

Seasonal and spatial variations of greenhouse gas (CO₂, CH₄ and N₂O) emissions from urban ponds in Brussels

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Highlights :

- First survey of GHG emissions from Brussels ponds
- Small ponds are more subject to edge effects and have higher pCO₂
- Macrophytes enhance methane production in ponds
- City center ponds have higher N₂O emissions due to atmospheric deposition
- Brussels ponds emissions were equivalent to the carbon sink estimated for the urban green spaces.

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Abstract:

Freshwaters have been recognized as important sources of greenhouse gases (GHG) to the atmosphere. However, urban ponds have received little attention even though their number is increasing due to expanding urbanisation globally. Ponds are frequently associated to urban green spaces that provide several ecosystemic services such as cooling local climate, regulating the water cycle, and acting as small carbon sinks. This study aims to identify and understand the processes producing GHGs (CO₂, CH₄, and N₂O) in the urban ponds of the temperate European city of Brussels in Belgium. 22 relatively small ponds (0.1-4.6 ha) surrounded by contrasted landscape (strictly urban, bordered by cropland or by forest), were sampled during four seasons in 2021-2022. The mean \pm standard deviation was $3,667 \pm 2,904$ ppm for the partial pressure of CO₂ (pCO₂), $2,833 \pm 4,178$ nmol L⁻¹ for CH₄, and 273 ± 662 % for N₂O saturation level (%N₂O). Relationships of GHGs with oxygen and water temperature suggest that biological processes controlled pCO₂, CH₄ concentration and %N₂O. However, pCO₂ was also controlled by external inputs as indicated by the higher values of pCO₂ in the smaller ponds, more subject to external inputs than larger ones. The opposite was observed for CH₄ concentration that was higher in larger ponds, closer to the forest in the city periphery, and with higher macrophyte cover. N₂O concentrations, as well as dissolved inorganic nitrogen, were higher closer to the city center, where atmospheric nitrogen deposition was potentially higher. The total GHG emissions

30 from the Brussels ponds were estimated to 1kT CO₂-eq per year and were equivalent to the
31 carbon sink of urban green spaces.

32 **Keywords:**

33 *carbon dioxide, methane, nitrous oxide, Brussels, urban ponds, urban ecology,*
34 *macrophytes*

35

36 **1. Introduction**

37

38 The three main greenhouse gases (GHGs) emitted to the atmosphere by human
39 activities are carbon dioxide (CO₂), methane (CH₄) with a global warming potential (GWP)
40 34 times greater than CO₂ on a 100-year time scale, and nitrous oxide (N₂O) with a GWP
41 298 times greater than CO₂ on a 100-year time scale (Myrhe et al., 2013). In inland waters,
42 the production of CO₂ and CH₄ are mainly due to the degradation of organic matter (OM)
43 through mainly aerobic processes for CO₂ (Del Giorgio et al., 1999; Cole and Caraco, 2001)
44 and by anaerobic archaeal methanogenesis for CH₄ (Conrad, 2020). The OM in inland
45 waters can be autochthonous or allochthonous. Autochthonous production of OM is from
46 phytoplankton (McClure et al., 2020; Bartosiewicz et al., 2021) or aquatic macrophytes
47 (Grasset et al., 2019; Desrosiers et al., 2022). The relation between autochthonous biomass
48 (phytoplankton and macrophytes) and CH₄ emissions is well established (DeISontro et al.,
49 2018; Borges et al., 2022; Bastviken et al., 2023). The impact of autochthonous biomass on
50 CO₂ depends on the phase of the bloom development, high concentrations of chlorophyll-*a*
51 (Chl-*a*) during the growth and peak phases coinciding with intense photosynthesis and CO₂
52 consumption (Grasset et al., 2020; Borges et al. 2022;), and phases of senescence leading
53 to CO₂ production. Allochthonous OM in small water bodies (e.g., ponds) comes from fallen
54 leaves of the surrounding vegetation and from particulate and dissolved OM from surface
55 runoff or from soil-water and groundwater inputs (Gasith and Hosier, 1976; Weyhenmeyer
56 et al., 2015). The N₂O production is mainly due to microbial nitrification and denitrification
57 and is dependent on the availability of dissolved inorganic nitrogen (DIN) and O₂ (Codispoti
58 and Christensen, 1985; Mengis et al., 1997). Anthropogenic nitrogen inputs from leakage of
59 fertilizers from croplands or from atmospheric deposition may enhance the N₂O emissions
60 in freshwater wetlands, lakes and rivers (McCrackin & Elser, 2011; Bettez & Groffman, 2013;
61 Sønderup et al., 2016; D'Acunha et al., 2019; Decina et al., 2020; Bonetti et al., 2022).

62 The emissions from inland waters (rivers-streams, lakes and reservoirs) of GHGs
63 were estimated at 14.3 Pg CO₂ yr⁻¹ for CO₂ (Drake et al., 2018), 6.03 Pg CO₂ equivalent
64 (CO₂-q) yr⁻¹ for CH₄ (Rosentreter et al., 2021) and 0.05 Pg CO₂-eq yr⁻¹ for N₂O (Lauerwald
65 et al., 2019). Yet, these estimates remain highly uncertain because of the low number of
66 measurements that is insufficient to account for the strong spatial and temporal variability of
67 the fluxes within a given system, as well as the high diversity of systems due to differences
68 in size, climate, catchment morphology, land cover, and anthropogenic pressures.

69 Several studies have shown the importance of small water bodies (surface area < 0.1
70 ha) as hotspots for GHG emissions (Holgerson and Raymond, 2016; Grinham et al., 2018;
71 Rosentreter et al., 2021; Peacock et al., 2021). Although only representing 9% of the total
72 area of lentic waters, small water bodies account for 15% and 40% of diffusive emissions
73 from lakes of CO₂ and CH₄, respectively according to Holgerson and Raymond (2016).
74 Urban areas have a large number of small water bodies in the form of ponds mostly
75 associated to green spaces such as public parks, and their number is increasing due to rapid
76 urbanisation worldwide (Brans et al., 2018; Audet et al., 2020). Yet, only a very limited
77 number of studies have investigated GHG emissions from urban ponds (Singh et al., 2000;
78 Natchimuthu et al., 2014; van Bergen et al., 2019; Audet et al., 2020; Peacock et al., 2021).
79 Urban ponds are small and thus have high ratio of perimeter to surface area (Hanson et al.,
80 2007), surrounded by impervious surfaces (Davidson et al., 2015; Peacock et al., 2021),
81 and a high stormwater runoff that combined result in high inputs of OM and DIN that should
82 sustain emissions of CO₂, CH₄, and N₂O to the atmosphere. Additionally, urban ponds are
83 usually very shallow so there is a strong influence on the water column of GHG production
84 in sediments.

85 Brussels is the most densely populated area in Belgium and the largest city of the
86 country and contains 158 ponds totalling a surface area of 101ha. The population is more
87 than 1 million inhabitants, equating to less than 1m² of water body per inhabitant, indicating
88 a high degree of anthropogenic pressure on these ecosystems. However, not all urban
89 ponds are subject to the same degree of anthropogenic pressure, as the periphery of the
90 city is bordered by cropland and forest. Yet, the ponds in Brussels are highly eutrophied with
91 in some cases high algal biomass but also macrophytes which compete with phytoplankton
92 for nutrients (Peretyatko et al., 2007 ; de Backer et al., 2010 ; Peretyatko et al., 2012 ; Descy
93 et al., 2016). Yet, the equilibrium between the prevalence of macrophytes and phytoplankton
94 entails a gradual shift rather than an abrupt process and is influenced by additional factors
95 other than nutrient levels (Van Nes et al., 2002; Davidson et al., 2023). It can be

96 hypothesized that water bodies with a dominance of phytoplankton and a dominance of
97 macrophytes should have different GHG dynamics, although this has seldom been tested
98 (Harpenslager et al., 2022 ; Baliña et al., 2023).

99 Here, we report CO₂, CH₄, and N₂O dissolved concentrations and complementary
100 variables (water temperature, chlorophyll-a (Chl-a), DIN, and O₂) sampled in 22 small urban
101 ponds (0.1-4.6 ha) of the city of Brussels, during the four seasons in 2021-2022. Potential
102 drivers of CO₂, CH₄, and N₂O dynamics are investigated by comparing pond size, presence
103 of macrophytes, relative distance to the city center, surrounding landscape (strictly urban,
104 bordered by cropland or by forest). We test if the most frequent accepted hypotheses of the
105 drivers of CO₂, CH₄ and N₂O seasonal and spatial variations in natural lakes and ponds also
106 apply to urban ponds The CO₂, CH₄, and N₂O diffusive emissions are computed and
107 compared to the official inventory of GHG emissions from the city of Brussels.

108

109 **2. Material and methods**

110

111 **2.1. Pond selection and sampling frequency**

112

113 22 ponds were selected within the Brussels region (Fig. 1). All of the ponds are
114 artificial and were built during the past century, the majority with the primary aim of
115 landscaping and embellishing parks. Recreational fishing activities are allowed in four of the
116 sampled ponds (Table S1). Some of the sampled ponds are connected to a river network
117 (flow-through ponds) and others are fed directly by groundwater and small watercourses
118 (overflow ponds), where water level is kept constant by an overflow outlet (Table S1). The
119 sampled ponds are in the four main catchment areas of the city of Brussels (Molenbeek
120 (n=2), Neerpedebeek-Vogelzangbeek (n=4), Maelbeek (n=6), and Woluwe (n=10)
121 catchments) (Table S1). Four sampling campaigns were carried out on the 22 ponds,
122 corresponding to the four seasons: November 2021 (Fall), February 2022 (Winter), May
123 2022 (Spring) and August 2022 (Summer).

124

125 **2.2. Meteorological data**

126

127 Air temperature, rainfall and wind speed were retrieved from <https://wvw.meteo.be/fr>
128 for the meteorological station of the Royal Meteorological Institute of St-Lambert (50.8408
129 °N, 4.4234 °E) in Brussels, located between 1 and 10 km from the sampled ponds. Wind
130 speeds and air temperatures were averaged over 24 hours to obtain a daily average value.
131 The rainfall was integrated on each day to obtain the daily rainfall.

132

133 **2.3. Sampling in field**

134

135 Water was sampled and collected from pontoons. Water pH, temperature,
136 conductivity and oxygen saturation level (%O₂) were measured by a VWR MU 6100 H probe.
137 Water was collected in 2L polypropylene bottles for subsequent sampling for Chl-*a*, total
138 suspended matter (TSM) and dissolved inorganic nutrients (ammonium (NH₄⁺), nitrite (NO₂⁻
139), nitrate (NO₃⁻), soluble reactive phosphorus (SRP)). Three 50mL falcon tubes were filled
140 with unfiltered water to which was added 200µL of HNO₃ (65%) for total phosphorus (P_{tot})
141 analysis.

142 Macrophytes were defined as vascular aquatic plants, excluding filamentous algae,
143 mosses and liverworts, (Bowden et al., 2017). The percentage of macrophyte cover was
144 estimated visually during sampling. Dominant species of macrophytes (*Potamogeton*
145 *pectinatus*, *Ceratophyllum demersum*, *Chara sp.*, *Nitella sp.*, *Lemna trisulca*, *Zannichellia*
146 *palustris*) were identified in August 2022, when macrophyte cover was maximal. This list of
147 species of macrophytes agreed with past studies in Brussels ponds (Peretyatko et al., 2009;
148 de Backer et al., 2010).

149

150 **2.4. Laboratory analysis**

151

152 Water was filtered on Whatman filters 0.7µm GF/F glass microfibers with diameter of
153 47mm for TSM and Chl-*a* (stored frozen). Filtered water was stored in 50 ml plastic vials and
154 stored frozen for dissolved nutrients analysis. Chl-*a* was measured on extracts with 90%
155 acetone by fluorimetry (Kontron SFM 25 model) (Yentsch and Menzel, 1963). NH₄⁺ was
156 determined with the nitroprusside-hypochlorite-phenol coloration method (Grasshoff and
157 Johannsen, 1972). NO₂⁻ and NO₃⁻ were determined before and after reduction of NO₃⁻ to

158 NO₂⁻ by a cadmium-copper column, with Griess' reagent in acidic medium coloration method
159 (Grasshoff et al., 1983). SRP was determined with ammonium molybdate, ascorbic acid and
160 potassium antimony tartrate coloration method (Koroleff, 1983).

161 P_{tot} was determined by inductively coupled plasma spectroscopy (ICP) on an ICP-
162 OES Perkin Elmer Avio 200 model. The assay protocol was based on the US EPA (1994)
163 method 200.7 for analysis of metals and trace elements in water by ICP with prior microwave
164 acid digestion based on US EPA, (2007) method 3015A.

165

166 **2.5. Measurement of GHG concentrations and computation of fluxes**

167

168 CO₂ measurements were carried out on the field with a Li-Cor Li-840 CO₂/H₂O gas
169 analyser using the headspace technique with 4 polypropylene syringes of 60mL. The
170 technique consisted in equilibrating inside the syringe 30mL of sample water with 30mL of
171 atmospheric air by vigorous shaking during 5 minutes. The headspace of each syringe was
172 injected sequentially in the Li-840 and a fifth syringe was used to measure atmospheric CO₂.
173 The final pCO₂ value was computed taking into account the partitioning of CO₂ between
174 water and the headspace, as well as equilibrium with HCO₃⁻ (Dickson et al. 2007) using
175 water temperature measured in-stream and after equilibration, and total alkalinity (data not
176 shown). Samples for total alkalinity were conditioned, stored and analysed as described by
177 Borges et al. (2019). The Li-Cor Li-840 was calibrated before and after each sampling period
178 with ultrapure N₂ and a suite of gas standards (Air Liquide Belgium) with CO₂ mixing ratios
179 of 388, 813, 3788 and 8300 ppm. The overall precision of pCO₂ measurements was ±2%.

180 Samples for dissolved CH₄ and N₂O were collected directly in surface waters with 60
181 ml plastic syringes and transferred with a plastic tube into two 60mL borosilicate serum
182 bottles and poisoned with 200µL of saturated HgCl₂ solution. The vials were sealed with a
183 butyl stopper and crimped with an aluminium cap. Measurements were carried out using the
184 headspace technique with a SRI 8610C gas chromatograph with a flame ionisation detector
185 (FID) for CH₄ and an electron capture detector (ECD) for N₂O, calibrated with CH₄:N₂O:N₂
186 gas mixtures (Air Liquide Belgium) with mixing ratios of 1, 10, 30, 509, and 2010 ppm for
187 CH₄, and 0.2, 2.0 and 6.0 ppm for N₂O. The precision of measurement was ±11% for CH₄
188 and ±6% for N₂O based on 176 replicates. Ebullitive CH₄ fluxes were not measured during
189 this study, but were measured with inverted funnels (e.g. Keller and Stallard, 1994) in four

190 ponds (Leybeek, Pêcheres Royales, Silex, Ten Reuken) for the period of 29/03/22 to
 191 09/09/23 (unpublished data).

192 The CO₂ concentration is expressed in terms of partial pressure of CO₂ (pCO₂ in
 193 ppm), the CH₄ dissolved concentration in nmol L⁻¹ corresponding to the usual convention in
 194 topical literature. N₂O concentrations oscillated around atmospheric equilibrium, so the data
 195 are presented as a percentage of the saturation level (%N₂O), where 100% corresponds to
 196 atmospheric equilibrium. The atmospheric pCO₂ was measured on the field with the Li-Cor
 197 Li-840. The equilibrium with atmosphere for N₂O was calculated from the average air mixing
 198 ratios of N₂O provided by the Global Monitoring Division (GMD) of the National Oceanic and
 199 Atmospheric Administration (NOAA) Earth System Research Laboratory (ESRL) (Dutton
 200 and Hall, 2023). A constant atmospheric concentration of 1.9 ppm was assumed for CH₄.

201 The exchange of CO₂, CH₄, and N₂O between surface water and the atmosphere was
 202 calculated according to:

$$203 \quad F_G = k_g \Delta C$$

204 F_G (mmol m⁻²d⁻¹) is the flux of a specific gas (G), k_g (cm h⁻¹) is the specific gas transfer
 205 velocity, and ΔC is the gas concentration gradient in water and the atmosphere (mmol L⁻¹).

206 The k_g values were derived from the gas transfer velocity normalized to Schmidt
 207 number of 600 (k_{600}) according to :

$$208 \quad k_g = k_{600} \times (Sc/600)^{-0.5}$$

209 where Sc is the Schmidt number of the given gas in freshwater at in-situ water
 210 temperature computed according the algorithms given by Wanninkhof (1992).

