeek ($178\pm82\%$, p<0.0001, Table S7), the Tenreuken ($140\pm77\%$, p<0.0001, Table S7), and the Silex ($144\pm113\%$,
- 322 p<0.0001, Table S7) ponds, and was lower in the Tenreuken pond than the Leybeek pond (p=0.0038, Table S7). When data
- 323 were pooled, $\%N_2O$ was influenced negatively by T_W and positively by DIN and NH_4^+ (Table S4). In individual ponds,
- 324 % N_2O was influenced negatively by T_W in the Leybeek, the Pêcheries, and the Tenreuken ponds (Table S5). % N_2O was
- influenced positively by NO₃ in the Leybeek pond and by NH₄ in the Pêcheries and Tenreuken ponds (Table S8). %N₂O
- 326 was influenced positively by Chl-a and TSM in the Tenreuken pond, and negatively by Chl-a in the Leybeek pond (Table
- 327 S5), probably reflecting the negative influence on Chl-a and TSM by T_W in the Tenreuken pond and the positive influence
- on Chl-a by T_w in the Leybeek pond (Table S6).

3.2. Drivers of bubble flux

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- 330 The F_{bubble} measured with inverted funnels in the four sampled ponds in the city of Brussels ranged between 0 and 2078 ml
- $m^{-2} d^{-1}$ and was influenced positively by T_W in all four systems (Fig. 4). The mean %CH₄ of the bubbles in the four sampled
- ponds in the city of Brussels was 31±21%, and values were influenced positively by T_w in the Silex pond (Fig. 4). The
- 333 %CH₄ of the bubbles was correlated with F_{bubble} (Fig. S3) as both variables were positively influenced by T_W (Fig. 4).
- 334 The time-series at the Silex pond allowed investigating in more detail the effects of T_w and atmospheric pressure variations
- on F_{bubble} (Fig. 5). In spring 2022, the F_{bubble} at the Silex pond increased during drops in atmospheric pressure (depressions)
- 336 (Fig. 5). There was no relation between wind speed and peaks of F_{bubble} ($r^2 = 0.01$, p=0.463), suggesting a more important
- 337 role of changes of atmospheric pressure than of wind speed in triggering bubble fluxes in the Silex pond in spring 2022. The
- 338 F_{bubble} values at the Silex pond was higher in summer (1152±433 mL m⁻² d⁻¹) than during spring (198±170 mL m⁻² d⁻¹) and
- 339 the temporal changes of F_{bubble} tracked those of T_W (Fig. 5). In order to evaluate the relative importance of changes of
- 340 atmospheric pressure and water temperature in triggering bubble fluxes, the F_{bubble} was modelled as function of T_W alone or
- as function of both T_W and Δp (Figs. 5, S4). For periods of low T_W (<15°C), the inclusion of the term of Δp in the model
- 342 improved the performance of the model by comparison to the measurements (Figs. 5, S4). But for warmer periods (>15°C),
- 343 when bubbling fluxes were quantitatively more important, the inclusion in the model of the term of Δp did not improve the
- 344 performance of the model (Figs. 5, S4). For the full T_W range (<15°C and >15°C), the inclusion of the term of Δp only
- improved the performance of the model very marginally (Fig. S4).

346 3.3. Drivers of methane ebullitive fluxes

- 347 The E_{CH4} values in the four ponds ranged between 0 and 59 mmol m⁻² d⁻¹ and were positively related to T_W (Fig. 6). The
- 348 fitted relations between E_{CH4} and T_{W} were specific to each pond and encompassed the fitted relations established in similar
- 349 systems: four small ponds in Québec (DelSontro et al., 2016) and a small urban pond in the Netherlands (Aben et al., 2017).
- 350 The Q₁₀ of CH₄ ebullition values ranged between 4.4 in the deeper Pêcheries pond and 26.9 in the shallower Leybeek pond
- 351 (Table S9). The Q₁₀ of CH₄ ebullition in the four studied ponds of the city of Brussels, in Québec (DelSontro et al., 2016),
- and in the Netherlands (Aben et al., 2017) were negatively related to water depth (Fig. 6).

3.4. Relative contribution of methane ebullitive and diffusive fluxes

- 354 Diffusive CH₄ fluxes computed from dissolved CH₄ concentration and k derived from wind speed ranged between 0.1 and
- 355 19.7 mmol m⁻² d⁻¹ (Fig. 7). The diffusive CH₄ fluxes tended to be higher in summer and spring than in fall and winter owing
- 356 to the strong positive influence on CH₄ dissolved concentration by T_W (Fig. 3; Tables S4, S5). In addition, wind speed only
- showed small seasonal variations during sampling (0.6±0.6m s⁻¹ in spring, 0.3±0.2 m s⁻¹ in summer, 0.7±0.7 m s⁻¹ in fall, and

 $0.6\pm0.2 \text{ m s}^{-1}$ in winter) (Fig. S2). E_{CH4} values were calculated from the relations with T_{W} for each pond given in Figure 6

from the T_W data coincident with the diffusive CH_4 fluxes (Fig. 7). The resulting calculated E_{CH4} values allowed to compare

and integrate seasonally both components of CH₄ emissions to the atmosphere, and to calculate the relative contribution of

361 E_{CH4} to total (diffusive+ebullitive) CH₄ emissions. The relative contribution of E_{CH4} to total CH₄ emissions ranged between 1

and 99% in the four sampled ponds in the city of Brussels (Fig. 7) and was influenced positively by T_W (Fig. S5). The values

of Q_{10} of diffusive CH₄ fluxes were lower than those for E_{CH4} in each pond, and less variable seasonally (1.2 in the Pêcheries

pond to 2.9 in the Silex pond) (Table S9).

365 The annually averaged diffusive and ebullitive fluxes of CH₄ in the four ponds in the city of Brussels were plotted against

annually averaged Chl-a concentration, total macrophyte cover in summer, water depth, and lake surface area (Fig. 8) that

367 are frequent predictors of variations of CH₄ fluxes among lakes and ponds (Holgerson and Raymond, 2016; DelSontro et al.,

2018, Deemer and Holgerson, 2021; Casas-Ruiz et al., 2021; Borges et al., 2022). The annually averaged E_{CH4} values were

significantly higher in the two clear-water ponds (7.3±2.9 and 13.4±3.7 mmol m⁻² d⁻¹ in the Tenreuken and the Silex ponds,

370 respectively) than the two turbid-water ponds (3.8±3.2 and 2.5±1.4 mmol m⁻² d⁻¹ in the Leybeek and the Pêcheries ponds,

371 respectively) (Table S7). The annually averaged E_{CH4} values were significantly higher in the Silex pond, that showed a

higher macrophyte cover during summer (100% in the Silex pond and 68% in the Tenreuken pond), than the Tenreuken pond

373 (p<0.0001, Table S7) and were not significantly different in the two turbid-water Leybeek and Pêcheries ponds (p=0.0617,

Table S7) that showed similar low macrophyte cover during summer (6 and 9% in the Leybeek and Pêcheries ponds,

375 respectively) (Fig. 8). The annually averaged E_{CH4} values were overall influenced positively by macrophyte cover and

376 negatively by Chl-*a* (Fig. 8).

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377 In the four sampled urban ponds, annually averaged CH₄ diffusive fluxes were higher in the pond with the highest total

378 macrophyte cover in the clear-water ponds, and higher in the pond with the highest Chl-a concentration in the turbid-water

379 ponds (Fig. 8). The annually averaged relative contribution of E_{CH4} to total CH_4 emissions were higher in the two clear-water

380 ponds than the two turbid-water ponds (Table S7). The relative contribution of E_{CH4} to the total CH₄ flux was influenced

positively by macrophyte cover and negatively by Chl-a (Fig. 8).

382 The annually averaged diffusive fluxes of CO₂ (F_{CO2}) and N₂O (F_{N2O}) in the four ponds in the city of Brussels were also

383 plotted against annually averaged Chl-a concentration, total macrophyte cover in summer, water depth, and lake surface area

384 (Fig. S6). Annually averaged F_{CO2} were lower in the Leybeek pond than the Pêcheries and the Silex ponds (Table S7). F_{CO2}

385 was not significantly influenced by the other variables (Chl-a concentration, total macrophyte cover, water depth, and lake

surface area) (Fig. S6). Annually averaged F_{N2O} was not significantly different between clear-water and turbid-water ponds.

387 F_{N2O} was significantly lower in the deeper Pêcheries pond than the two shallower Leybeek and Silex ponds (Table S7), and

 F_{N2O} showed a significant negative relationship with water depth (Fig. S6).

3.5. Relative contribution of CO₂, CH₄ and N₂O emissions

390 The emissions in CO₂-eq for the 3 GHGs averaged per season for both 2022 and 2023 peaked seasonally in summer in the

391 Silex (2.9 mg CO_2 -eq m⁻² d⁻¹), the Tenreuken (1.7 mg CO_2 -eq m⁻² d⁻¹), and the Leybeek (1.1 mg CO_2 -eq m⁻² d⁻¹) pond (Fig.

392 9), but peaked in fall in the Pêcheries pond (1.3 mg CO_2 -eq m^{-2} d^{-1}). The higher value of the total GHG emissions in fall

393 compared to other seasons in the Pêcheries pond was due to an increase of CO₂ emissions in fall that surpassed the peak of

394 CH₄ emissions in summer. The GHG fluxes were lowest in winter in the Silex (1.3 9 mg CO₂-eq m⁻² d⁻¹), the Tenreuken (0.9

395 mg CO₂-eq m⁻² d⁻¹), the Pêcheries (0.8 mg CO₂-eq m⁻² d⁻¹), and the Leybeek (0.6 mg CO₂-eq m⁻² d⁻¹) ponds. The relative

- 396 contribution of E_{CH4} peaked in summer in the Silex (73.8%), the Tenreuken (70.9%), the Pêcheries (23.6%), and the Leybeek
- 397 (58.3%) ponds. The relative contribution of E_{CH4} was lowest in winter in the Silex (22.1%), the Tenreuken (10.0%), the
- 398 Pêcheries (6.7%), and the Leybeek (1.0%) ponds.
- 399 The annual emissions in CO₂-eq of the three GHGs (CO₂, CH₄, and N₂O) in 2022 and 2023 were higher in the two clear-
- water ponds (1.3±0.5 and 1.8±0.9 mg CO₂-eq m⁻² d⁻¹ in the Tenreuken and Silex ponds, respectively) than in the two turbid-
- water ponds (1.0±0.2 and 0.9±0.5 mg CO₂-eq m⁻² d⁻¹ in the Leybeek and Pêcheries ponds, respectively) (Fig. 9) due to higher
- 402 total CH₄ emissions (diffusive+ebullitive) in clear-water ponds (0.7±0.4 and 1.2±0.5 mg CO₂-eq m⁻² d⁻¹ in the Tenreuken and
- Silex ponds, respectively) than in turbid-water ponds (0.2±0.2 and 0.4±0.3 mg CO₂-eq m⁻² d⁻¹ in the Leybeek and Pêcheries
- 404 ponds, respectively). The contribution of N₂O to the total GHG emissions was marginal and did not affect the differences in
- 405 total GHG fluxes between ponds, with the highest contribution observed in the Leybeek pond, with a contribution of 1.7%.
- 406 The majority of GHG emissions in CO₂-eq was related to CO₂ and CH₄ (diffusive+ebullitive) in the four ponds. In turbid-
- 407 water ponds CO₂ represented the largest fraction of GHG emissions (68.5% (2022) and 79.3% (2023) in the Pêcheries pond,
- 408 and 49.0% (2022) and 58.3% (2023) in the Leybeek pond). In clear-water ponds CH₄ represented the largest fraction of
- 409 GHG emissions (66.5% (2022) and 63.3% (2023) in the Silex pond, and 60.8% (2022) and 50.0% (2023) in the Tenreuken
- 410 pond). The higher annual GHG emissions in CO₂-eq from the two clear-water ponds than from the turbid-water ponds were
- 411 related to the higher contribution of E_{CH4} .
- 412 The annual GHG fluxes increased from 2022 to 2023 due to an increase in the relative contribution of CO₂ diffusive
- emissions 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
- 414 mg CO₂ m⁻² d⁻¹ in 2023. Diffusive CO₂ emissions were 2.1 times higher in summer 2023 than in summer 2022, and 2.5 times
- 415 higher in fall 2023 than in fall 2022, and showed similar values between 2023 and 2022 in spring and winter (1.1 higher and
- 416 1.1 lower, respectively).