211 k_{600} was computed from wind speed using the parameterisation of Cole and Caraco
 212 (1998):

$$213 \quad k_{600} = 2.07 + 0.215 u^{1.7}$$

214 where u is the wind speed (m s⁻¹).

215 The average daily u value on the day of sampling of each sampled pond was used.
 216 The u data were measured at the meteorological station of the Royal Meteorological Institute
 217 of St-Lambert (50.8408 °N, 4.4234 °E) in Brussels (located between 1 and 10 km from the
 218 sampled ponds) (retrieved from wow.meteo.be). The area-weighted average of F_G from the
 219 22 sampled ponds (totaling a surface area of 34 ha) for each of the 4 seasons was averaged

220 to obtain an annual value that was extrapolated to the total surface area (101 ha) of the all
221 of the 158 urban ponds of the city of Brussels.

222 The emissions of CH₄ and N₂O were expressed in CO₂ equivalents (CO₂-eq) by
223 multiplying the respective fluxes by global warming potential (GWP) values of 34 for CH₄
224 and 298 for N₂O corresponding to a time horizon of 100 years (Myrhe et al., 2013).

225

226 **2.6. Statistical analysis**

227

228 The statistical analyses were conducted using R version 4.1.1 (2022) (R Core Team,
229 2022) and graphs were produced with GraphPad Prism v9. Prior to analysis, data underwent
230 logarithmic or square root transformations to ensure normality. Shapiro tests were performed
231 to assess the normality of the distribution.

232 To investigate relationships between variables, quantile regressions were performed
233 with the quantreg package v5.95 (Koenker, 2005). Quantile regressions were chosen over
234 linear regressions due to their ability to provide a more comprehensive understanding of
235 potential causal relationships in ecological processes (Cade and Noon, 2003). Significance
236 levels for the quantile regression were determined using Wald statistical tests.

237 For comparisons of medians in the boxplots, permutational multivariate analysis of
238 variance (PERMANOVA) on each Bray-Curtis distance matrix with 999 permutations was
239 performed. A post-hoc test was performed to establish significant differences between
240 sample pairs. For multiple comparisons, *p*-values were adjusted using the Bonferroni
241 correction. Simultaneously, a test for homogeneity of multivariate dispersions (PERMDISP)
242 was conducted. This test allowed for the examination of dispersion effects independent of
243 location effects, as significant differences could be attributed to variations in within-group
244 dispersion rather than differences in medians values among the groups (Anderson, 2006).
245 PERMANOVA and PERDISP were performed using the vegan package v2.6.4 (Oksanen et
246 al., 2013).

247

248 **2.7. Data availability**

249

250 The time-sampled and georeferenced data-set presented in this paper is publicly
251 available (Bauduin et al., 2024).

252

253 **3. Results**

254

255 **3.1. Seasonal variations of meteorological conditions and GHGs**

256

257 In 2022, the average air temperature in Brussels was 12.2°C, above the average of
258 11.0°C for the 1991-2020 period. With the exception of April, September and December, air
259 temperature monthly averages were above the reference value for the 1991-2020 period. A
260 total of 701 mm of rain fell in Brussels (837 mm for the 1991-2020 period), during 149 days
261 (190 days for the 1991-2020 period). The year 2022 was the fourth driest and the first (ex-
262 aequo with 2020) warmest year of the current reference period. During the sampling period
263 from 22/11/21 to 18/08/22, the air temperature and wind speed averaged $11.9 \pm 6.3^\circ\text{C}$ and
264 $0.6 \pm 0.6 \text{ m s}^{-1}$, respectively, and the total precipitation was 321mm in the city of Brussels
265 (Fig. 2A). Air temperature was highest in summer ($22.2 \pm 1.6^\circ\text{C}$) and lowest in fall ($6.1 \pm 0.7^\circ\text{C}$).
266 The highest wind speed was in spring (Fig. 2B) ($1.7 \pm 0.5 \text{ m s}^{-1}$). In the 15 days prior to
267 sampling, it rained a total of 13mm in fall, 53mm in winter, 13mm in spring, and less than 1
268 mm in summer.

269 During the four sampling periods, pCO₂ averaged $3,667 \pm 2,904$ ppm, CH₄
270 concentration averaged $2,833 \pm 4,178 \text{ nmol L}^{-1}$, and %N₂O averaged 273 ± 662 % in the 22
271 sampled ponds in the city of Brussels. The maximum pCO₂ value was observed in fall
272 (15,029 ppm) and the minimum value in summer (121 ppm). Median pCO₂ was higher in fall
273 than during the other seasons and was lower in winter than spring (Fig. 3A). The maximum
274 and minimum CH₄ concentration values were both observed in summer, 29,190 and 10 nmol
275 L⁻¹, respectively (Fig. 3B). The CH₄ concentrations showed seasonal variations, with
276 significantly lower median concentration in winter than other seasons, and higher
277 concentrations during spring and summer (Fig. 3B). The minimum and maximum values of
278 %N₂O were both observed during summer (0 and 10,354%) (Fig. 3C). The median
279 concentration of %N₂O was lower during summer than fall and winter. The %N₂O values
280 showed a higher dispersion during summer than other seasons (Fig. 3C). Undersaturation
281 of CO₂ with respect to atmospheric equilibrium was observed only on 7 occurrences (out of
282 88 observations), during spring (2 times) and summer (5 times). For N₂O, undersaturation

283 with respect to atmospheric equilibrium was observed at each season, 3 times in fall, 5 times
 284 in winter, 7 times in spring and 10 times during summer. CH₄ concentrations were always
 285 above saturation.

286 The average, median, minimum and maximum values of ancillary variables (%O₂,
 287 Chl-*a*, SRP, P_{tot}, NH₄⁺, NO₃⁻ and NO₂⁻) are given in Table S2. The median %O₂ values were
 288 significantly lower in fall than in the other seasons (Table S2). The median Chl-*a*
 289 concentration value was significantly lower in winter than in fall and the highest
 290 concentrations were measured in summer, when values were the most variable. There were
 291 no differences between seasons for SRP. P_{tot} had higher concentrations in spring and
 292 summer than in fall and winter. DIN concentrations were significantly higher in fall and winter
 293 than during spring and summer. NH₄⁺, NO₃⁻ and NO₂⁻ concentrations were significantly
 294 higher in fall than during the other seasons.

295 pCO₂, CH₄ concentration and %N₂O were negatively related to %O₂ (Figs. 4A, 4G,
 296 4M). CH₄ concentration was positively related to water temperature (Fig. 4H) and %N₂O was
 297 negatively related to water temperature (Fig. 4N). There was no relation between pCO₂ and
 298 water temperature (Fig. 4B), nor Chl-*a* concentration (Fig. 4C). pCO₂ was positively related
 299 to SRP (Fig. 4E). CH₄ was negatively related to Chl-*a* (Fig. 4I). There was a positive
 300 correlation between CH₄ and P_{tot} (Fig. 4L), that was also observed during each individual
 301 season, except in winter (Figs. S2A, S2B, S2C, S2D). The negative relation observed
 302 between CH₄ and DIN (Fig. 4J) could be spurious and be attributed to parallel seasonal
 303 changes of both variables, as there were no correlations between CH₄ and DIN when
 304 analysed independently for each season, except in spring (Figs. S2E, S2F, S2G, S2H).
 305 %N₂O was positively correlated to DIN for the whole dataset (Fig. 4P) as well as for each
 306 season analysed independently (Figs. S2I, S2J, S2K, S2L). %N₂O was statistically more
 307 strongly correlated to NO₂⁻ (Fig. 5C) and NO₃⁻ (Fig. 5D) than to NH₄⁺ (Fig. 5B). %N₂O was
 308 negatively related to O₂:DIN (Fig. 5E), O₂:NO₂⁻ (Fig. 5G) and O₂:NO₃⁻ (Fig. 5H) but unrelated
 309 to O₂:NH₄⁺ (Fig. 5F).

310

311 **3.2. Spatial variations of GHGs**

312

313 As pond perimeter ($r = 0.90$, $p < 0.0001$) and depth ($r = 0.45$, $p = 0.014$) were correlated
 314 to surface area ; hereafter, only relationships between GHGs and pond surface area are

315 presented (Figs. 6A, 6E, 6I). Both $p\text{CO}_2$ and $\%\text{N}_2\text{O}$ were negatively related to pond surface
 316 area (Figs. 6A, 6I) and CH_4 was positively related to pond surface area (Fig. 6E). The ponds
 317 closer to the city center had lower CH_4 concentrations (Fig. 6F) and higher $\%\text{N}_2\text{O}$ (Fig. 6J)
 318 than the ponds closer to the periphery of the city. $\%\text{N}_2\text{O}$ values showed a higher dispersion
 319 in small ponds than larger ponds and in ponds closer to the city center compared to the
 320 periphery of the city. $p\text{CO}_2$ did not show a clear pattern with regards to the city center (Fig.
 321 6B). DIN , NO_3^- and NO_2^- concentrations decreased with distance from the city center but not
 322 NH_4^+ (Fig. 7). Atmospheric nitrogen dioxide (NO_2) concentration decreased with distance
 323 from the city center (Fig. 7E). $\%\text{N}_2\text{O}$ was positively correlated to the atmospheric NO_2
 324 concentration (Fig. 7F). Elevated CH_4 concentrations were observed in ponds with higher
 325 macrophyte cover (Fig. 6G). Ponds with high macrophyte cover were located farther away
 326 from the city center (Fig. S3A), had larger surface areas (Fig. S3B), and lower residence
 327 times (Fig. S3C). Ponds with high macrophyte cover exhibited lower concentrations of DIN
 328 and Chl-a than ponds with low macrophyte cover (Fig. S3D,E). Ponds with high and low
 329 macrophyte cover had no difference in SRP and P_{tot} concentration (Figs. S3F,G). No
 330 significant correlation was observed between Chl-a and distance from city center (Fig. S4).

331

332 3.3. Fluxes of GHGs

333

334 The mean and median concentrations of the three GHGs in the sampled ponds were
 335 consistently higher than their respective equilibrium concentrations (Fig. 3). Consequently,
 336 the sampled ponds were sources of GHGs to the atmosphere during all four season,
 337 although negative flux values were observed on some rare occasions (Fig. S5). Seasonal
 338 variations of the GHG fluxes mirrored those of the respective concentrations because of
 339 modest variations of wind speed and computed k_{600} , although higher fluxes were observed
 340 during spring characterized by the highest daily wind speeds (Fig. S5A).

341 In terms of GHG emissions expressed in CO_2 equivalents, CO_2 was the dominant
 342 GHG emitted, with a median annual emission to the atmosphere of $732 \text{ g CO}_2 \text{ m}^{-2} \text{ yr}^{-1}$.
 343 Median annual emission of CH_4 and N_2O was 97 and 9 $\text{g CO}_2\text{-eq m}^{-2} \text{ yr}^{-1}$, respectively. CO_2
 344 contributed to 88% of the total CO_2 equivalent emissions, CH_4 contributed to 11%, and N_2O
 345 contributed to 1%. The total annual diffusive emissions of CO_2 , CH_4 , and N_2O amounted to
 346 0.8 kT of $\text{CO}_2\text{-eq}$ for all of the ponds in the city of Brussels ($n=158$) (Fig. 8).

347

348 **4. Discussion**

349

350 **4.1. Seasonal variations**

351

352 Seasonal variations of dissolved CO₂ in lacustrine systems at mid and high latitudes
353 depend on the seasonal alternation between periods of net ecosystem heterotrophy and
354 periods when the aquatic ecosystem evolves to more autotrophic conditions. This seasonal
355 alternation depends on light conditions, with a higher aquatic primary production in spring
356 and summer. Light conditions co-vary with temperature, and warmer conditions coincide
357 seasonally with the sunnier conditions in summer. Allochthonous organic matter inputs from
358 the watershed can also vary seasonally and are usually stronger in fall and winter than spring
359 and summer.

360 In the sampled ponds of the city of Brussels, the overall negative relation between
361 pCO₂ and %O₂ (Figs. 4A) reflected an alternation between autotrophy-heterotrophy
362 (balance between photosynthesis and community respiration), as also observed in other
363 studies in rivers and lakes (Borges et al., 2015, 2018). This confirmed by the observed
364 relationship between pCO₂ and SRP (Fig. 4F) that may reflect the uptake of nutrients by
365 autotrophic groups leading to the lowering of SRP and pCO₂, and, conversely, high SRP
366 and pCO₂ resulting from the degradation of OM in the water column and sediments (Alleson
367 et al., 2020). The pCO₂ values in the sampled ponds of the city of Brussels were highest in
368 fall and lowest in summer. In fall, the senescence of photosynthetic organisms, induced by
369 a decrease in light intensity and water temperatures, will lead to a large amount of potentially
370 degradable autochthonous OM (Bartosiewicz et al., 2021), which will be respired and lead
371 to higher pCO₂ in fall than in winter and summer. The input in fall of allochthonous OM to
372 ponds is increased with leaf fall from surrounding trees (Sønderup et al., 2016) and with
373 runoff due to higher precipitation . In spring, the occurrence of aquatic primary production in
374 response to the increase of the photoperiod and water temperature, lead to a CO₂
375 consumption and O₂ production.

376 In summer, the median (average) pCO₂ was not significantly different compared to
377 other seasons, but there was a higher dispersion of the values that explained the lack of
378 relation between pCO₂ and water temperature and Chl-a (Figs. 4B, 4C), unlike CH₄ and

379 %N₂O that correlated to water temperature (Figs. 4H, 4N). The high dispersion of pCO₂
380 values in summer may be attributed to sampling different stages of phytoplankton
381 development in various ponds during the summer campaign (Fig. S4). Low pCO₂ values
382 were observed in ponds where phytoplankton was blooming and high pCO₂ values were
383 observed in ponds where phytoplankton was senescent. The relation of pCO₂ and Chl-*a*
384 was further complicated by the variable density of macrophyte cover among the different
385 sampled ponds. High macrophyte cover corresponded to low Chl-*a* concentrations, while
386 low macrophyte cover corresponded to higher Chl-*a* levels (Fig. S3D).

387 Low %O₂ values in the water column indicate a generalized net heterotrophy
388 within the sampled ponds including the sediments and therefore also a high
389 methanogenesis, leading to an overall negative relation between CH₄ and %O₂ (Fig. 4G),
390 as also observed in other urban ponds (Audet et al., 2020; Peacock et al., 2021). Since
391 methanogenesis occurs in sediments rich in organic matter, seasonal variations of
392 methanogenesis are very strongly dependent on temperature variations (Zeikus and
393 Winfrey, 1976; Yvon-Durocher et al., 2014; Chen et al., 2021). Consequently, there was also
394 a highly significant positive correlation between CH₄ and water temperature (Fig. 4H). There
395 is a highly significant positive correlation between P_{tot} and CH₄, as in other urban ponds
396 (Peacock et al., 2019; Rabaey and Cotner, 2022), probably reflecting the delivery of fresh,
397 labile organic matter to sediments enhancing methanogenesis (Grasset et al., 2021; Gruca-
398 Rokosz and Cieśla, 2021 ; Nijman et al., 2022). P_{tot} was similar in ponds with high and low
399 macrophyte cover (Fig. S3G), indicating P_{tot} gives a measure of potentially degradable OM
400 from both phytoplankton and macrophytes. Consequently, P_{tot} was higher in spring and
401 summer (Table S2), when primary production from phytoplankton and macrophytes is
402 highest.