417 4. Discussion

- 418 The Leybeek and Pêcheries ponds are turbid-water systems (high Chl-a and TSM values, low submerged macrophyte cover)
- and the Tenreuken and Silex ponds are clear-water systems (low Chl-a and TSM values, high submerged macrophyte cover)
- 420 (Figs. 1, 3). All four ponds have a relatively similar size (0.7 to 3.2 ha) and depth (0.5 to 1.4 m) and are uniformly located in
- 421 an urban landscape in the city of Brussels. It can be assumed that, among the four systems, the major difference that is
- 422 expected to affect GHG emissions is the dominance of aquatic primary producer, either phytoplankton or macrophytes,
- 423 corresponding to two alternative states sensu Scheffer et al. (1993). Our data-set provides the opportunity to investigate the
- 424 effect of the two alternative states on GHG emissions from small lentic systems.
- The reported pCO₂ values (40 to 13,804 ppm) (Fig. 3) in the four ponds in the city of Brussels were within the range of
- values typically observed in ponds (Holgerson and Raymond, 2016; Peacock et al., 2019; Audet et al., 2020) (Fig. 3). The
- 427 pCO₂ values were influenced negatively by %O₂ and positively by DIN and SRP across seasons (Tables S4, S5) showing
- 428 that their seasonal variability was driven by aquatic primary production and degradation of organic matter (e.g. Holgerson
- 429 2015). Accordingly, low values of pCO₂ were generally observed in spring and summer probably due to uptake of CO₂ by
- 430 primary production from either phytoplankton or submerged macrophytes. High values of pCO₂ were observed in fall in the
- 431 four ponds and probably reflect the release of CO₂ from degradation of organic matter due to the senescence of
- 432 phytoplankton or macrophytes (Fig. 3). In all four ponds, pCO₂ values were influenced positively by precipitation (Tables

433 S4, S5) suggesting an additional control of external inputs of carbon either as organic carbon sustaining internal degradation of organic matter or as soil CO₂ (e.g. Marotta et al., 2010; Ojala et al., 2011; Rasilo et al., 2012; Vachon and del Giorgio, 434 435 2014; Holgerson, 2015). The %N₂O values (32 to 826%) (Fig. 3) in the four ponds were within the range of values typically observed in ponds (Audet et al., 2020; Rabaey and Cotner, 2022). When all the data were pooled, the %N₂O was influenced 436 437 positively by DIN (Table S4) as also frequently reported by other studies in ponds and interpreted as a control of nitrification 438 and/or denitrification (hence N₂O production) by DIN levels (Audet et al., 2020; Webb et al., 2021; Wang et al., 2021; Xie 439 et al., 2024). The negative influence on %N₂O by T_W (Table S4) might reflect the effect of the inhibition at low temperatures 440 of the final step of denitrification leading to an accumulation of N₂O (Velthuis and Veraart, 2022) but could also indirectly 441 result from the higher DIN concentrations observed at low Tw values (Table S6). The CH4 dissolved concentrations (194 to 48,380 nmol L⁻¹) (Fig. 3) in the four ponds were within the range of values typically observed in ponds (Natchimuthu et al., 442 443 2014; Holgerson and Raymond, 2016; Peacock et al., 2019; Audet et al., 2020; Rabaey and Cotner, 2022; Ray et al., 2023), 444 and were influenced positively by Tw in all four ponds individually and when pooled (Tables S4, S5), most probably 445 reflecting the increase of sedimentary methanogenesis with temperature (Schulz and Conrad, 1996).

446 Temperature also exerted a strong control on bubble flux from sediments and ebullitive CH₄ emissions. The F_{bubble} values (0 and 2078 ml m⁻² d⁻¹) in the four sampled ponds (Fig. 4) were within the range of values reported in lentic systems of 447 equivalent size by Wik et al. (2013) (0 to 2772 mL m⁻² d⁻¹), Delsontro et al. (2016) (11 to 748 mL m⁻² d⁻¹), and Ray and 448 Holgerson (2023) (0 to 2079 mL m⁻² d⁻¹). The F_{bubble} was influenced positively by T_W (Fig. 4) in agreement with previous 449 450 studies (e.g. Wik et al., 2013; DelSontro et al., 2016; Aben et al., 2017; Ray and Holgerson, 2023). Bubbling events from 451 lake sediments are known to also be triggered by a decrease of hydrostatic pressure on the sediments due to water level 452 fluctuations or drops in atmospheric pressure (Tokida et al., 2007; Scandella et al., 2011; Varadharajan and Hemond, 2012; 453 Wik et al., 2013; Taoka et al., 2020; Zhao et al., 2021). In the Silex pond, in spring 2022, some peaks in F_{bubble} were related 454 to drops in atmospheric pressure (Fig. 5) but unrelated to wind speed as shown in Gatun Lake (Keller and Stallard, 1994). A 455 statistical model of F_{bubble} that included the contributions of T_W and Δp was used to quantify the relative importance of each 456 of these two drivers (Fig. S4) and showed that air pressure drop seemed quantitatively important only at low T_W and that the 457 intensity of bubble flux was mainly driven by temperature change at yearly scales, in agreement with previous studies (e.g. Wik et al., 2013; DelSontro et al., 2016; Aben et al., 2017; Ray and Holgerson, 2023). 458

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The mean %CH₄ of the bubbles (31±21%) in the four sampled ponds in the city of Brussels was comparable to the values obtained by Wik et al. (2013) (35±25%), Delsontro et al. (2016) (58±25%), and Ray and Holgerson (2023) (25±13%) in lentic systems of similar size. The increasing pattern of %CH₄ of the bubbles with T_W (Fig. 4) was most probably related to the strong dependence of methanogenesis on temperature (Schulz and Conrad, 1996). The increase of methanogenesis with temperature leads to the build-up of gas bubbles in sediments that are richer in CH₄, and consequently to higher bubble fluxes with a higher CH₄ content at higher temperatures (Figs. 4, S3). Since both F_{bubble} and the %CH₄ of the bubbles increased with T_W (Fig. 4), E_{CH^4} in the four ponds were also positively related to T_W (Fig. 6) as shown previously in other small lentic systems (*e.g.* Wik et al., 2013; DelSontro et al., 2016; Natchimuthu et al., 2016; Aben et al., 2017; Ray and Holgerson, 2023; Rabaey and Cotner, 2024). Yet, the dependency of CH₄ ebullition on temperature (Q₁₀) was different among the four ponds and was negatively related to depth including data from systems in Québec (DelSontro et al., 2016) and The Netherlands (Aben et al., 2017) (Fig. 6). This implies that an increase in T_W leads to a smaller increase in CH₄ ebullitive fluxes (lower Q₁₀) in deeper ponds as the impact of hydrostatic pressure on sediments is higher in deeper ponds compared to shallow ponds, restricting bubble formation and release (*e.g.* DelSontro et al., 2016). This dependence of Q₁₀ of

- 472 CH₄ ebullition to depth suggests that the response of CH₄ ebullition to heatwaves (or longer-term warming) might be more
- 473 intense the shallower the pond, in addition to other effects from heat-waves on GHG emissions (e.g. Audet et al., 2017).
- The values of Q₁₀ for diffusive CH₄ fluxes in the four ponds were lower than those for ebullitive CH₄ fluxes (Table S9) as
- 475 reported by other studies in lentic systems (DelSontro et al., 2016; Xun et al., 2024). The lower dependence to T_W of
- 476 diffusive CH₄ fluxes compared to ebullitive CH₄ fluxes might be related to a lower relative change of CH₄ concentrations
- 477 and k with the variation of T_W. Dissolved CH₄ concentrations in surface waters of lentic systems are strongly affected by
- 478 microbial methane oxidation (e.g. Bastviken et al., 2002). A relative increase of methanogenesis in sediments might lead to a
- 479 stronger increase of CH₄ emission by ebullition than by diffusion because of a reduction of CH₄ diffusive emissions resulting
- 480 from microbial methane oxidation. Additionally, k depends on wind speed, but in the four ponds, the warmer periods of the
- 481 year (summer) tended to be less windy (~0.3 m s⁻¹) than the other seasons (>0.6 m s⁻¹) also contributing to a lower
- 482 dependence on T_W of CH₄ diffusive fluxes compared to ebullitive fluxes (lower Q₁₀ values).
- 483 The difference in the Q₁₀ of diffusive and ebullitive CH₄ fluxes was consistent with a variable contribution of the diffusive
- 484 and ebullitive CH₄ fluxes seasonally as a function of T_W . The contribution of E_{CH4} to total (diffusive+ebullitive) CH₄
- 485 emissions strongly increased with T_W in the four ponds (Fig. S5). At annual scale, E_{CH4} represented between 55% and 83%
- 486 of the total CH₄ emissions in the Leybeek and Silex ponds, respectively. This finding is consistent with other studies
- showing that ebullitive CH₄ fluxes can account for more than half of total CH₄ emissions in small and shallow lentic systems
- 488 (e.g. Wik et al., 2013; Deemer and Holgerson, 2021; Ray and Holgerson, 2023; Rabaey and Cotner, 2024). The averaged
- 489 E_{CH4} values were higher in the two clear-water ponds (10.4 mmol m⁻² d⁻¹) than the two turbid-water ponds (3.2 mmol m⁻² d⁻¹)
- 490 (Fig 7). The averaged E_{CH4} values in the four ponds were influenced positively by macrophyte cover and negatively by Chl-a
- 491 (Fig. 8). The higher E_{CH4} values from the two clear-water ponds would suggest that the delivery of organic matter to
- 492 sediments from macrophytes sustained a quantitatively larger methane production than from phytoplankton. This finding is
- 493 consistent with the notion that vegetated littoral zones of lakes are hot spots of CH₄ production and emission (e.g. Hyvönen
- 494 et al., 1998; Huttunen et al., 2003; Juutinen et al., 2003; Desrosiers et al., 2022). CH₄ fluxes in lentic systems have been
- 495 extrapolated at global scale assuming a dependency on aquatic productivity using Chl-a as a predictor (e.g. DelSontro et al.,
- 496 2018). The negative relation between E_{CH4} values with Chl-a (Fig. 8) shows that Chl-a concentration alone fails to predict
- 497 ebullitive fluxes in macrophyte-dominated clear-water ponds.
- 498 The annually averaged diffusive CH₄ emissions in the four ponds seemed to respond positively to both increasing
- 499 phytoplankton and macrophyte biomass resulting in a U-shaped relation between diffusive CH₄ emissions and Chl-a as well
- 500 as macrophyte cover (Fig. 8). Higher values of annually averaged CH₄ diffusive fluxes occurred at the extreme values of
- 501 Chl-a or of macrophyte cover (minimum or maximum), and lower values occurred at the intermediate values of Chl-a or
- 502 macrophyte cover. Such U-shape relation resulted from the inverse relationship between macrophyte cover and Chl-a
- 503 (alternative states) and is consistent with reported positive relation between diffusive CH₄ fluxes with both macrophyte cover
- 504 (e.g. Ray et al., 2023; Theus et al., 2023) as well as with phytoplankton biomass (e.g. DelSontro et al., 2018; Yan et al.,
- 505 2019; Bartosiewicz et al., 2021). The relative contribution of E_{CH4} to the total annual CH₄ flux increased with the macrophyte
- 506 cover (Fig. 8), in agreement with the idea of an increase of CH₄ ebullition relative to diffusive CH₄ emissions in vegetated
- 507 sediments compared to unvegetated sediments (e.g. Desrosiers et al., 2022; Ray et al., 2023; Theus et al., 2023).
- 508 Fluxes of CH₄ and CO₂ have been reported to be negatively related to surface area and depth by numerous studies in ponds
- 509 (e.g. Holgerson, 2015; Holgerson and Raymond, 2016; Ray et al., 2023; Theus et al., 2023) and lakes (e.g. Kankaala et al.,
- 510 2013; DelSontro et al., 2018, Deemer and Holgerson, 2021; Casas-Ruiz et al., 2021; Borges et al., 2022). Annual diffusive