403 N₂O in lakes and ponds results from nitrification and denitrification that show seasonal
404 variations, mainly as a function of DIN availability and O₂ conditions (in particular in thermally
405 stratified systems), as well as water temperature (Myrstener et al., 2016; Palacin-Lizarbe et
406 al., 2018; Mander et al., 2021). Additionally, the yield of N₂O production from both nitrification
407 and denitrification also depends on O₂ levels and water temperature. In the sampled ponds,
408 %N₂O was higher in fall than summer, as observed by Mander et al. (2021) in artificial
409 wetlands. During fall, high heterotrophic degradation of organic matter and ammonification,
410 combined with lower uptake of NH₄⁺ by phytoplankton, resulted in high concentrations of
411 NH₄⁺ (Table S2). This led to high N₂O production from nitrification due to higher NH₄⁺
412 availability, and the N₂O yield from nitrification was possibly enhanced by low O₂ conditions

413 (Rosamond et al., 2012; Soued et al., 2016). In spring and summer, the NH_4^+ was
414 assimilated by phytoplankton and there was substrate limitation for N_2O production by
415 nitrification in the water column. The combination of high and low nitrification in fall and
416 spring, respectively explained the observed overall relationship between $\%\text{N}_2\text{O}$ and NH_4^+
417 (Fig. 5B). NO_2^- and NO_3^- are, respectively, intermediate and final product of the nitrification,
418 which explained the observed overall relationship between $\%\text{N}_2\text{O}$ and NO_2^- and NO_3^- (Figs.
419 5C, 5D) that was statistically more significant than the relation between $\%\text{N}_2\text{O}$ and NH_4^+ .
420 Low O_2 levels will favour N_2O production over NO_2^- production during nitrification (Goreau
421 et al., 1980; Ni et al., 2011), leading to a negative relation between $\%\text{N}_2\text{O}$ and $\text{O}_2:\text{DIN}$,
422 $\text{O}_2:\text{NO}_2^-$ and $\text{O}_2:\text{NO}_3^-$ (Figs. 5E, 5G, 5H). While the correlations between N_2O and NH_4^+ ,
423 NO_3^- and NO_2^- (Figs. 5B, 5C, 5D) in the sampled ponds of the city of Brussels indicate the
424 importance of nitrification in driving the variability of N_2O , denitrification should also have
425 played a role in N_2O dynamics. An indication of this is given by the overall negative
426 relationship between $\%\text{N}_2\text{O}$ and water temperature (Fig. 4N), as the last step of
427 denitrification is inhibited at low temperatures (Liao et al., 2018; Velthuis and Veraart, 2022),
428 leading to an accumulation of N_2O in the water column during fall and winter. A pattern of
429 higher N_2O levels during wintertime was also observed in boreal lakes and attributed to the
430 effect of temperature on the N_2O yield from denitrification (Kortelainen et al., 2020).

431

432 **4.2. Spatial variations in GHGs**

433

434 The levels of CO_2 and CH_4 in lakes vary as a function of lake size (Lapierre and del
435 Giorgio, 2012; Kankaala et al., 2013; Raymond et al., 2013; Holgerson and Raymond, 2016;
436 Casas-Ruiz et al., 2021; Borges et al., 2022; Chiriboga et al. 2024) as well as land cover
437 (Maberly et al., 2013). Combined lake size and land cover determine relative levels of
438 allochthonous organic matter inputs to aquatic systems and the general balance of
439 autotrophy and heterotrophy largely determining CO_2 levels (del Giorgio and Peters, 1994),
440 as well as benthic methanogenesis. The size of the water body will also largely determine
441 the relative importance of soil-water and ground-water inputs of CO_2 and CH_4
442 (Weyhenmeyer et al., 2015; Olid et al. 2022). The size of the water body will also determine
443 to a larger extent its depth, with smaller systems being in general shallower (Wetzel, 2001).
444 Depth will in turn determine the degree of coupling between sedimentary processes and
445 surface waters, as well as the possibly development of macrophytes. Yet, given the low

446 number of studies, it has not been investigated if these general patterns established from
447 studies in natural lakes and ponds can be applied to urban ponds. In urban ponds the
448 catchment cover is obviously very different from more natural systems, and there is a very
449 strong atmospheric nitrogen deposition. Inter-lake variations of N₂O have been less
450 investigated than CO₂ and CH₄, although DIN levels and depth are expected to exert a
451 strong influence on N₂O lacustrine levels (Lauerwald et al., 2019; Borges et al., 2022).

452 The surface area of the sampled ponds was overall low and the range of variation
453 was small (0.1-4.6 ha), so that other properties than pond size might have driven inter-
454 system differences. The analysis of spatial patterns of GHGs in the sampled ponds was
455 complicated by the fact that the smaller systems were located towards the center of city, and
456 the larger ones at the periphery in contact with either cropland to the West and the Sonian
457 forest to the East. Furthermore, the larger systems were characterized by a more important
458 macrophyte cover.

459 The pCO₂ values in the sampled ponds did not show a significant relation to the
460 distance from the city center or macrophyte cover (Figs. 6B, 6C). However, the pCO₂ values
461 were significantly higher in the smaller systems than the larger ones (Fig. 6A). This agrees
462 with several studies at local, regional or global scales showing a negative relation between
463 pCO₂ and lake surface area (Lapierre and Del Giorgio, 2012; Kankaala et al., 2013;
464 Raymond et al., 2013; Holgerson and Raymond, 2016; Casas-Ruiz et al., 2021). The causes
465 of such relations are related to the overall influence of lake size on inputs from the landscape
466 and lake carbon cycling and have been discussed in detail in several other studies (del
467 Giorgio and Peters, 1994; Sand-Jensen and Staehr, 2007; Borges et al., 2022).

468 The CH₄ concentration in the sampled ponds was significantly higher in the larger
469 systems than the smaller ones (Fig. 6E). Several studies reported a negative correlation
470 between CH₄ concentration and lake size (surface and/or depth) across spatial scales (local
471 to global) (Borges et al., 2011; Kankaala et al., 2013; Holgerson and Raymond, 2016; Borges
472 et al., 2022; Chiriboga et al. 2024), yet one study reported no relationship (Peacock et al.,
473 2019) and another a positive correlation (Rabaey et al., 2022). The positive relationship
474 between CH₄ concentration and pond size in the ponds of the city of Brussels was probably
475 indirect and reflected another driver. The relationship of CH₄ concentrations in relation to the
476 distance from the city center and to macrophyte cover was statistically more significant than
477 to surface area (Figs. 6E, 6F, 6G). We hypothesize that macrophyte cover was the
478 predominant driver (given the stronger statistical significance) that was itself a function of

479 distance from the city center. The statistically weak negative relationship between Chl-*a* and
480 CH₄ (Fig. 4J) could reflect the fact that ponds with high macrophyte cover tended to exhibit
481 lower Chl-*a* concentrations (Fig. S3D) but had significantly higher CH₄ concentrations (Fig.
482 6G).

483 Submerged macrophytes have a complex impact on CH₄ dynamic in lakes and
484 ponds, as synthesized by Bastviken et al. (2023), with both positive and negative effects on
485 the CH₄ emissions. In the urban ponds of the city of Brussels, the CH₄ concentration was
486 significantly higher in ponds with a high macrophyte cover (Fig. 6G), located in the periphery
487 of the city and closer to the Sonian Forest (Figs. 6F, 6H). In addition to the potential positive
488 impact of macrophytes on CH₄, the Sonian forest likely transfers allochthonous carbon as
489 plant litter promoting CH₄ production compared to other ponds. Litter from trees can induce
490 sediment anoxia (Mehring et al., 2014) enhancing CH₄ production. Yet, this potential
491 additional input of allochthonous organic carbon from the Sonian Forest did not seem to
492 affect the pCO₂ in ponds for which the effect of size seemed more important.

493 The %N₂O levels were significantly higher but also more variable in the sampled
494 urban ponds closer to the city center than at the periphery of the city (Figs. 6J, 6L). %N₂O
495 was strongly correlated to DIN, NO₂⁻ and NO₃⁻ concentrations (Fig. 5) that were higher close
496 to the city center (Fig. 7). The higher DIN, NO₂⁻ and NO₃⁻ concentrations in the city center
497 could have resulted from higher N deposition as indicated by the pattern of atmospheric NO₂
498 (Fig. 7E). No correlation was found with macrophyte cover (Fig. 6K), although the presence
499 of macrophytes strongly influences nitrogen cycling in sediments (Barko et al., 1991;
500 Choudhury et al., 2018; Dan et al., 2021; Ni et al., 2022) and could in theory potentially have
501 affected N₂O levels.

502

503 **4.3. Comparison with other urban and natural ponds**

504

505 There are only a few equivalent studies of GHG emissions in urban ponds for
506 comparison (Table S3). We restricted the comparison to dissolved GHG concentrations
507 rather than fluxes across the air-water interface. Fluxes measured with floating chambers
508 and computed fluxes from *k* can provide diverging results, so are not necessarily
509 comparable (Duchemin et al., 1999; Guérin et al., 2007; Klaus and Vachon, 2020; Erkkilä et

510 al., 2018; Perolo et al., 2021). This is possibly related to measurement biases using floating
511 chambers (Belanger and Korzun, 1991; Lorke et al., 2015; Vingiani et al., 2021), although
512 the computation of k can be made with different parameterizations derived from several
513 methods (including floating chambers) and that provide different values particularly at high
514 wind speeds. It is unclear which wind parametrization is the most appropriate (Klaus and
515 Vachon, 2020), in particular with regards to effects of size and fetch limitation (Wanninkhof,
516 1992) which might become very important in small water bodies such as ponds (Holgerson
517 et al., 2017; Jansen et al., 2020). We computed the fluxes from dissolved concentrations of
518 CO₂, CH₄ and N₂O and the gas transfer velocity computed from the parameterization as a
519 function of wind speed of Cole and Caraco (1998). The parametrization of Cole and Caraco
520 (1998) was built from a large compilation of deliberate tracer based estimates of k_{600} in 11
521 lakes with a wide range of maximum depth (1-109m) and surface area (0.2-487 km²). The
522 parametrization of Cole and Caraco (1998) parameterization as function of wind speed
523 intrinsically integrates other sources of turbulence (night-time convection due to cooling) and
524 fetch limitation, because it is built on compilation of repeated tracer measurements in several
525 lakes with a wide range of size and depth. Yet, the parameterization of Cole and Caraco
526 (1998) is based on k_{600} data acquired in water bodies with a surface area larger (>4 ha) than
527 the ponds we sampled (0.1-4.6 ha), which admittedly introduces uncertainty in the
528 computation of k_{600} . The computed k_{600} from the sampled ponds with the parametrization of
529 Cole and Caraco (1998) ranged from 2.1 to 2.6 cm h⁻¹ in summer and spring, respectively.
530 The overall annual average k_{600} value of 2.3 cm h⁻¹ was within the range of k_{600} values (0.9-
531 3.0 cm h⁻¹) reported by Holgerson et al. (2017) in four ponds located in a dense forest, with
532 surface area ~0.02 ha. These four ponds located in a dense forest were smaller than the
533 ponds we sampled (0.1-4.6 ha) that were located in recreational urban parks, hence, less
534 sheltered from wind than in a dense forest. The values of computed k_{600} from the sampled
535 ponds with the parameterization of Cole and Caraco (1998) are also comparable to the
536 values of k_{600} used by Holgerson and Raymond (2016), in a global analysis of CO₂ and CH₄
537 emissions from ponds, of 1.5 cm h⁻¹ for ponds with a surface area <0.1 ha and of 2.0 cm h⁻¹
538 for ponds with a surface area of 0.1-1.0 ha.

539 The average of CH₄ dissolved concentrations (annual or summer average) in the
540 sampled ponds of Brussels was comparable to the values reported in other urban ponds in
541 the cities of Silkeborg (Denmark, Audet et al., 2020), Uppsala (Sweden, Peacock et al.,
542 2019; Ray et al., 2023), Linköping (Sweden, Natchimuthu et al., 2014), Minnesota (United
543 States, Rabaey & Cotner, 2022; Ray et al., 2023) and in Salaspils (Latvia, Ray et al., 2023).

544 The average dissolved CO₂ concentrations in the sampled ponds was also equivalent to the
545 values in the ponds of Silkeborg (average of 4 seasons) (Audet et al., 2020) and possibly
546 lower in summer than the ponds in Uppsala (Peacock et al., 2019).

547 The extreme variability of CH₄ and CO₂ concentrations seasonally and among ponds
548 indicated by the large values of the standard deviations was observed in all systems
549 (Brussels, Silkeborg, Uppsala, Minnesota and Salaspils) and probably obscured differences
550 among different studies that could have emerged from differences in inputs of allochthonous
551 carbon and nitrogen, despite the differences in the size of the cities, as Brussels (> 1 million
552 inhabitants) is larger than Silkeborg (<50,000 inhabitants) and Uppsala (140,000
553 inhabitants). The N₂O concentration in the sampled ponds of Brussels was higher than
554 values in the ponds of Silkeborg (average of 4 seasons) (Audet et al., 2020), which might
555 be a consequence of a larger anthropogenic input of nitrogen from atmospheric deposition
556 due to the larger size of the city of Brussels compared to Silkeborg. This was consistent with
557 a higher value of the annual average NH₄⁺ concentration in the ponds of Brussels (27±63
558 μmol L⁻¹) compared to the ponds of Silkeborg (9±14 μmol L⁻¹, Audet et al., 2020). A stronger
559 eutrophication (nitrogen levels) sustaining a higher primary production, might also explain
560 potentially lower CO₂ concentrations in Brussels in summer than in Silkeborg and Uppsala
561 (Table S3).

562 Concentrations of CO₂ and CH₄ in the sampled ponds of Brussels were >3 times
563 higher than those in natural (non-urban) ponds of similar size reported by Holgerson and
564 Raymond (2016) (Fig. 9). This confirms the emerging idea that artificial inland water bodies
565 (agricultural reservoirs, urban ponds, storm-water retention basins, ...) have higher emission
566 rates per m² than natural systems (Martinez-Cruz et al., 2017; Grinham et al., 2018; Herrero
567 Ortega et al., 2019; Gorsky et al., 2019; Ollivier et al., 2019; Peacock et al., 2019, 2021;
568 Webb et al., 2019). This seems to result from more important anthropogenic allochthonous
569 inputs of carbon and nitrogen in artificial systems but might also reflect other differences
570 compared to natural systems such as in hydrology (Clifford and Heffernan, 2018).

571

572 **4.4. Contribution of urban ponds to the GHG budget of the city of Brussels**

573

574 The CO₂, CH₄ and N₂O air-water fluxes were converted into CO₂ equivalents and the
575 average scaled to total surface area of ponds in the city of Brussels (101 ha). The diffusive

576 emissions were dominated by CO₂ (88%), followed by CH₄ (11%) and N₂O (1%). The GHG
577 emissions officially accounted from the city of Brussels are overwhelmingly dominated by
578 CO₂ (99.1%) with a very small contribution from N₂O (0.6%) and CH₄ (0.3%). The relative
579 additional contribution of emissions of GHG from ponds to the total urban emissions is
580 different for each of the three GHGs individually (Table S4). The diffusive emissions from
581 ponds correspond to a fraction of the other urban emissions of 1.2% for CH₄, 0.05% for N₂O
582 and 0.03% for CO₂. The total annual GHG diffusive emissions from ponds of 0.8 kTCO₂eq
583 is several orders of magnitude lower and very marginal compared to the three major
584 anthropogenic emissions in the city of Brussels such as energy (1804 kTCO₂eq yr⁻¹),
585 transport (807 kTCO₂eq yr⁻¹), and industries (277 kTCO₂eq yr⁻¹) (Fig. 8). Ebullitive fluxes of
586 CH₄ were measured in four ponds (Leybeek, Pêcherries Royales, Silex, Ten Reuken) during
587 2021, 2022, and 2023 and ranged from 0.0 to 59.2 mmol CH₄ m⁻² d⁻¹ (unpublished data).
588 Based on an exponential relationship between ebullitive CH₄ and temperature in the 4
589 sampled ponds (n=39, r²=0.54, not shown), we scaled the ebullitive CH₄ fluxes to all of
590 ponds in the city of Brussels that summed correspond to an additional GHG emission of 0.2
591 kTCO₂-eq yr⁻¹. When additionally accounting for CH₄ ebullition, the total CH₄ emissions from
592 the urban ponds (0.2 kTCO₂-eq yr⁻¹) correspond to 3.7% of the total urban emissions of
593 CH₄. Yet, the total CH₄ emissions (including CH₄ ebullition) from the urban ponds only
594 represented 0.01% of the total GHG emissions of the city of Brussels (2885 kTCO₂-eq yr⁻¹).
595 This was consistent with the contribution of CH₄ emissions from streams and ponds of
596 0.004% to the total GHG emissions from the city of Berlin reported by Herrero Ortega et al
597 (2019). The total annual GHG emissions from urban ponds, (1.0 kTCO₂-eq yr⁻¹ including
598 CH₄ ebullition) is of the same order of magnitude as smaller GHG emissions such as waste
599 (5.3 kTCO₂eq yr⁻¹) and agriculture (1.3 kTCO₂eq yr⁻¹). The total annual GHG emissions from
600 urban ponds was nearly identical than the estimated sink of GHGs from urban green spaces
601 (1.2 kTCO₂eq yr⁻¹). Urban green spaces (parks and gardens) provide several ecosystem
602 services reviewed by Pataki et al. (2011) including reducing or offsetting GHG emissions.
603 Here, we showed that the emissions of GHGs from ponds almost fully balance the reported
604 GHG sink reported for green spaces in the city of Brussels. As such, GHG emissions from
605 ponds should be accounted among the negative consequences or tradeoffs of implementing
606 green infrastructure, the so-called “ecosystem disservices” according to Pataki et al. (2011).

607

608 **5. Conclusions**

609 Our study showed that small urban ponds in the public parks of the city of Brussels
610 emit CO₂, CH₄ and N₂O to the atmosphere. The GHG concentrations and emissions in the
611 sampled ponds showed geographical variations that were partly related to differences in
612 morphology (size) but also the position with regards to the city center, and the surrounding
613 landscape for those in the urban periphery (cropland versus forest), as well as the presence
614 of macrophytes. Smaller ponds received larger amounts of allochthonous carbon, either
615 directly as CO₂ or as OM resulting in larger CO₂ production from respiration. Larger ponds
616 emitted more CH₄ contrary to the pattern most commonly described in the literature. This
617 effect was due to the presence of macrophytes and the proximity of the forest in larger ponds
618 at the periphery of the city compared to the smaller ponds located closer to the city center.
619 N₂O levels were higher closer to the city center that we hypothesize resulting from a stronger
620 atmospheric nitrogen deposition based on the patterns of atmospheric NO₂.

621 Biological processes were the main drivers of GHG production, as showed from
622 relationships of the three GHGs with water temperature, oxygen saturation and nutrients.
623 CH₄ concentrations were positively related to temperature on seasonal time scales due to
624 the dependence of methanogenesis on temperature, as usually observed in aquatic systems
625 (Yvon-Durocher et al., 2014), including urban ponds (Bartosiewicz et al., 2016). On the
626 contrary the N₂O levels were negatively related to temperature possibly reflecting
627 incomplete denitrification at low temperatures, although DIN levels also increased in winter,
628 so it is difficult to separate the impact of both potential drivers.

629 The GHG emissions from the studied ponds were lower than the other GHG sources
630 from the city such as road transport and building heating. However, urban green spaces are
631 described as important carbon sinks by Pataki et al. (2011), but this sink was offset by
632 emissions from the ponds within these same green spaces in the city of Brussels. CH₄
633 emissions from inland waters were incorporated in the latest IPCC assessment report
634 (IPCC, 2019) and should now be included in current national inventories. We show that the
635 total CH₄ emissions from the urban ponds (including diffusive and ebullitive components)
636 correspond to a larger fraction of the total respective urban emissions (3.73%) than CO₂
637 (0.05%) and N₂O (0.03%). The emissions of GHGs from the urban ponds were equivalent
638 to the reported GHG sink reported for green spaces in the city of Brussels.

639 Audet et al. (2020) proposed to consider the GHG emissions from ponds in the overall
640 GHG assessment of cities to provide a more comprehensive quantification of a city's carbon
641 footprint and fully quantify the ecosystem services provided by urban green spaces. Yet, the

642 total CO₂, N₂O, and CH₄ emissions (including CH₄ ebullition) from the urban ponds only
643 represented 0.04% of the total GHG emissions of the city of Brussels (2885 kTCO₂-eq yr⁻¹).

644

645 **Acknowledgements**

646 TB received a funding from the Brussels-Capital Region's institute for the
647 encouragement of scientific research and innovation (Innoviris) as part of the Smartwater
648 project (ref. Innoviris RBC/2020-EPF-6h) and also from the "Fonds pour la formation à la
649 Recherche dans l'Industrie et dans l'Agriculture" (FRIA, Belgium). CH₄ and N₂O analyses
650 were done by Thomas Bousmanne and Ozan Efe. The analysis of total phosphorus was
651 performed at the Institut Meurice, within the Department of Physical Chemistry and Catalysis
652 at Labiris, under the supervision of Vincent Dubois and Grégory Ploeglaerts. AVB is a
653 Research Director at the Fonds National de la Recherche Scientifique.

654

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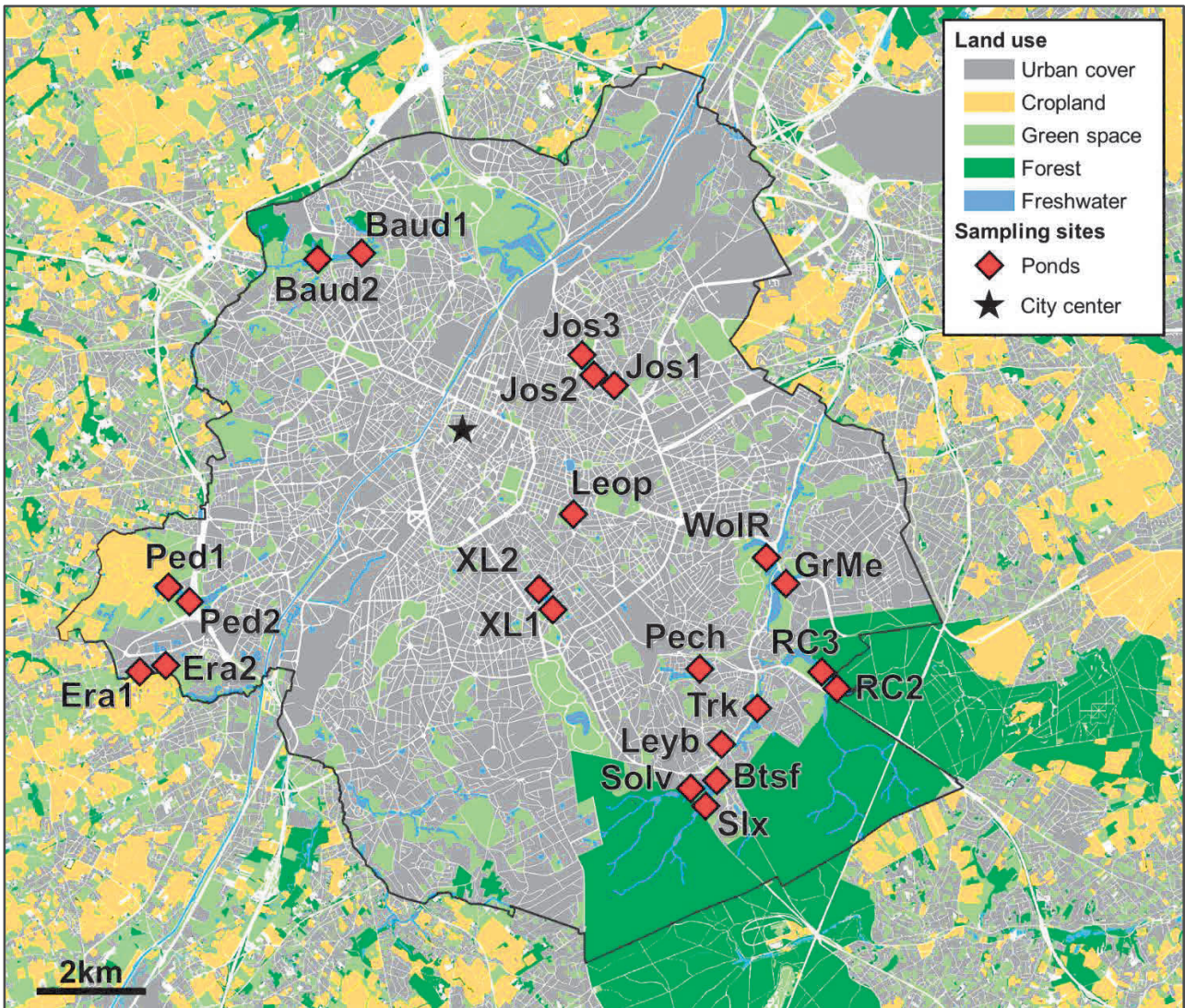


Figure 1 : Map of the metropolitan area of the region of Brussels delineated by the black line and surrounding region of Flanders in Belgium showing land cover and sampling locations in ponds (red diamonds). The star corresponds to the center of the city (50.8504°N, 4.3487°E). The acronyms correspond to the official names of the ponds of Bruxelles Environnement, the regional water management agency (Baud 1 : Etang Parc Roi Baudouin Phase I ; Baud2 : Etang Parc Roi Baudouin Phase II ; Btsf : Etang de Boitsfort ; Era 1 : Etang Erasme amont ; Era 2 : Etang Erasme aval ; GrMe : Grand étang Mellaerts ; Jos1 : Parc Josaphat - Etang aux canards ; Jos2 : Parc Josaphat – Etang aux pigeons ; Jos3 : Parc Josaphat – Etang de la laiterie ; Leop : Etang parc Leopold ; Leyb : Etang du Leybeek ; Pech : Etang Pêcheuries Royales ; Ped1 : Etang de la Pède Grand Etang ; Ped2 : Etang de la Pède ; RC2 : Grand étang des Clabots (Etang 2) ; RC3 : Etang du moulin (Etang 3) ; Slx : Etang du Silex ; Solv : Grand étang – Parc Tournay Solvay ; Trk : Etang Parc Ten Reuken ; WolR : Etang rond – Parc de Woluwe ; XL1 : Etang d’Ixelles Nord ; XL2 : Etang d’Ixelles Sud). The shape of each pond and connection to the hydrological network is shown in Figure S1.

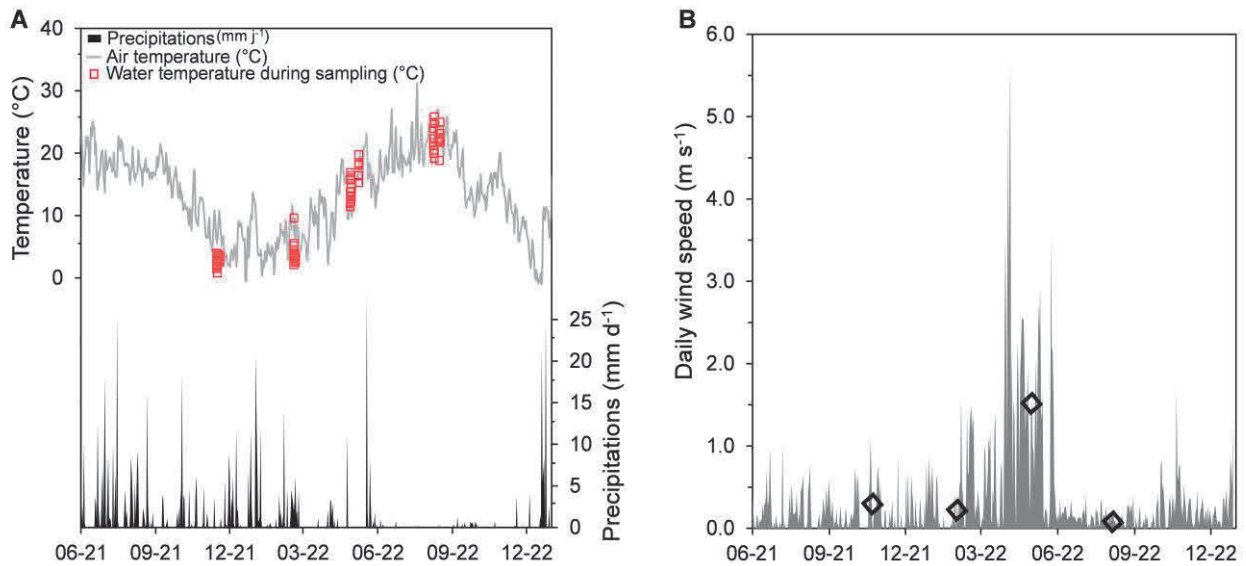


Figure 2 : Daily mean air temperature (light grey line), daily precipitation (black line) (A) and daily wind speed (dark grey, right) (B). The empty red squares on (A) indicate the in-situ water temperature during sampling and the empty black diamonds on (B) indicate the average wind during each sampling period. Meteorological data were obtained from crowdsourcing platform of the Royal Belgian Meteorological Institute (wow.meteo.be) at the Brussels–Woluwe Saint Lambert station (50.8408 °N, 4.4234 °E).

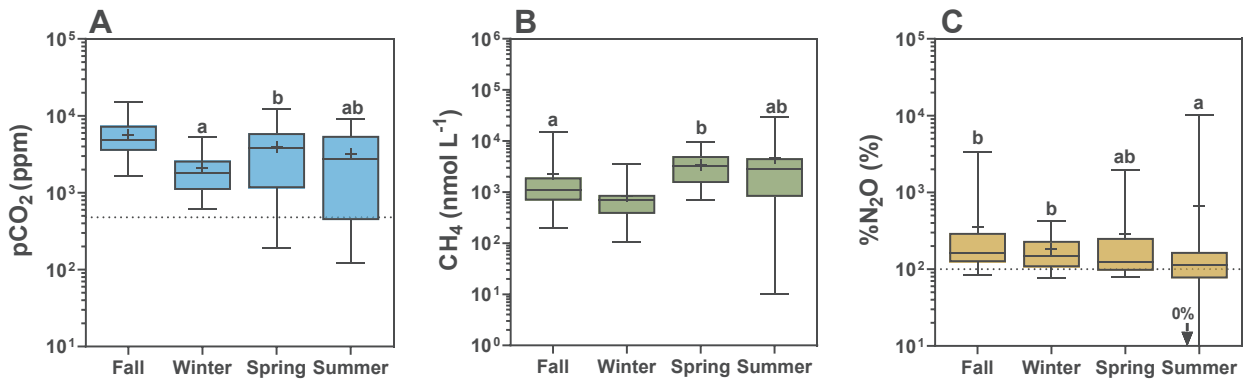


Figure 3 : Seasonal variations of partial pressure of CO₂ (pCO₂ in ppm) (A); dissolved CH₄ concentration (CH₄ in nmol L⁻¹) (B) and N₂O saturation level (%N₂O in %) (C) in 22 ponds in the city of Brussels during 4 seasons. Dashed lines indicate the equilibrium with atmosphere (493 ppm for pCO₂ (mean of measurements) and 100% for %N₂O). Box plots show median (horizontal line), mean (cross), and 25-75% percentiles (box limits). Whiskers extend from min to max. The common letters above the box plots indicate groups that are not significantly different (Permanova). Statistical results of Permanova and Betadisper are summarized in Table S5.

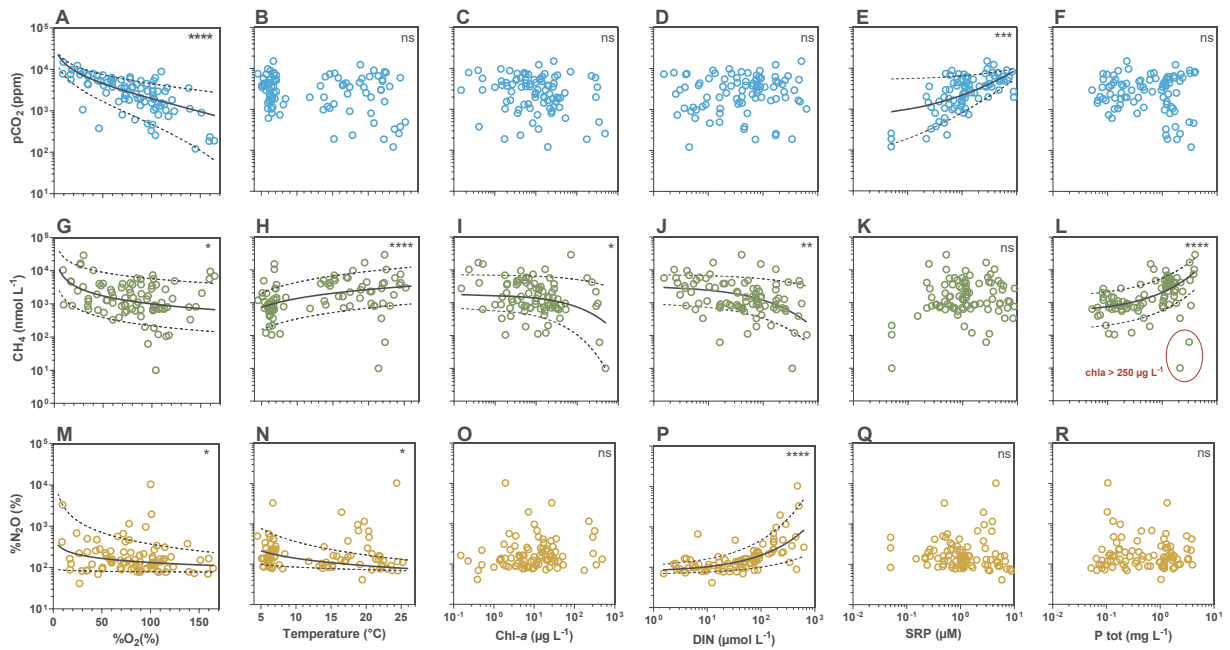


Figure 4 : Partial pressure of CO₂ (pCO₂, ppm), dissolved CH₄ concentration (CH₄, nmol L⁻¹), and N₂O saturation level (%N₂O, %), versus oxygen saturation level (%O₂, %), water temperature (°C), concentration of chlorophyll-a (Chl-a, in µg L⁻¹), concentration of dissolved inorganic nitrogen (DIN= NH₄⁺ + NO₂⁻ + NO₃⁻, in µmol L⁻¹), concentration of soluble reactive phosphorus (SRP, in µmol L⁻¹) and total phosphorus concentration (P tot, mg L⁻¹) in 22 ponds in the city of Brussels during 4 seasons. Median quantile regression shown as solid lines, 25th-75th quantile regression as dashed lines. Significance indicated by ns (non-significant) * (p<0.05), ** (p<0.01), ****(p<0.0001). Statistical results of quantile regressions are summarized in Table S7.