F_{CH4} and F_{CO2} were both unrelated to surface area and depth in the four studied ponds (Figs. 8, S6) resulting from the narrow range of variation of water depth (0.6 to 1.4 m) and surface area (0.7 to 3.2 ha). The lack of relationship between annual F_{CO2} and both Chl-*a* and macrophyte cover in the four ponds (Fig. S6) might be surprising since other studies have reported lower CO₂ fluxes in more productive lentic systems (*e.g.* Sand-Jensen and Staehr, 2007; Borges et al., 2022). We hypothesize that given that the four systems were either phytoplankton-dominated or macrophyte-dominated (alternative states), the ponds had an important submerged productivity, in both cases, resulting in a relatively invariant F_{CO2} as function of either Chl-*a* or macrophyte cover.

Global average emissions of GHGs in CO_2 -eq from inland waters are dominated by CO_2 followed by CH_4 with a small contribution from N_2O according to Lauerwald et al. (2023). However, in small lentic systems such as ponds, the CO_2 -eq emissions from CH_4 can match or dominate those of CO_2 (*e.g.* Webb et al., 2023; Ray and Holgerson, 2023; Rabaey and Cotner, 2024). The meta-analysis of Holgerson and Raymond (2016) suggested that the CO_2 and CH_4 emissions in CO_2 -eq are numerically close in small lentic systems such as ponds but become increasingly dominated by CO_2 emissions in larger lentic systems. In the four studied ponds, the GHG emissions in CO_2 -eq were dominated by CO_2 and CH_4 with a marginal contribution (<2%) from N_2O (Fig. 9). Annually, CO_2 represented the largest fraction of GHG emissions in CO_2 -eq (~60%) in turbid-water ponds (Leybeek and Pêcheries), while CH_4 represented the largest fraction of GHG emissions in CO_2 -eq (~60%) in clear-water ponds (Silex and Tenreuken) as a result of higher E_{CH4} values in the clear-water ponds (Fig. 7).

The annual GHG emissions in CO₂-eq increased from 2022 to 2023 due to an increase in the relative contribution of CO₂ 527 528 diffusive emissions in all four ponds (Fig. 9) as a result of higher precipitation in 2023 (Fig. 2). Air temperatures were 529 similar in both years (annual average of 12.2°C in 2022 and 12.1°C in 2023), and precipitation was 1.5 times higher in 2023 than in 2022. Higher precipitation is likely to increase the inputs of organic and inorganic carbon from soils to ponds, as 530 previously shown in other lentic systems (e.g. Marotta et al., 2011; Ojala et al., 2011; Rasilo et al., 2012; Vachon and del 531 532 Giorgio, 2014; Holgerson, 2015). This hypothesis is only based on the comparison of two years but the increase of the 533 relative contribution of CO₂ diffusive emissions in 2023 was observed in all four ponds. The synchronicity of the increase of CO₂ diffusive emissions in 2023 compared to 2022 suggests a common uniform driver in all four ponds that would be 534 535 consistent with a large variation in weather between the two years such as annual precipitation. The strong El Niño event in 536 2023 induced low-level cyclonic wind anomalies and higher precipitation over Western Europe, including Belgium (Chen et 537 al., 2024).

5. Conclusions

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539 Ebullitive CH₄ emissions in 2022-2023 were higher in the two clear-water, macrophyte-dominated ponds (Tenreuken and 540 Silex) than in the two turbid-water, phytoplankton-dominated ponds (Pêcheries and Leybeek) of the city of Brussels, 541 although, the diffusive CH₄ fluxes were not significantly different between the clear-water ponds and the turbid-water ponds. 542 The annually averaged diffusive N₂O and CO₂ fluxes were not significantly different in the two clear-water ponds from those 543 in the two turbid-water ponds. Other studies have found no difference in N₂O sedimentary production in lakes with high and 544 low density of submerged macrophytes. We hypothesize that CO₂ fluxes were relatively invariant among the four sampled ponds because of their similar size, depth, and putatively productivity (either from phytoplankton or submerged 545 546 macrophytes). The total (diffusive and ebullitive) CH₄ emissions represented 58% of total annual GHG emissions in CO₂-eq 547 in the two clear-water ponds compared to 41% in the two turbid-water ponds. CO₂ represented nearly all the remainder of 548 total annual GHG emissions in CO₂-eq, and N₂O represented a very marginal fraction (<2%).

The seasonal variations of GHG emissions were mainly driven by CH_4 ebullitive emissions that peaked in summer (both quantitatively and relatively), as CH_4 ebullition was related positively to T_W resulting from an increase in both flux of bubbles and CH_4 content of bubbles with warming. The pCO_2 values in the four sampled ponds increased with precipitation at seasonal scale, probably in relation to higher inputs of organic and inorganic carbon from soils. Years 2022 and 2023 were abnormally dry and wet, respectively. The GHG emissions were higher in 2023 than 2022 mainly due to an increase in the relative contribution of CO_2 emissions probably due to higher precipitation in response to a strong El Niño event of 2023. This would suggest that variations of precipitation also affected year-to-year variations of CO_2 emissions in addition to partly regulating seasonal variations of CO_2 emissions from the four studied ponds.