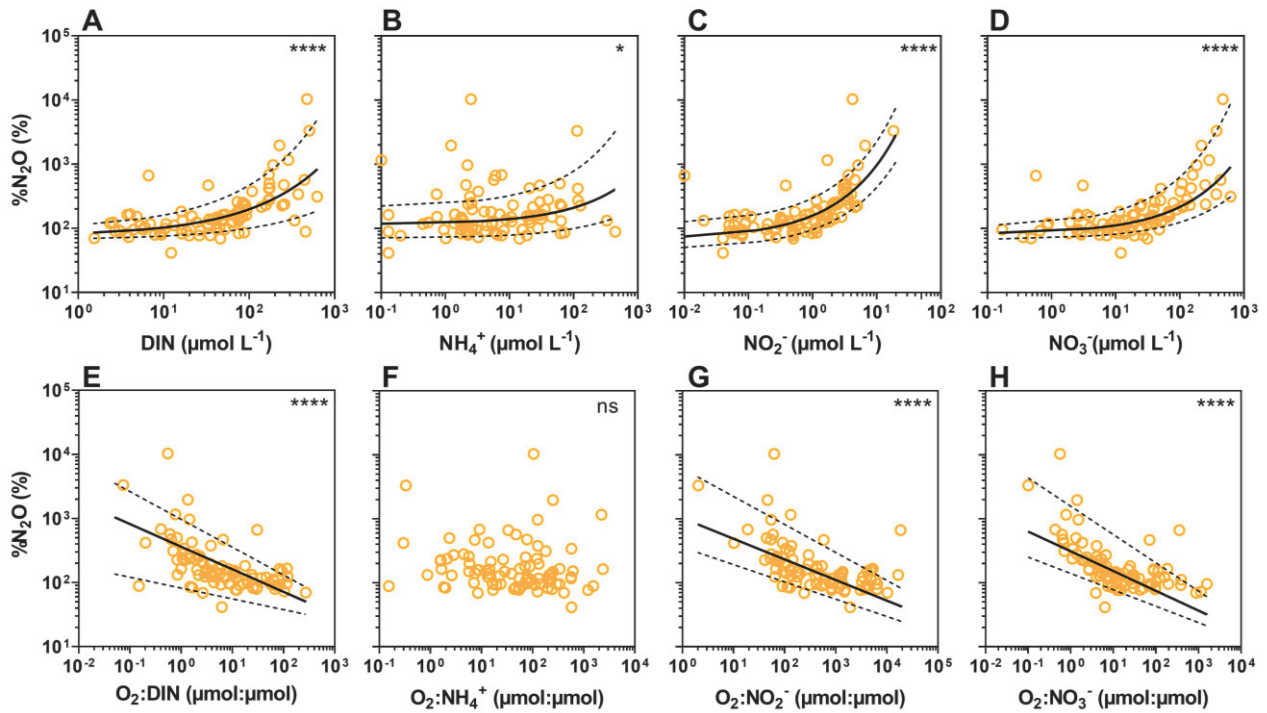


Figure 5 : N₂O saturation level (%N₂O, %) versus dissolved inorganic nitrogen (DIN= NH₄⁺ + NO₂⁻ + NO₃⁻, in μmol L⁻¹) (A), ammonium concentration (NH₄⁺, μmol L⁻¹) (B), nitrite concentration (NO₂⁻, μmol L⁻¹) (C), nitrate concentration (NO₃⁻, μmol L⁻¹) (D), ratio of dissolved inorganic nitrogen (DIN= NH₄⁺ + NO₂⁻ + NO₃⁻, in μmol L⁻¹) to dissolved oxygen (O₂, in μmol L⁻¹) (E), ratio of NH₄⁺ to O₂ (μmol:μmol) (F), ratio of NO₂⁻ to O₂ (μmol:μmol) (G), ratio of NO₃⁻ to O₂ (μmol:μmol) (H) in 22 sampled ponds in the city of Brussels during 4 seasons. Median quantile regression shown as solid lines, 25th-75th quantile regression as dashed lines. Significance indicated by * (p<0.05), ****(p<0.0001). Statistical results of quantile regressions are summarized in Table S6.

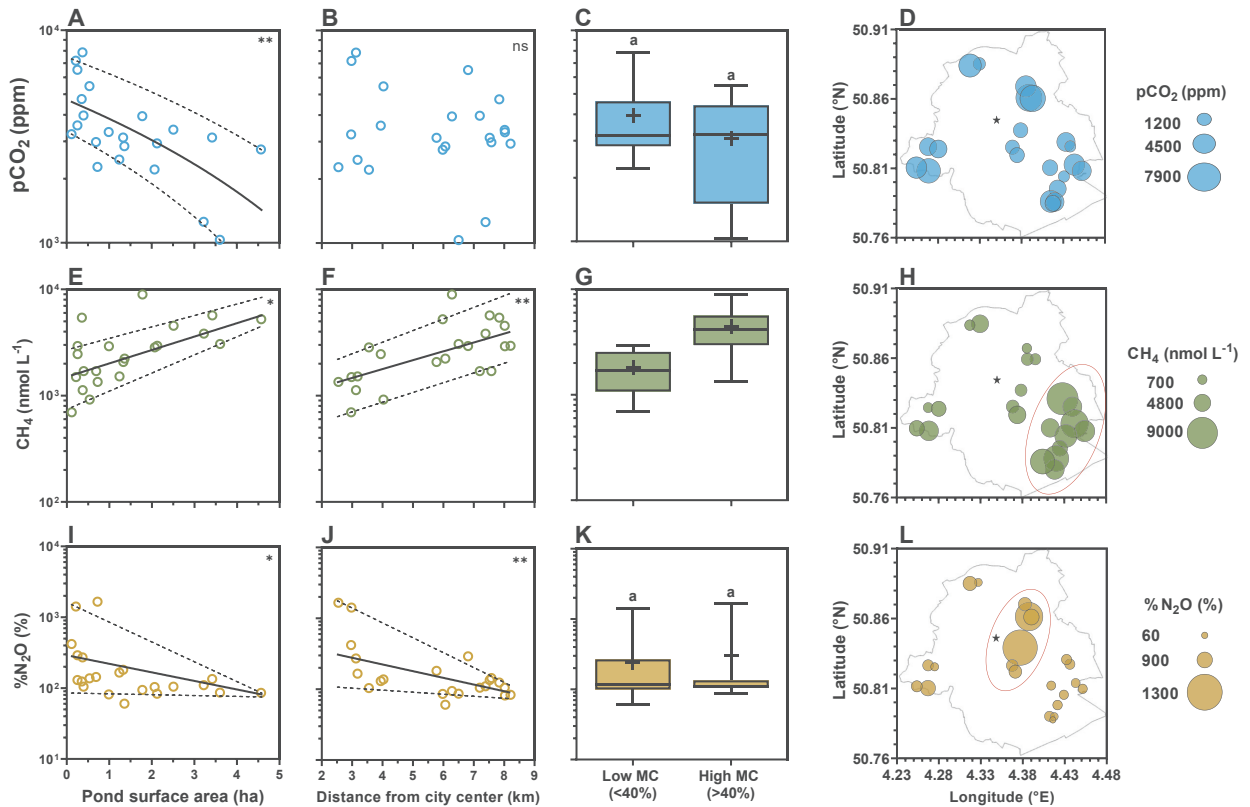


Figure 6 : Median quantile regressions of mean partial pressure of CO₂ (pCO₂, ppm), dissolved CH₄ concentration (CH₄, nmol L⁻¹), and N₂O saturation level (%N₂O, %) as function of surface area (A, E, I) and distance from city center (B,F, J) and boxplots of mean GHG with low (<40%, n=14) and high (>40%, n=8) macrophyte cover (MC) (C,G, K). Median quantile regression shown as solid lines, 25th-75th quantile regression as dashed lines. Significance indicated by ns (non-significant) * (p<0.05), ** (p<0.01). Box plots show median (horizontal line), mean (cross), and 25-75% percentiles (box limits). Whiskers extend from min to max. The common letters above the box plots indicate groups that are not significantly different (Permanova). Graphs D, H, L represent respectively mean pCO₂ (ppm), CH₄, (nmol L⁻¹) and %N₂O (%) for each pond during the four spatial surveys plotted against coordinates (°E, °N). City perimeter shown in light grey, city center (50.8504°N, 4.3487°E) marked with star. Red circles evidence zones with highest values of CH₄ concentrations and %N₂O. Statistical results of Permanova and Betadisper are summarized in Table S6. Statistical results of quantile regressions are summarized in Table S7.

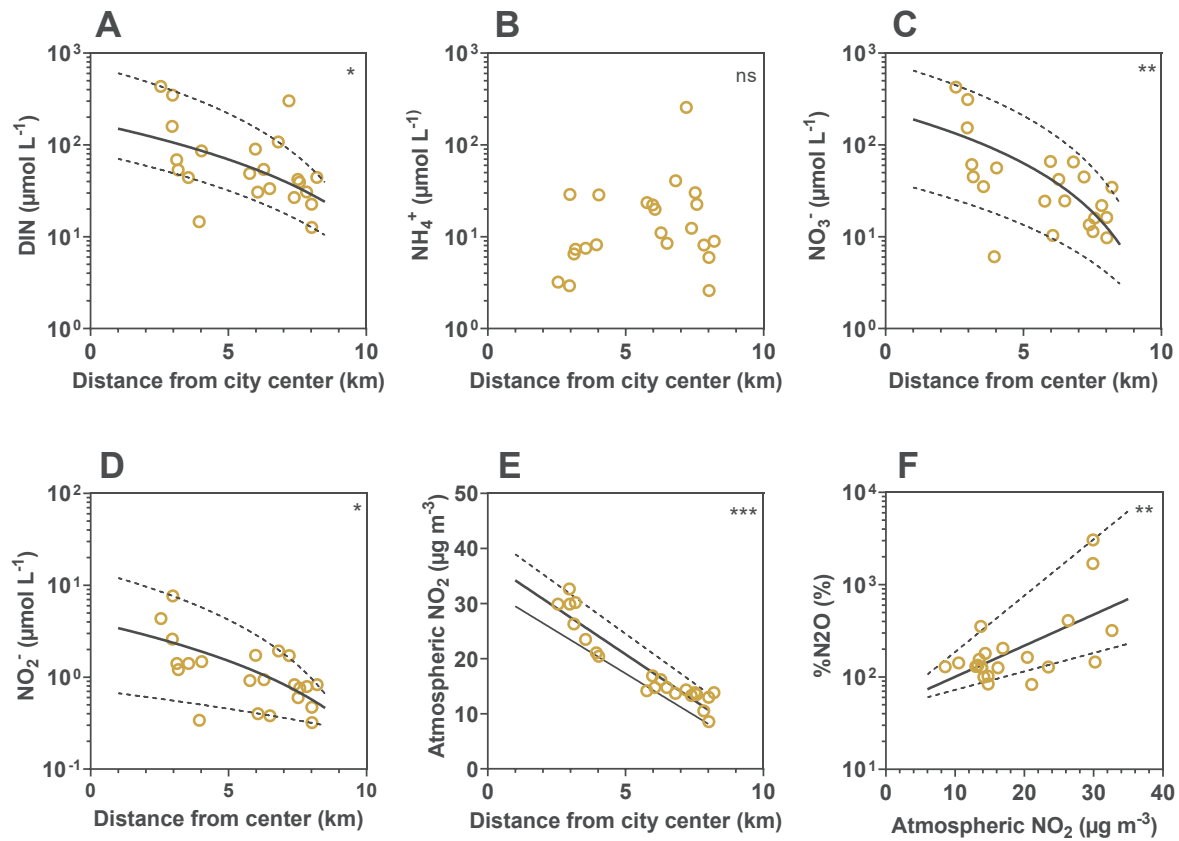


Figure 7 : Mean dissolved inorganic nitrogen (DIN= $\text{NH}_4^+ + \text{NO}_2^- + \text{NO}_3^-$, in $\mu\text{mol L}^{-1}$) (A), ammonium (NH_4^+ , in $\mu\text{mol L}^{-1}$) (B), nitrite (NO_2^- , in $\mu\text{mol L}^{-1}$) (C), nitrate (NO_3^- , in $\mu\text{mol L}^{-1}$) (D), and atmospheric NO_2 ($\mu\text{g m}^{-3}$) (E) concentrations as function of the distance from city center (50.8504°N , 4.3487°E) in 22 ponds in the city of Brussels during 4 seasons. N_2O saturation level (% N_2O , %) as function of atmospheric NO_2 concentration ($\mu\text{g m}^{-3}$) (F). Median quantile regression shown as solid lines, 25th-75th quantile regression as dashed lines. Significance (ns non-significant, * $p < 0.05$, ** $p < 0.01$) indicated top right. Statistical results of quantile regressions are summarized in Table S6. The atmospheric nitrogen dioxide (NO_2) concentration ($\mu\text{g m}^{-3}$) was extracted from the Curieuzenair initiative, which analyzed 2483 air samples in September 2021 covering the whole of the city of Brussels with a homogeneous distribution (<https://curieuzenair.brussels/en/the-results/>). Each data point corresponds to the average value of the six atmospheric NO_2 stations closest to each sampled pond, a distance between 50 and 450m.

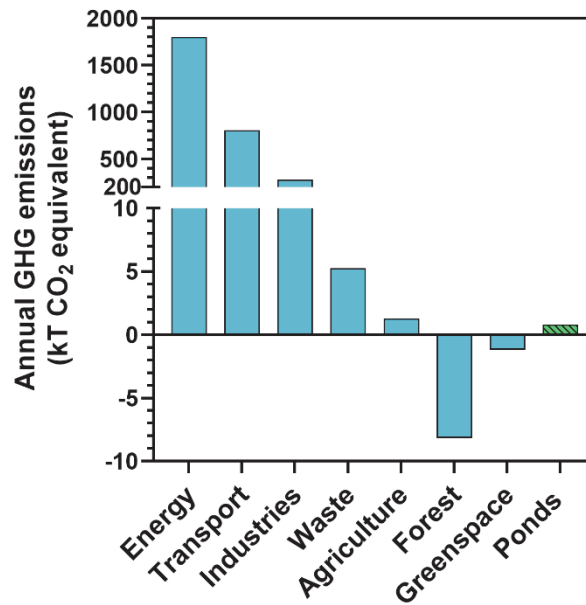


Figure 8 : Official inventory of annual emissions of greenhouse gases (kT CO₂ equivalents per year) for the city of Brussels in 2020 (<https://environnement.brussels>) Energy (residential/tertiary buildings), Transport (road), Industry (industrial processes and product use), Waste (mainly wastewater treatment), Agriculture (few soils), Forest (mainly Sonian forest), greenspace (private and public gardens, grassland) and diffusive emissions estimated from Brussels ponds (Ponds) in this study (2021-2022). The specific budget for each GHG is given in Table S4.