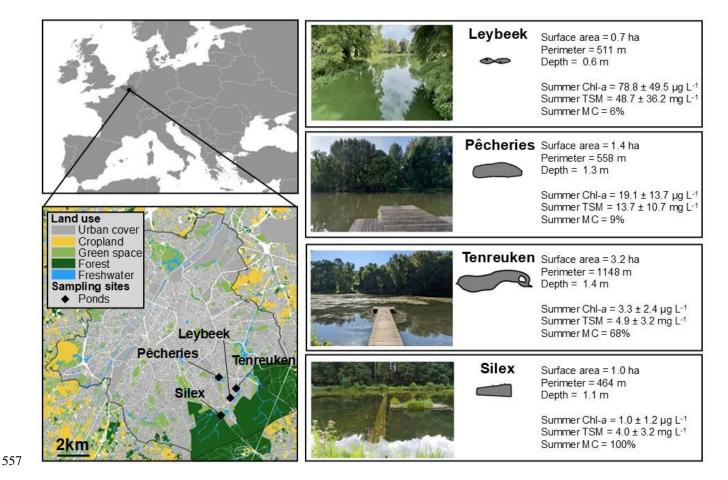


Figure 1: Location of the four sampled urban ponds (black diamonds) in city of Brussels (Belgium) delineated by the black line. Right panels indicate for each pond the shape of the ponds, surface area (ha), perimeter (m), average depth (m), mean±standard deviation of chlorophyll-a (Chl-a, in µg L⁻¹) and total suspended matter (TSM, in mg L⁻¹) in summer (21 June to 21 September in 2021, 2022, 2023), and summer total macrophyte cover (MC, in %) (Table S1).

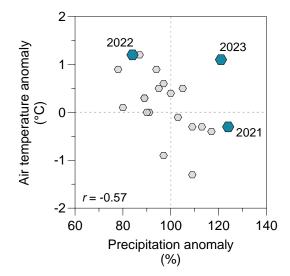


Figure 2: Anomaly of annual air temperature (°C) as a function of anomaly of annual precipitation (%) from 2003 to 2023 with respect to average of the 1991-2020 period (11 °C and 837 mm, respectively). Small grey hexagons represent anomaly values for years 2003-2020 and larger blue hexagons represent anomaly values for years 2021-2023. The air temperature anomaly was correlated to the precipitation anomaly for years 2003-2020 (Pearson r = -0.57, p = 0.0147, n = 18). Note the anomalous rainy year in 2023 relative to the pattern of precipitation as function of temperature anomalies for the other years, possibly in response to the strong El Niño event of 2023 (Chen et al., 2024).

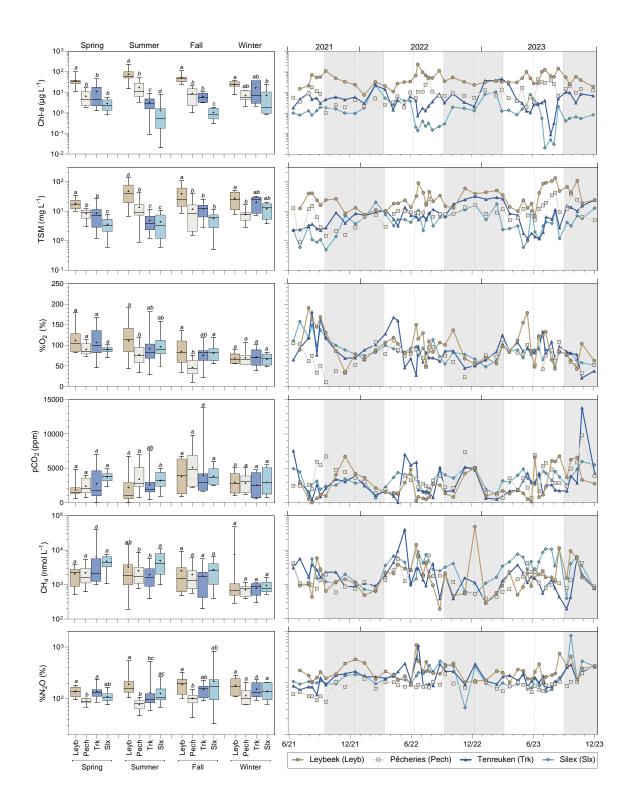


Figure 3: Seasonal variations of Chlorophyll-a (Chl-a, in μg L⁻¹), total suspended matter (TSM, in mg L⁻¹), oxygen saturation (%O₂, in %), partial pressure of CO₂ (pCO₂ in ppm), dissolved CH₄ concentration (CH₄, in nmol L⁻¹), and N₂O saturation level (%N₂O, in %) in four urban ponds (Leybeek (Leyb), Pêcheries (Pech), Tenreuken (Trk), and Silex (Slx)) in the city of Brussels (Belgium) from June 2021 to December 2023. Box plots show median (horizontal line), mean (cross), and 25–75% percentiles (box limits). Whiskers extend from minimum to maximum values. Grey and white bands in the plots on the right correspond to the autumn/winter and spring/summer periods, respectively, and dotted vertical bars indicate the first day of each season. Lower case letters indicate significant differences between ponds (Tables S3 and S4).

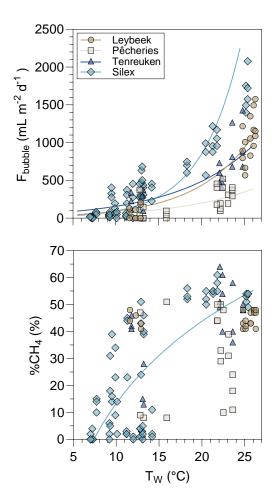


Figure 4: Bubble flux $(F_{bubble}$ in ml m⁻² d⁻¹) and the relative CH_4 content in bubbles (%CH₄, in %) as a function of surface water temperature $(T_w \text{ in }^{\circ}C)$ in four urban ponds (Leybeek, Pêcheries, Tenreuken, and Silex) in the city of Brussels (Belgium) from June 2021 to December 2023. Bubbles fluxes were measured with three bubble traps in spring, summer, and fall of 2022 and 2023, totalling 8 days in the Leybeek, Pêcheries, and Tenreuken ponds and 24 days in the Silex pond. Given the shallowness of the sampled systems (<1.5 m, Fig. 1), we assumed that sediments experience the same temperature as surface waters. In the upper plot, solid lines represent linear regression of $log_{10}(F_{bubble})$ as a function of T_W for the Leybeek $(log_{10}(F_{bubble}) = 0.0664 \text{ x } T_W + 1.3095, n=22, p<0.0001)$, the Pêcheries $(log_{10}(F_{bubble}) = 0.0486 \text{ x } T_W + 1.3257, n=22, p=0.0146)$, the Tenreuken $(log_{10}(F_{bubble}) = 0.0492 \text{ x } T_W + 1.7039, n=19, p<0.0001)$, and the Silex $(log_{10}(F_{bubble}) = 0.0945 \text{ x } T_W + 1.0373, n=72, p<0.0001)$ ponds. In the lower plot, solid line represents the linear regression of %CH₄ as a function of $log_{10}(T_W)$ for the Silex pond (%CH₄=101.11 × $log_{10}(T_W)$ - 87.8, n=72, p<0.0001).

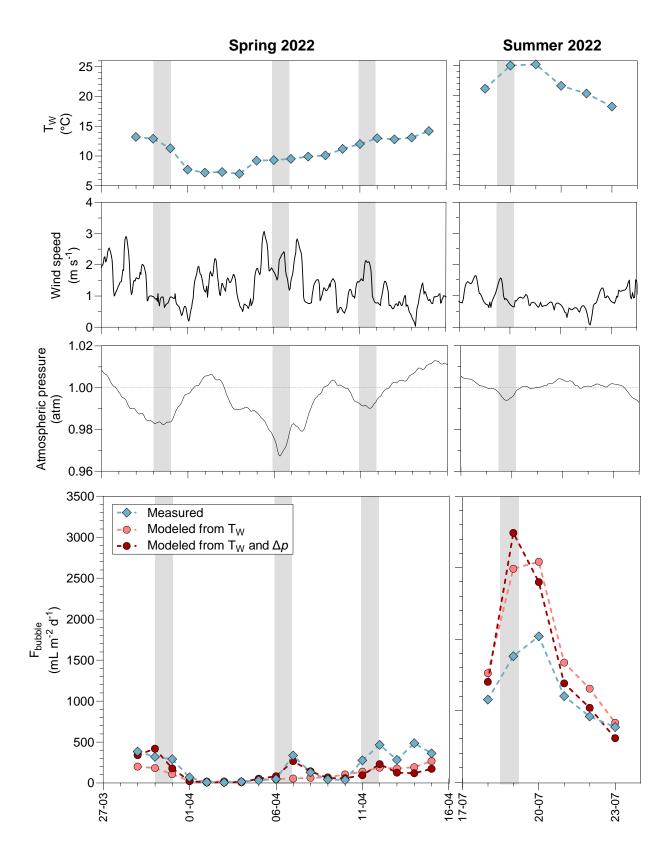
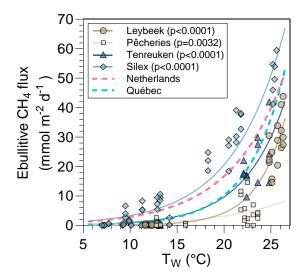
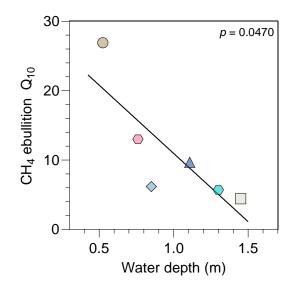


Figure 5: Time-series of surface water temperature (T_W , $^{\circ}$ C), wind speed (m s⁻¹), atmospheric pressure (atm), measured and modeled bubble flux (F_{bubble} in ml m⁻² d⁻¹) in the Silex pond from the 29 March 2022 to the 15 April 2022 and from the 18 July 2022 to the 23 July 2022. The bubble flux was modelled from a fit to data based on Tw alone ($\log_{10}(F_{bubble}) = 3.973 \times \log_{10}(T_w) - 2.15$, p<0.0001, n=72) and based on both T_W and drops in atmospheric pressure (Δp) ($\log_{10}(F_{bubble}) = 4.551 \times \log_{10}(T_w) + 1.962 \times \Delta p - 3.006$, p < 0.0001, n = 72).





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Figure 6: Measured ebullitive CH₄ fluxes (mmol m⁻² d⁻¹) as function of surface water temperature (°C) in four urban ponds (Leybeek, Pêcheries, Tenreuken, and Silex) in the city of Brussels (Belgium), in spring, summer, and fall of 2022 and 2023, totalling 8 days in the Leybeek, Pêcheries, and Tenreuken ponds and 24 days in the Silex pond, with three bubble traps. Solid lines represent exponential fit for the Leybeek ($Y = 0.01 \cdot e^{0.32 \cdot X}$, n=22), Pêcheries ($Y = 0.16 \cdot e^{0.15 \cdot X}$, n=22), Tenreuken ($Y = 0.10 \cdot e^{0.23 \cdot X}$, n=19), Silex ($Y = 0.54 \cdot e^{0.18 \cdot X}$, n=72) ponds (Table S7). Dashed lines represent published exponential fit established in similar systems: four 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 ($Y = 0.51 \cdot e^{0.17 \cdot X}$) (Aben et al., 2017). Each exponential curve allows to determine a Q₁₀ of CH₄ ebullition, plotted against water depth; solid line represents linear regression ($Y = 30.64 - 19.67 \cdot X$, n = 6).