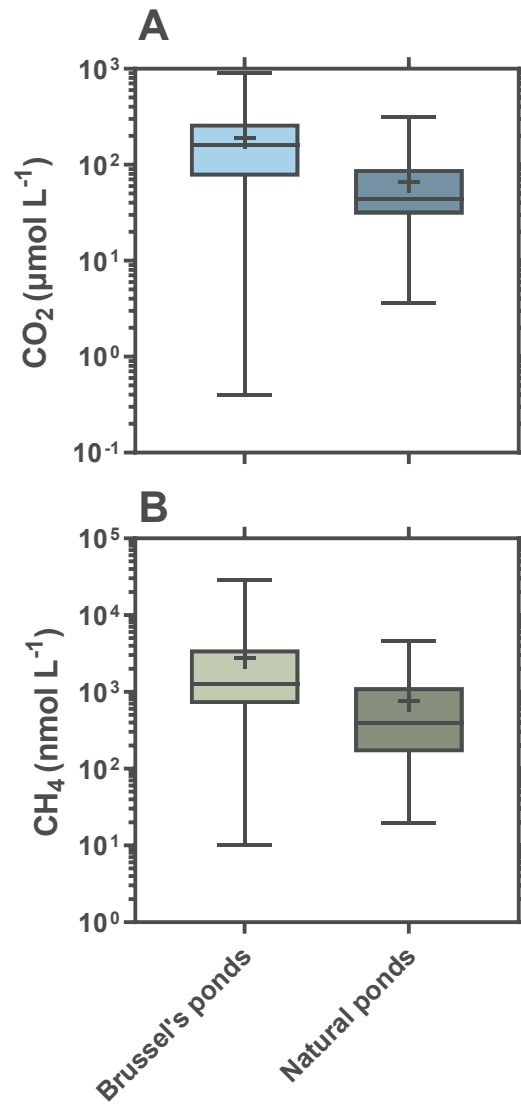


Figure 9: dissolved CO₂ concentration (CO₂, μmol L⁻¹) (A) and dissolved CH₄ concentration (CH₄, nmol L⁻¹) (B) in 22 sampled ponds in Brussels during 4 seasons (n=88) and natural ponds of similar size reported by Holgerson and Raymond (2016) (n=81 for CO₂, n=53 for CH₄). Box plots show median (horizontal line), mean (cross), and 25-75% percentiles (box limits). Whiskers extend from min to max. Both comparisons are significant based on Permanova test. Statistical results of Permanova and Betadisper are summarized in Table S5.

Seasonal and spatial variations of greenhouse gas (CO₂, CH₄ and N₂O) emissions from urban ponds in Brussels

(Supplementary Tables and figures)

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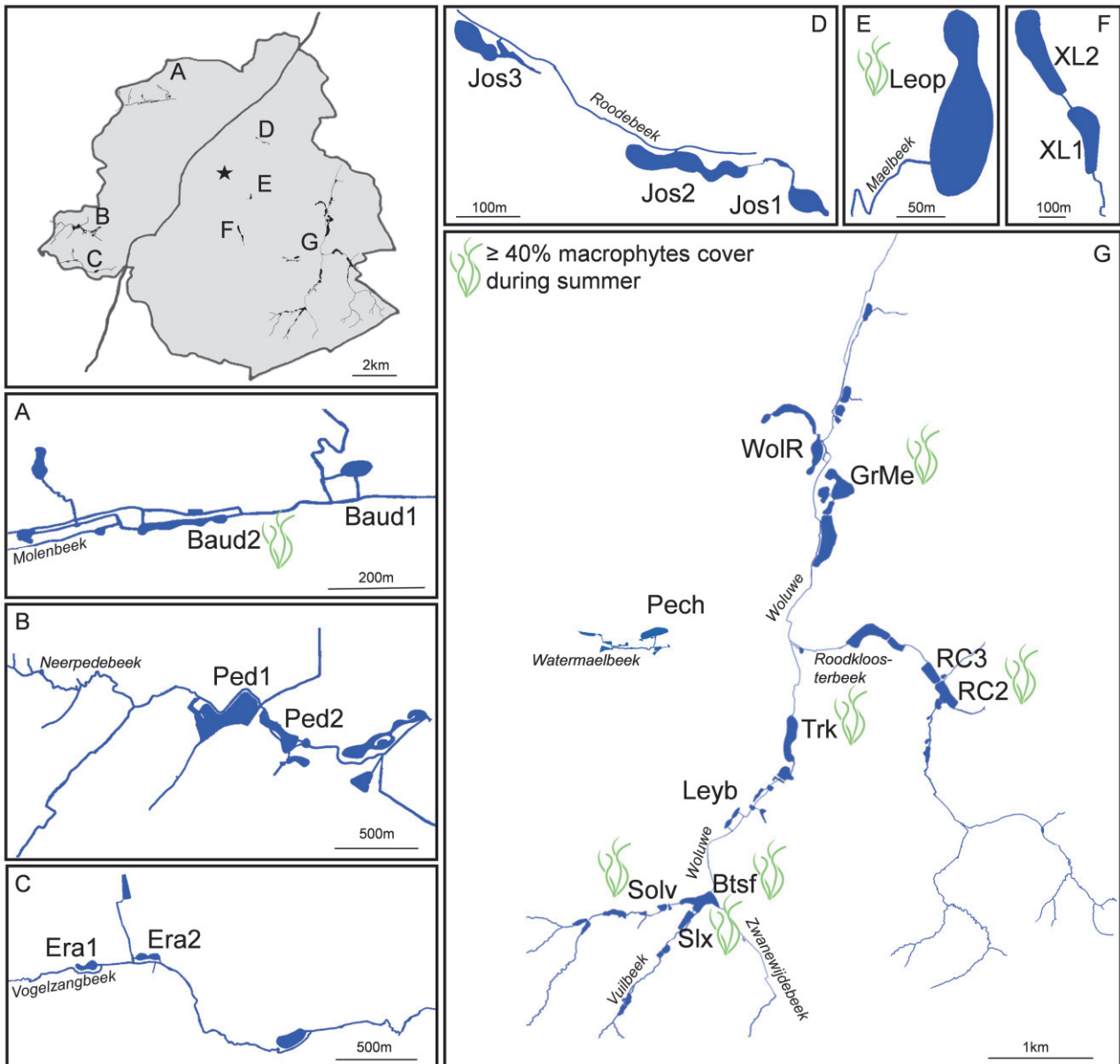


Figure S1 : Map of the city of Brussels and the sampled ponds. The acronyms correspond to the official names of the ponds of Bruxelles Environnement, the regional water management agency (Baud 1 : Etang Parc Roi Baudouin Phase I ; Baud2 : Etang Parc Roi Baudouin Phase II ; Btsf : Etang de Boitsfort ; Era 1 : Etang Erasme amont ; Era 2 : Etang Erasme aval ; GrMe : Grand étang Mellaerts ; Jos1 : Parc Josaphat - Etang aux canards ; Jos2 : Parc Josaphat – Etang aux pigeons ; Jos3 : Parc Josaphat – Etang de la laiterie ; Leop : Etang parc Leopold ; Leyb : Etang du Leybeek ; Pech : Etang Pêcheries Royales ; Ped1 : Etang de la Pède Grand Etang ; Ped2 : Etang de la Pède ; RC2 : Grand étang des Clabots (Etang 2) ; RC3 : Etang du moulin (Etang 3) ; Six : Etang du Silex ; Solv : Grand étang – Parc Tournay Solvay ; Trk : Etang Parc Ten Reuken ; WolR : Etang rond – Parc de Woluwe ; XL1 : Etang d’Ixelles Nord ; XL2 : Etang d’Ixelles Sud). The stream names are italicized. The city center (50.8504°N, 4.3487°E) is marked with star.

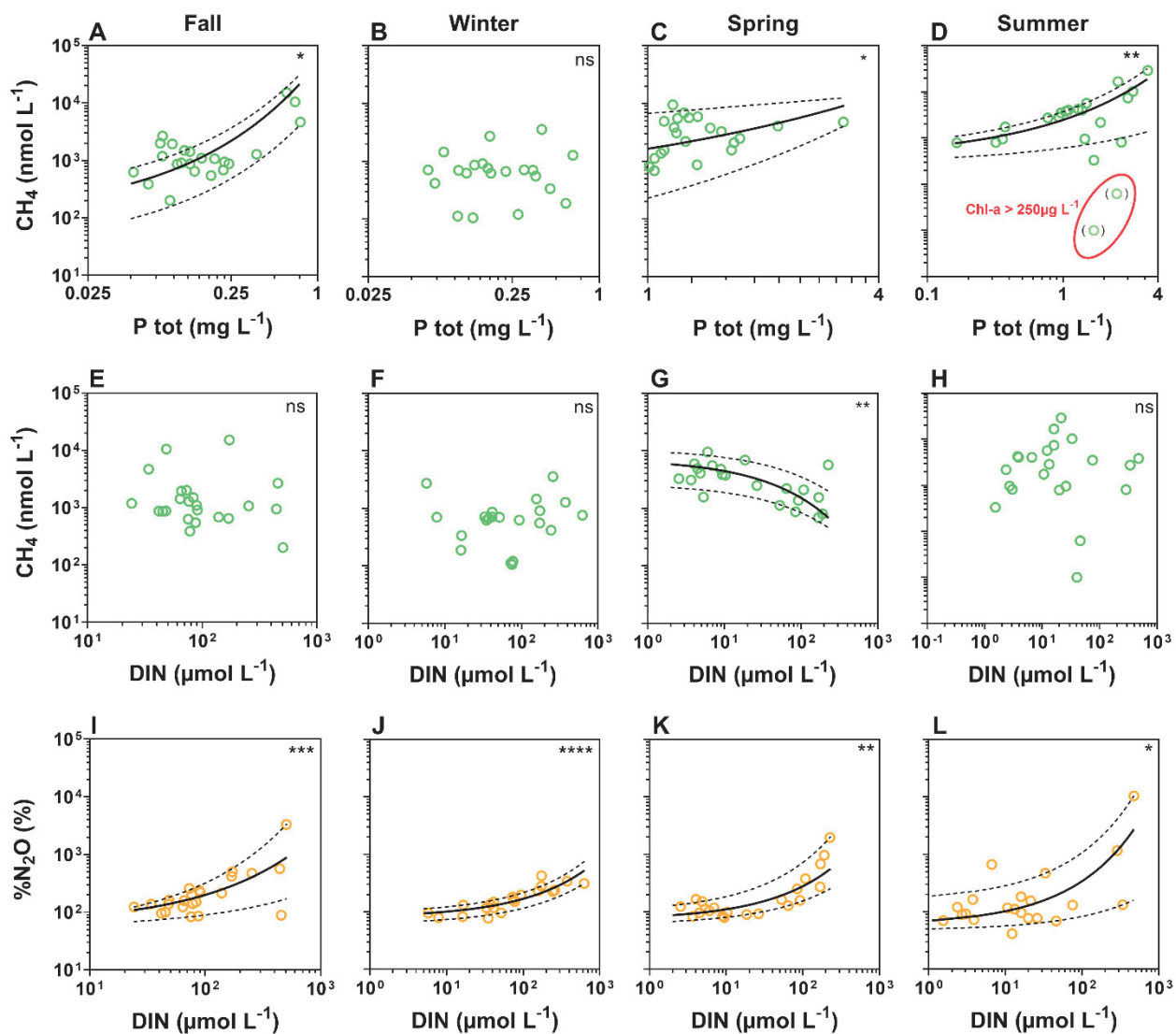


Figure S2: Dissolved CH₄ concentration (CH₄, nmol L⁻¹) versus the total phosphorus concentration (P tot, mg L⁻¹) (A,B,C,D) and versus dissolved inorganic nitrogen (DIN= NH₄⁺ + NO₂⁻ + NO₃⁻, in μmol L⁻¹) (E,F,G,H), N₂O saturation level (%N₂O, %) versus dissolved inorganic nitrogen (DIN= NH₄⁺ + NO₂⁻ + NO₃⁻, in μmol L⁻¹) (I,J,K,L) in 22 sampled ponds in the city of Brussels during 4 seasons. Median quantile regression shown as solid lines, 25th-75th quantile regression as dashed lines. Significance indicated by ns (non-significant), * (p<0.05), ** (p<0.01), *** (p<0.001), **** (p<0.0001). Statistical results of quantile regressions are summarized in Table S6.

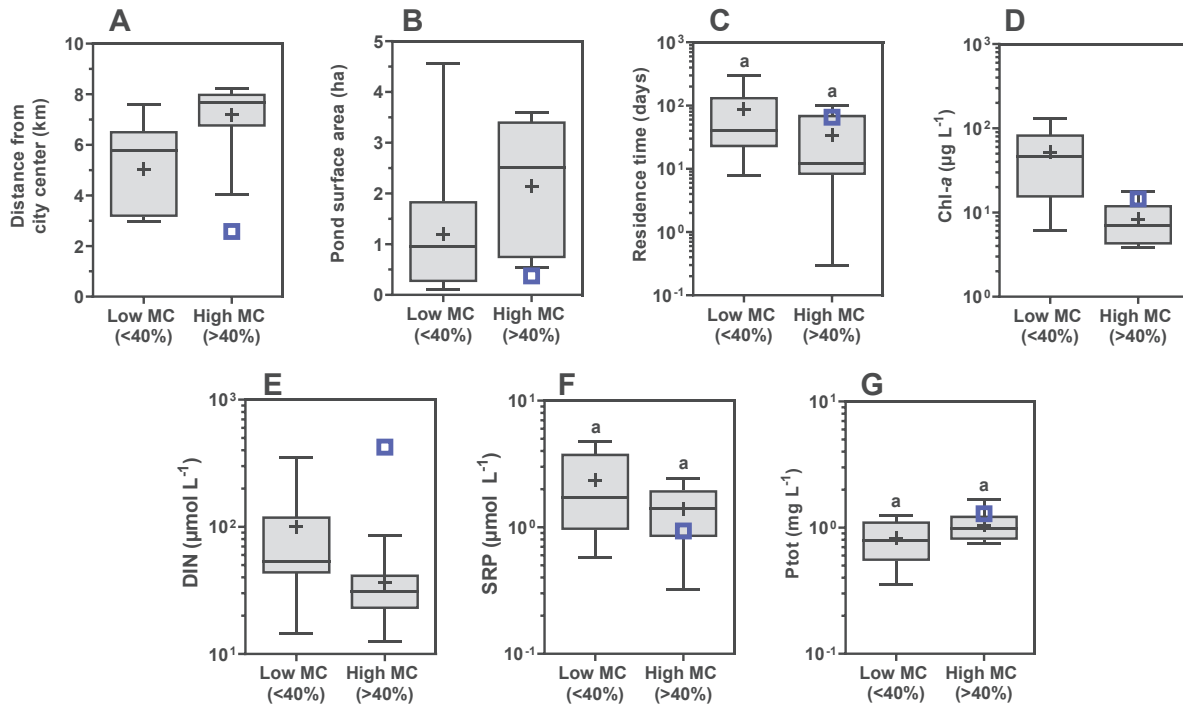


Figure S3: Distance from city center (km) (A), pond surface (ha) (B), pond water residence time (days) (C), mean chlorophyll-a ($\mu\text{g L}^{-1}$) (D), dissolved inorganic nitrogen (DIN= NH_4^+ + NO_2^- + NO_3^- , in $\mu\text{mol L}^{-1}$) (E), soluble reactive phosphorus (SRP, in $\mu\text{mol L}^{-1}$) (F), total phosphorus (Ptot, in mg L^{-1}) (G), in 22 ponds with macrophyte cover below and above 40% during four seasons in the city of Brussels. Leopold Pond is indicated by an empty blue square and was excluded from the analysis due to extreme values for some variables. Box plots show median (horizontal line), mean (cross), and 25-75% percentiles (box limits). Whiskers extend from min to max. The common letters above the box plots indicate groups that are not significantly different (Permanova). Statistical results of Permanova and Betadisper are summarized in Table S6.

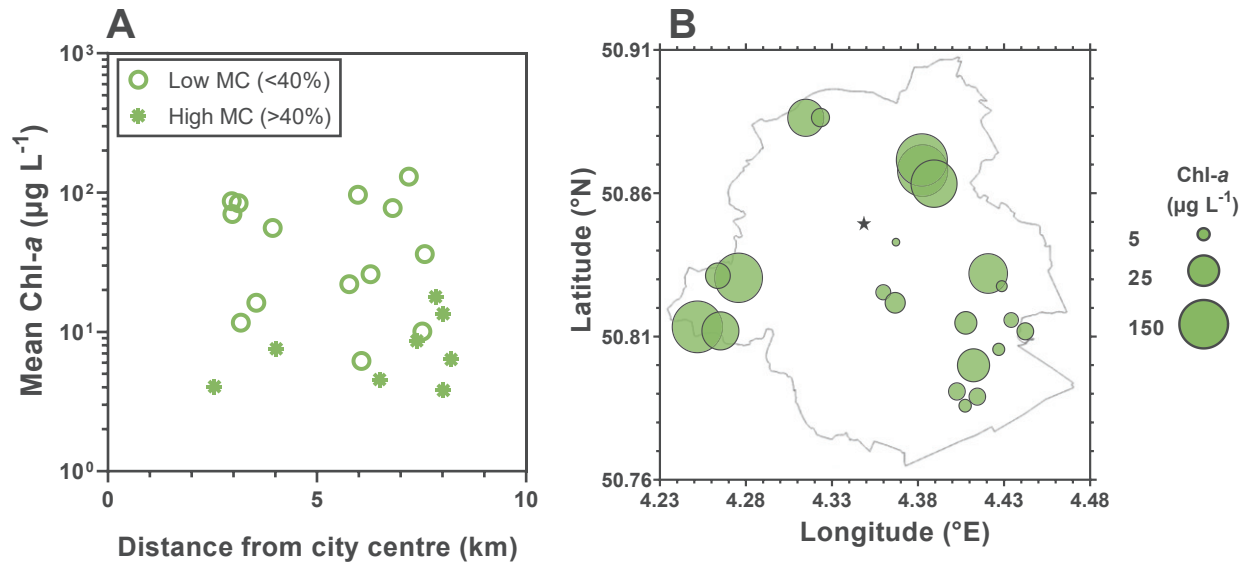


Figure S4: Mean chlorophyll-a concentration (Mean Chl-a, in $\mu\text{g L}^{-1}$) in 22 sampled ponds during the four seasons in the city of Brussels as a function of the distance from city center (50.8504°N , 4.3487°E) (A) and on a map showing the city perimeter (line) and center (star, 50.8504°N , 4.3487°E) (B).