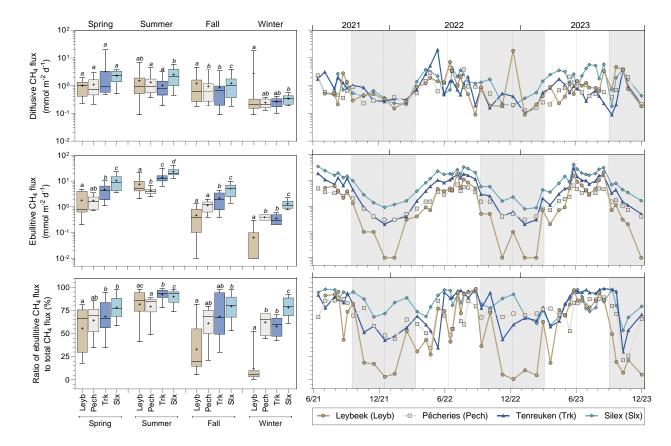


Figure 7: Seasonal variations of diffusive and ebullitive CH₄ fluxes (mmol m⁻² d⁻¹), and the ratio of ebullitive CH₄ flux to total (ebullitve+diffusive) CH₄ flux (%) in four urban ponds (Leybeek (Leyb), Pêcheries (Pech), Tenreuken (Trk), and Silex (Slx)) in the city of Brussels (Belgium) from June 2021 to December 2023. Diffusive fluxes were calculated from CH₄ concentration and gas transfer velocity derived from wind speed. Ebullitive CH₄ fluxes were calculated from the relations with water temperature for each pond (Fig. 6; Table S7) from the water temperature data coincident with the diffusive CH₄ fluxes. Note that the relations of ebullitive CH₄ fluxes as a function of water temperature were established over a temperature range (7.0 to 26.3°C) that is consistent with the range of water temperature values (2.0-25.9°C) over which the ebullitive CH₄ fluxes were modelled. Box plots show median (horizontal line), mean (cross), and 25–75% percentiles (box limits). Whiskers extend from minimum to maximum values. Grey and white bands in the plots on the right correspond to the autumn/winter and spring/summer periods, respectively, and dotted vertical bars indicate the first day of each season. Lower case letters indicate significant differences between ponds (Tables S3 and S4).

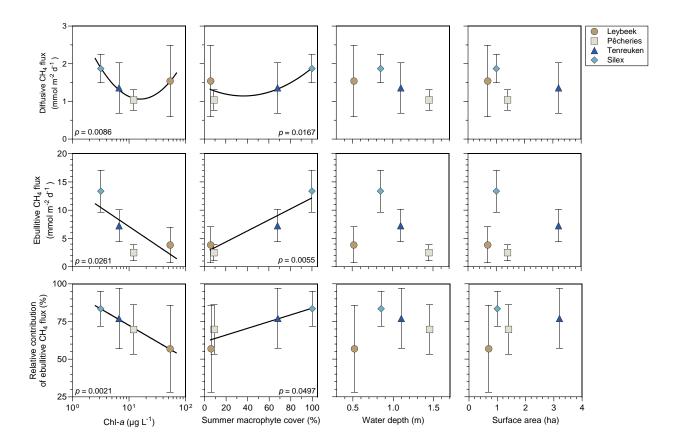


Figure 8: Mean diffusive and ebullitive CH_4 fluxes (mmol m^{-2} d^{-1}) and mean ratio of ebullitive CH_4 flux to total (diffusive+ebullitive) CH_4 flux (%) versus chlorophyll-a (Chl-a, in μg L^{-1}), total macrophyte cover in summer (%), water depth (m), and lake surface area (ha) in four ponds (Leybeek, Pêcheries, Tenreuken, and Silex) in the city of Brussels (Belgium) from June 2021 to December 2023. Error bars indicate the standard deviation. Solid lines indicate either linear or polynomial fits. Statistical comparisons between the four ponds are summarized in Table S3.

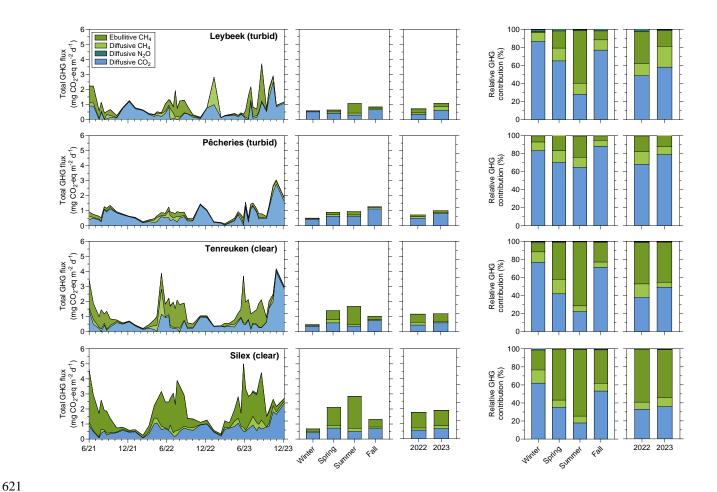


Figure 9: Seasonal and year-to-year variations of the emissions to the atmosphere of CO_2 (diffusive), CH_4 (diffusive and ebullitive), and N_2O (diffusive) expressed in CO_2 equivalents (in mg CO_2 -eq m⁻² d⁻¹) and their relative contribution in %, in four urban ponds (Leybeek, Pêcheries, Tenreuken, and Silex) in the city of Brussels (Belgium) from June 2021 to December 2023. Seasonal averages include data from 2021, 2022, and 2023. The annual precipitation was higher in 2023 (1011 mm) than in 2022 (701 mm).

- **Data availability.** The full data-set is available at 10.5281/zenodo.11103556.
- 627 Author contributions. AVB and NG conceived the study; TB collected field samples; TB and AVB made the laboratory
- 628 analysis; TB and AVB jointly interpreted data and drafted the manuscript with substantial inputs from NG.
- 629 **Competing interests.** The authors declare that they have no conflict of interest.
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