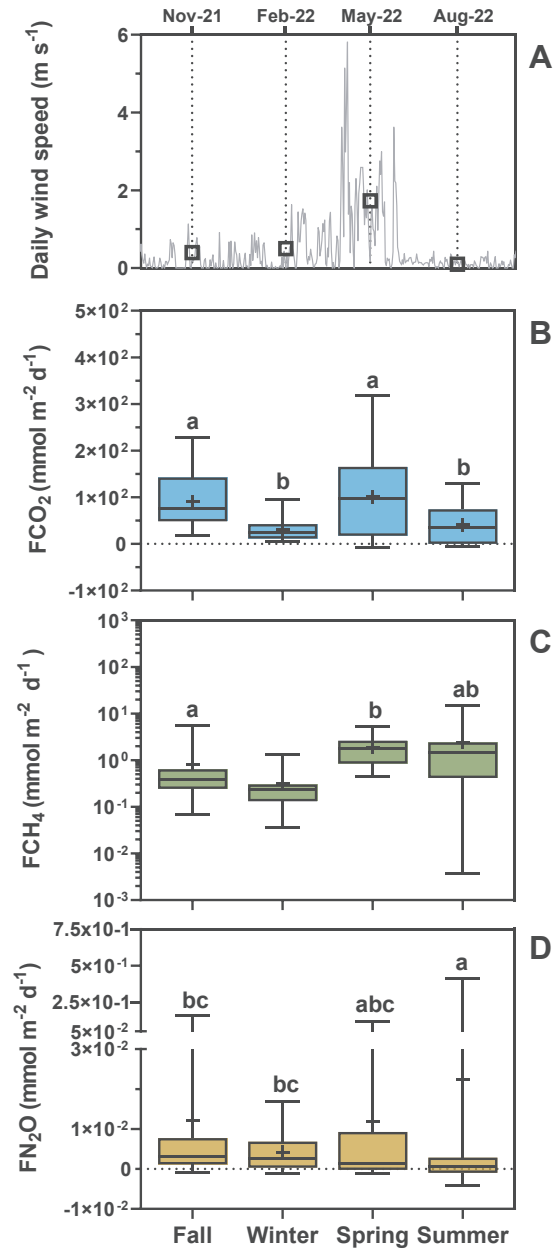


Figure S5 : Daily wind speed (dark grey, m s⁻¹) obtained from crowdsourcing platform of the Royal Belgian Meteorological Institute (wow.meteo.be) at the Brussels – Woluwe Saint Lambert station (50.8408 °N, 4.4234 °E) (A). squares indicate the mean wind for all samplings during a campaign. Seasonal comparisons of diffusive fluxes of CO₂ (B), CH₄ (C) and N₂O (D) (mmol m⁻² d⁻¹) in 22 ponds in the city of Brussels during 4 seasons. Box plots show median (horizontal line), mean (cross), and 25-75% percentiles (box limits). Whiskers extend from min to max. The common letters above the box plots indicate groups that are not significantly different (Permanova). Statistical results of Permanova and Betadisper are summarized in Table S4.

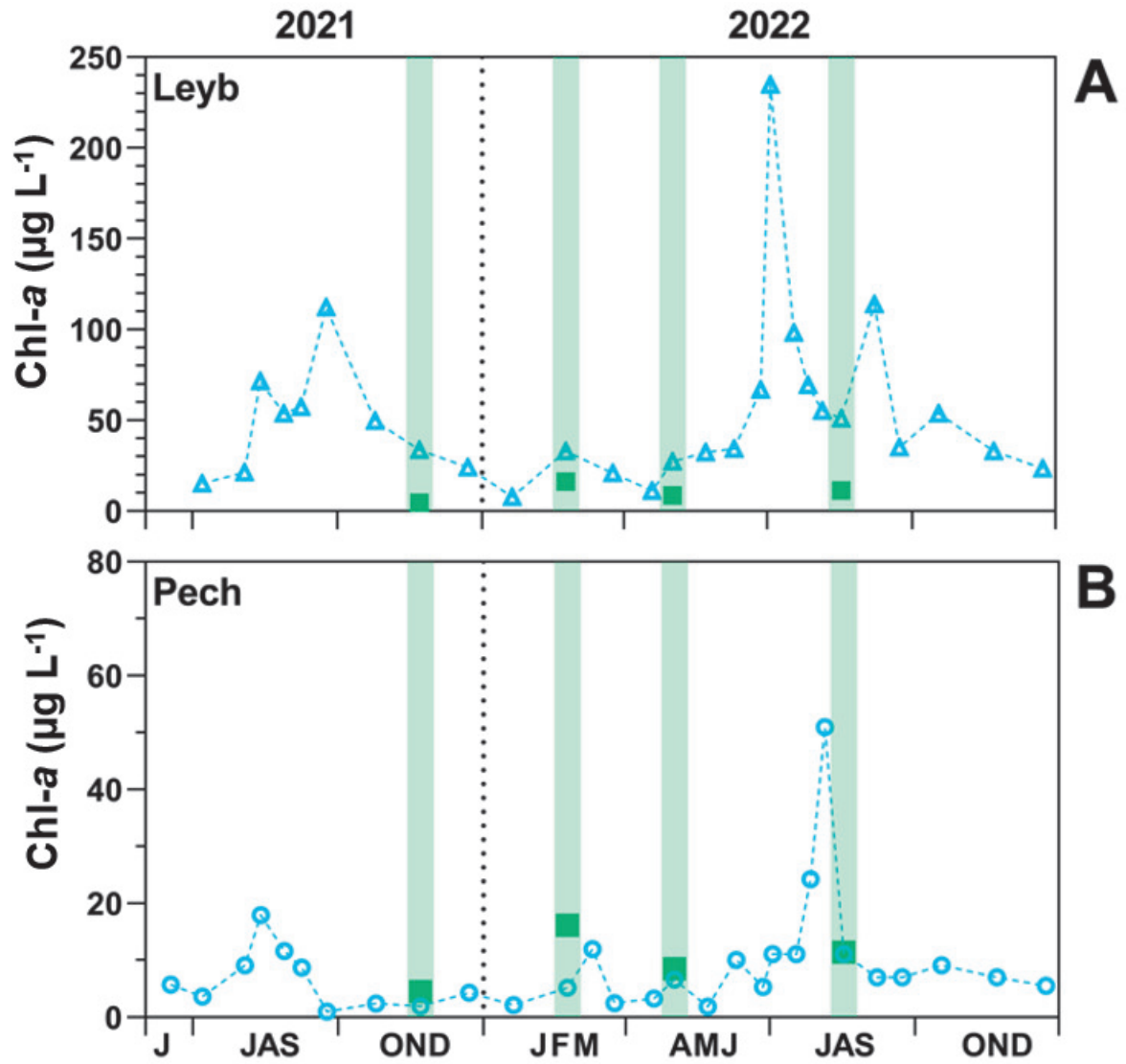


Figure S6: Chlorophyll-a concentration in (Chl-a, in $\mu\text{g L}^{-1}$) in the Leybeek (Leyb) (A) and Pêcheries (Pech) (B) ponds from June 2021 to December 2022. The solid squares indicate the median values of the 22 ponds sampled at four different seasons.

Table S1: Name, acronym, catchment, geographical coordinates, surface area, depth, perimeter, residence time, distance from the city center (50.8504°N, 4.3487°E), summer macrophyte cover, over-flow or flow-through regime, fishing activity, and storm-water regulation in 22 ponds in the city of Brussels. Surface area and perimeter data were calculated from satellite images in Google Earth and the other data were obtained from data resources from Brussels Environment, the regional water management agency.

Pond name (in French, according to Bruxelles Environnement)	Abbreviation	Catchment	Latitude (°N)	Longitude (°E)	Average depth (m)	Surface (ha)	Perimeter (m)	Residence time (days)	Distance from city center (km)	Summer macrophyte cover (%)	Regime	Fishing pond	Build as Stormwater Pond
Etang Parc Roi Baudouin Phase I (Sacré Cœur)	Baud1	Molenbeek	50.8833	4.3276	0.8	0.3	211	30.0	3.9	25	Overflow	No	No
Etang Parc Roi Baudouin Phase II - Sud	Baud2	Molenbeek	50.8813	4.3185	0.8	0.5	671	100.0	4.0	60	Overflow	No	No
Etang de Boitsfort ou du Moulin	Btsf	Woluwe	50.7930	4.4171	0.3	2.5	904	7.0	8.0	60	Flow-through	No	No
Etang Erasme Amont	Era1	Neerpedebeek	50.8118	4.2666	0.7	0.4	316	45.0	7.2	25	Overflow	No	Yes
Etang Erasme Aval	Era2	Neerpedebeek	50.8123	4.2722	0.7	0.3	283	30.0	6.8	5	Overflow	No	Yes
Grand étang Mellaerts	GrMe	Woluwe	50.8274	4.4347	1.2	3.6	939	78.0	6.5	50	Overflow	No	No
Parc Josaphat - Etang aux canards	Jos1	Maelbeek	50.8606	4.3903	0.5	0.4	438	115.0	3.1	0	Overflow	No	No
Parc Josaphat - Etang aux pigeons	Jos2	Maelbeek	50.8611	4.3878	0.6	0.2	206	66.4	3.0	15	Overflow	No	No
Parc Josaphat - Etang de la laiterie	Jos3	Maelbeek	50.8629	4.3830	0.7	0.1	481	35.9	3.0	15	Overflow	No	No
Etang Parc Leopold	Leop	Maelbeek	50.8390	4.3802	1.0	0.7	515	150.0	2.5	50	Overflow	No	No
Etang Parc du Leybeek	Leyb	Woluwe	50.8000	4.4210	0.6	0.7	511	300.0	7.6	0	Overflow	No	No
Etang Pêcheries Royales - Parc de la Héronnière	Pech	Woluwe	50.8127	4.4119	1.3	1.4	558	7.8	6.1	15	Flow-through	Yes	No
Etang de la Pède - Grand Etang	Ped1	Neerpedebeek	50.8262	4.2726	1.3	4.6	1084	25.0	6.0	30	Overflow	No	Yes
Etang de la Pède	Ped2	Neerpedebeek	50.8254	4.2766	0.7	1.3	713	100.0	5.8	20	Overflow	No	Yes
Grand étang des Clabots (Etang 2) - Rouge-Cloître	RC2	Woluwe	50.8105	4.4477	2.0	2.1	664	13.0	8.2	25	Flow-through	No	No
Etang du Moulin (Etang 3) - Rouge-Cloître	RC3	Woluwe	50.8118	4.4463	1.1	3.4	989	11.0	7.5	40	Flow-through	Yes	No
Etang du Silex	Six	Woluwe	50.7917	4.4151	1.1	1.0	464	0.3	8.0	90	Flow-through	No	No
Grand étang - Parc Tournay-Solvay	Solv	Woluwe	50.7925	4.4123	1.4	0.4	331	12.0	7.8	80	Flow-through	No	No
Etang Parc Ten Reuken	Trk	Woluwe	50.8066	4.4278	1.4	3.2	1148	12.0	7.4	80	Flow-through	No	No
Etang rond - Parc de Woluwe	WolR	Woluwe	50.8300	4.4317	1.4	1.8	600	10.7	6.3	15	Overflow	No	No
Etang d'Ixelles Nord	XL1	Maelbeek	50.8229	4.3743	1.8	2.1	774	200.0	3.5	20	Flow-through	Yes	No
Etang d'Ixelles Sud	XL2	Maelbeek	50.8257	4.3716	1.2	1.2	589	250.0	3.2	15	Flow-through	Yes	No

Table S2: Mean, standard deviation (SD), minimal value (min), maximal value (max), median and interquartile range (IQR) and number of samples (n) of Water temperature (in °C), pH, conductivity (in $\mu\text{S cm}^{-1}$), Oxygen saturation (in %), Ammonium, Nitrite, Nitrate, Soluble reactive phosphorus (SRP) concentrations (in $\mu\text{mol L}^{-1}$), total phosphorus concentration (in mg L^{-1}), Chlorophyll-a concentration (in $\mu\text{g L}^{-1}$), total suspended matter concentration (in mg L^{-1}), partial pressure of CO_2 (pCO_2 in ppm); dissolved CH_4 concentration (CH_4 in nmol L^{-1}) and N_2O saturation level (in %) measured at 22 ponds in the city of Brussels during 4 seasons. Statistical results of Permanova and Betadisper for comparisons between seasons are summarized in Table S4.

November 2021					
Parameter	Units	mean \pm SD	(min;max)	median (IQR)	n
Water temperature	°C	6.1 \pm 0.7	(4.5 ; 7.1)	6.2 (5.5 ; 6.6)	22
pH		7.51 \pm 0.42	(7.12 ; 8.23)	7.39 (7.19 ; 7.74)	22
Conductivity	$\mu\text{S cm}^{-1}$	858 \pm 256	(599 ; 1507)	795 (675 ; 918)	22
Oxygen saturation	%	52 \pm 29	(9 ; 110)	51 (34.6 ; 70.9)	22
Ammonium	$\mu\text{mol L}^{-1}$	57.7 \pm 90.4	(2.1 ; 447.0)	29.8 (20.4 ; 62.6)	22
Nitrite	$\mu\text{mol L}^{-1}$	2.7 \pm 3.6	(0.6 ; 18.3)	1.7 (1.2 ; 2.9)	22
Nitrate	$\mu\text{mol L}^{-1}$	77.2 \pm 115.4	(9.3 ; 432.8)	34.6 (21.1 ; 47.4)	22
SRP	$\mu\text{mol L}^{-1}$	1.8 \pm 1.2	(0.5 ; 4.8)	1.2 (1.0 ; 2.8)	22
Total phosphorus	mg L^{-1}	0.1 \pm 0.2	(0.1 ; 0.8)	0.1 (0.1 ; 0.3)	22
Chlorophyll-a	$\mu\text{g L}^{-1}$	9.5 \pm 10.0	(0.5 ; 33.6)	4.5 (2.1 ; 15.2)	22
Total suspended matter	mg L^{-1}	8.3 \pm 14.2	(1.2 ; 72.4)	4.6 (3.9 ; 7.6)	22
pCO_2	ppm	5453 \pm 3058	(1655 ; 15029)	4650 (3568 ; 7094)	22
CH_4	nmol L^{-1}	2260 \pm 3523	(201 ; 15132)	1073 (773 ; 1714)	22
N_2O saturation	%	360 \pm 677	(84 ; 3324)	161 (123 ; 247)	22
February 2022					
Water temperature	°C	6.9 \pm 1.3	(5.6 ; 11.8)	6.5 (6.2 ; 7.1)	21
pH		8.11 \pm 0.47	(7.05 ; 8.75)	8.13 (7.93 ; 8.49)	21
Conductivity	$\mu\text{S cm}^{-1}$	796 \pm 265	(520 ; 1364)	685 (585 ; 942)	21
Oxygen saturation	%	96 \pm 23	(48 ; 139)	102 (80 ; 111)	21
Ammonium	$\mu\text{mol L}^{-1}$	12.3 \pm 27.9	(0.7 ; 127.6)	2.6 (1.9 ; 7.5)	21
Nitrite	$\mu\text{mol L}^{-1}$	1.6 \pm 1.5	(0.1 ; 5.0)	1.1 (0.5 ; 3.1)	21
Nitrate	$\mu\text{mol L}^{-1}$	110.3 \pm 146.6	(4.6 ; 619.3)	65.2 (30.8 ; 129.5)	21
SRP	$\mu\text{mol L}^{-1}$	2.2 \pm 2.9	(0.2 ; 8.9)	0.6 (0.4 ; 2.0)	21
Total phosphorus	mg L^{-1}	0.3 \pm 0.2	(0.1 ; 0.7)	0.2 (0.2 ; 0.4)	21
Chlorophyll-a	$\mu\text{g L}^{-1}$	19.7 \pm 14.3	(1.8 ; 52.8)	16.2 (7.2 ; 30.7)	21
Total suspended matter	mg L^{-1}	9.5 \pm 4.5	(3.7 ; 20.3)	9.2 (5.7 ; 11.1)	21
pCO_2	ppm	2064 \pm 1295	(610 ; 5283)	1822 (1121 ; 2524)	21
CH_4	nmol L^{-1}	860 \pm 841	(105 ; 3552)	697 (416 ; 861)	21
N_2O saturation	%	182 \pm 95	(77 ; 425)	150 (117 ; 231)	20
May 2022					
Water temperature	°C	16.2 \pm 2.0	(13.3 ; 20.2)	16.1 (14.6 ; 17.4)	22
pH		8.06 \pm 0.50	(7.37 ; 9.12)	7.91 (7.72 ; 8.44)	22
Conductivity	$\mu\text{S cm}^{-1}$	776 \pm 272	(402 ; 1424)	709 (604 ; 835)	22
Oxygen saturation	%	90 \pm 41	(17 ; 164)	87 (61 ; 104)	22
Ammonium	$\mu\text{mol L}^{-1}$	10.9 \pm 25.0	(0.5 ; 117.6)	2.9 (1.8 ; 7.5)	22
Nitrite	$\mu\text{mol L}^{-1}$	1.3 \pm 1.8	(0.1 ; 6.6)	0.4 (0.1 ; 1.8)	22
Nitrate	$\mu\text{mol L}^{-1}$	45.5 \pm 64.9	(0.9 ; 218.0)	10.7 (2.9 ; 69.5)	22
SRP	$\mu\text{mol L}^{-1}$	1.6 \pm 1.5	(0.2 ; 5.6)	1.0 (0.8 ; 1.2)	22
Total phosphorus	mg L^{-1}	1.4 \pm 0.5	(1.0 ; 3.3)	1.3 (1.1 ; 1.6)	22
Chlorophyll-a	$\mu\text{g L}^{-1}$	11.9 \pm 8.8	(1.7 ; 29.0)	8.5 (5.2 ; 19.2)	22
Total suspended matter	mg L^{-1}	10.7 \pm 7.4	(3.2 ; 38.5)	8.3 (7.3 ; 12.1)	22
pCO_2	ppm	3892 \pm 2993	(190 ; 12329)	3787 (1218 ; 5805)	22
CH_4	nmol L^{-1}	3456 \pm 2301	(688 ; 9507)	3180 (1546 ; 4790)	22
N_2O saturation	%	289 \pm 432	(79 ; 1963)	125 (96 ; 250)	22
August 2022					
Water temperature	°C	22.2 \pm 1.6	(19.4 ; 25.2)	22.1 (20.8 ; 23.3)	22
pH		7.99 \pm 0.56	(7.08 ; 9.10)	7.90 (7.65 ; 8.25)	22
Conductivity	$\mu\text{S cm}^{-1}$	667 \pm 270	(435 ; 1557)	537 (499 ; 792)	22
Oxygen saturation	%	87 \pm 33	(27 ; 159)	93 (71 ; 101)	22
Ammonium	$\mu\text{mol L}^{-1}$	23.7 \pm 70.1	(0.1 ; 330.6)	3.0 (0.7 ; 15.0)	22
Nitrite	$\mu\text{mol L}^{-1}$	0.4 \pm 0.9	(0.0 ; 4.2)	0.1 (0.0 ; 0.4)	22
Nitrate	$\mu\text{mol L}^{-1}$	42.1 \pm 112.4	(0.2 ; 467.7)	7.8 (1.7 ; 11.8)	22
SRP	$\mu\text{mol L}^{-1}$	2.6 \pm 2.6	(0.1 ; 8.4)	1.4 (0.4 ; 4.2)	22
Total phosphorus	mg L^{-1}	2.0 \pm 1.4	(0.2 ; 5.3)	1.8 (1.2 ; 2.8)	22
Chlorophyll-a	$\mu\text{g L}^{-1}$	105.9 \pm 149.9	(0.1 ; 488.4)	11.4 (2.8 ; 201.9)	22
Total suspended matter	mg L^{-1}	33.2 \pm 37.4	(3.2 ; 110.2)	12.1 (6.1 ; 59.3)	22
pCO_2	ppm	3185 \pm 2902	(121 ; 8970)	2737 (469 ; 5218)	22
CH_4	nmol L^{-1}	4691 \pm 6713	(10 ; 29190)	2790 (883 ; 4956)	22
N_2O saturation	%	656 \pm 2182	(0 ; 10354)	113 (77 ; 159)	22
All seasons					
Water temperature	°C	12.8 \pm 6.9	(4.5 ; 25.2)	12.6 (6.4 ; 19.5)	87
pH		7.91 \pm 0.53	(7.05 ; 9.12)	7.87 (7.55 ; 8.20)	87
Conductivity	$\mu\text{S cm}^{-1}$	775 \pm 271	(402 ; 1557)	699 (580 ; 903)	87
Oxygen saturation	%	81 \pm 36	(9 ; 164)	83 (54 ; 103)	87
Ammonium	$\mu\text{mol L}^{-1}$	26.7 \pm 62.9	(0.1 ; 447.0)	5.6 (2.1 ; 24.0)	87
Nitrite	$\mu\text{mol L}^{-1}$	1.5 \pm 2.3	(0.0 ; 18.3)	0.8 (0.2 ; 2.1)	87
Nitrate	$\mu\text{mol L}^{-1}$	68.4 \pm 114.6	(0.2 ; 619.3)	24.8 (7.0 ; 66.7)	87
SRP	$\mu\text{mol L}^{-1}$	2.0 \pm 2.2	(0.1 ; 8.9)	1.1 (0.6 ; 2.9)	87
Total phosphorus	mg L^{-1}	1.0 \pm 1.1	(0.1 ; 5.3)	0.6 (0.15 ; 1.4)	87
Chlorophyll-a	$\mu\text{g L}^{-1}$	36.6 \pm 84.6	(0.1 ; 488.4)	10.0 (3.6 ; 24.9)	87
Total suspended matter	mg L^{-1}	15.5 \pm 22.8	(1.2 ; 110.2)	8.0 (5.2 ; 11.8)	87
pCO_2	ppm	3667 \pm 2904	(121 ; 15029)	3016 (1394 ; 5343)	87
CH_4	nmol L^{-1}	2833 \pm 4178	(10 ; 29190)	1276 (705 ; 3604)	87
N_2O saturation	%	273 \pm 662	(0 ; 10354)	138 (96 ; 232)	86

Table S3: Dissolved concentrations of CO₂, CH₄ and N₂O in urban ponds reported in literature and the present study in 22 urban ponds in the city of Brussels during 4 seasons.

Region	n	Coverage	CO ₂	CH ₄	N ₂ O	Reference
			($\mu\text{mol L}^{-1}$)	(nmol L^{-1})	(nmol L^{-1})	
Brussel (Belgium)	22	4 seasons	189 ± 155	2,745 ± 4,120	39 ± 95	This study
Brussel (Belgium)	22	Summer	123 ± 112	4,691 ± 6,713	8 ± 158	This study
Linköping (Sweden)	1	4 seasons	-	1,733	-	Natchimuthu et al. (2014)
Minnesota (United States)	7	Summer	6 ± 180	2,530 ± 32,360		Ray et al. (2023)
Minnesota (United States)	18	Summer	170 ± 180	15,794 ± 23,107	18 ± 12	Rabaey and Cotner (2022)
Salaspils (Latvia)	1	Summer	133 ± 17	4,914 ± 3,779		Ray et al. (2023)
Silkeborg (Denmark)	37	4 seasons	162 ± 184	3,667 ± 16,500	28 ± 64	Audet et al. (2020)
Silkeborg (Denmark)	37	Summer	202 ± 302	6,000 ± 32,200	-	Audet et al. (2020)
Uppsala (Sweden)	40	Summer	157 ± 153	4,528 ± 5,712	-	Peacock et al. (2019)
Uppsala (Sweden)	2	Summer	66 ± 74	804 ± 2,378		Ray et al. (2023)

Table S4 : Reported annual emissions of CO₂, CH₄, and N₂O in CO₂ equivalents from the Brussels region for the year 2020 (<https://environnement.brussels>) and the CO₂, CH₄ and N₂O emissions from the urban ponds (n=158, total area = 101ha), and their relative contribution. The CH₄ emissions have a diffusive component (Diff.) computed from the diffusive flux estimates in the 22 ponds during the 4 seasons, and an ebullitive component (Ebull.) computed from water temperature in the 22 ponds during the 4 seasons based on a relation between CH₄ ebullition and water temperature collected in 4 ponds (Leybeek, Pêcheries, Silex and Ten Reuken) from 2021-2023 (unpublished data).

		CO ₂	CH ₄	N ₂ O
		kT CO ₂ eq y ⁻¹		
Officially reported	Energy	1796.18	6.15	1.63
	Transport	799.01	0.73	6.87
	Industries	275.45	0.04	2.23
	Waste	0.00	0.29	4.98
	Agriculture	0.04	0.53	0.71
	Forest	-8.68	0.00	0.50
	Greenspace	-1.19	0.00	0.00
This study	Ponds	0.74	0.09 (Diff.) 0.21 (Ebull.)	0.01
		Relative contribution (%)		
Ponds (expressed as fraction of the total)		0.03	1.12 (Diff.) 2.61 (Ebull.)	0.05

Table S5: Outcomes of the Betadisper and Permanova statistical tests for pairs of seasons on the medians of Water temperature (Temperature, in °C), Oxygen saturation (%O₂, in %), Soluble reactive phosphorus (SRP, in μmol L⁻¹), total phosphorus concentration (P_{tot}, in mg L⁻¹), ammonium concentration (NH₄⁺, in μmol L⁻¹); nitrite concentration (NO₂⁻, in μmol L⁻¹), nitrate concentration (NO₃⁻, in μmol L⁻¹), dissolved inorganic nitrogen (DIN= NH₄⁺ + NO₂⁻ + NO₃⁻, in μmol L⁻¹), Chlorophyll-a concentration (Chl-a, in μg L⁻¹), total suspended matter concentration (TSM, in mg L⁻¹), dissolved CH₄ concentration (CH₄, in nmol L⁻¹), partial pressure of CO₂ (pCO₂, in ppm); N₂O saturation level (%N₂O, in %), diffusive fluxes of CO₂ (FCO₂), CH₄ (FCH₄) and N₂O (FN₂O) (mmol m⁻² d⁻¹). Transformations have been applied to the data to ensure a normal distribution, and are indicated in the Transformation column.

Figure or table	Variable	Transformation	p-values of the Permanova test					
			Fall vs Winter	Fall vs Spring	Fall vs Summer	Winter vs Spring	Winter vs Summer	Spring vs Summer
Tab. S2	Temperature (°C)	logarithmic	0.005 **	0.001 ***	0.001 ***	0.001 ***	0.001 ***	0.001 ***
	%O ₂ (%)	logarithmic	0.001 ***	0.004 **	0.003 **	0.241	0.170	0.925
	SRP (μmol L ⁻¹)	square root	0.847	0.486	0.610	0.771	0.615	0.342
	P _{tot} (mg L ⁻¹)	square root	0.452	0.001 ***	0.001 ***	0.001 ***	0.001 ***	0.172
	NH ₄ ⁺ (μmol L ⁻¹)	square root	0.002 **	0.001 ***	0.004 **	0.804	0.729	0.582
	NO ₂ ⁻ (μmol L ⁻¹)	square root	0.041 *	0.006 **	0.001 ***	0.218	0.001 ***	0.04 *
	NO ₃ ⁻ (μmol L ⁻¹)	square root	0.354	0.119	0.022 *	0.015 *	0.004 **	0.419
	DIN (μmol L ⁻¹)	square root	0.532	0.003 **	0.008 **	0.037 *	0.038 *	0.864
	Chl-a (μg L ⁻¹)	square root	0.007 **	0.076	0.088	0.100	0.928	0.413
TSM (mg L ⁻¹)	square root	0.039*	0.044*	0.003 **	0.333	0.106	0.237	
Fig. 3	CH ₄ (nmol L ⁻¹)	logarithmic	0.008 **	0.007 **	0.092	0.001 ***	0.014 *	0.368
	%N ₂ O (%)	logarithmic	0.262	0.477	0.024*	0.764	0.047*	0.422
	pCO ₂ (ppm)	square root	0.001 ***	0.031 *	0.002 **	0.039 *	0.373	0.337
Fig. S5	FCO ₂ (mmol m ⁻² d ⁻¹)	logarithmic (value+median)	0.001 ***	0.238	0.002 **	0.018*	0.147	0.041*
	FCH ₄ (mmol m ⁻² d ⁻¹)	logarithmic	0.019*	0.001***	0.092	0.001***	0.002**	0.124
	FN ₂ O (mmol m ⁻² d ⁻¹)	logarithmic (value+median)	0.262	0.954	0.028*	0.279	0.047*	0.881
Figure or table	Variable	Transformation	p-values of the Betadisper test					
			Fall vs Winter	Fall vs Spring	Fall vs Summer	Winter vs Spring	Winter vs Summer	Spring vs Summer
Tab. S2	Temperature (°C)	logarithmic	0.689	0.515	0.093	0.987	0.121	0.093
	%O ₂ (%)	logarithmic	0.005 **	0.381	0.127	0.068	0.134	0.503
	SRP (μmol L ⁻¹)	square root	0.102	0.858	0.002 **	0.105	0.565	0.071
	P _{tot} (mg L ⁻¹)	square root	0.908	0.879	0.001 ***	0.779	0.103	0.002 **
	NH ₄ ⁺ (μmol L ⁻¹)	square root	0.143	0.138	0.629	0.942	0.439	0.397
	NO ₂ ⁻ (μmol L ⁻¹)	square root	0.801	0.492	0.392	0.584	0.078	0.057
	NO ₃ ⁻ (μmol L ⁻¹)	square root	0.412	0.665	0.852	0.614	0.364	0.596
	DIN (μmol L ⁻¹)	square root	0.407	0.722	0.912	0.624	0.523	0.829
	Chl-a (μg L ⁻¹)	square root	0.721	0.340	0.001 ***	0.194	0.002 **	0.002 **
TSM (mg L ⁻¹)	square root	0.933	0.814	0.159	0.799	0.006 **	0.004 **	
Fig. 3	CH ₄ (nmol L ⁻¹)	logarithmic	0.934	0.842	0.066	0.919	0.083	0.045
	%N ₂ O (%)	logarithmic	0.445	0.861	0.022*	0.326	0.026*	0.031*
	pCO ₂ (ppm)	square root	0.119	0.197	0.122	0.049 **	0.059	0.417
Fig. S5	FCO ₂ (mmol m ⁻² d ⁻¹)	logarithmic (value+median)	0.903	0.064	0.031*	0.019*	0.017*	0.969
	FCH ₄ (mmol m ⁻² d ⁻¹)	logarithmic	0.928	0.508	0.084	0.566	0.075	0.095
	FN ₂ O (mmol m ⁻² d ⁻¹)	logarithmic (value+median)	0.273	0.941	0.046*	0.174	0.036*	0.027*

Table S6: Outcomes of the Betadisper and Permanova statistical tests for variables shown in box plots from the listed Figures. Comparisons marked with a square (■) are not shown in boxplots but were conducted using quantile regressions. These additional comparisons were performed to assess a statistical value of the dispersion, which is not provided in quantile regressions. Transformations have been applied to the data to ensure a normal distribution, and are indicated in the Transformation column.

	Variable	Transformation	Betadisper	Permanova
Low (<40%) vs high (>40%) macrophyte cover				
Fig. 6	pCO ₂ (ppm)	square root	0.641	0.136
	CH ₄ (nmol L ⁻¹)	logarithmic	0.989	0.005**
	%N ₂ O (%)	logarithmic	0.524	0.361
Fig. S3	Chl-a (µg L ⁻¹)	logarithmic	0.294	0.003**
	P _{tot} (mg L ⁻¹)	none	0.833	0.975
	SRP (µmol L ⁻¹)	none	0.117	0.127
	Residence time (days)	none	0.263	0.144
	Distance (km)	logarithmic	0.63	0.048*
	Surface (ha)	none	0.669	0.041*
	DIN (µmol L ⁻¹)	square root	0.1472	0.028*
Natural (Holgerson and Raymond, 2016) vs Brussels ponds (This Study)				
Fig. 9	pCO ₂ (µmol L ⁻¹)	logarithmic	0.125	0.014*
	CH ₄ (nmol L ⁻¹)	logarithmic	0.973	0.001***
Center vs Periphery (■)				
None	%N ₂ O (%)	logarithmic	0.02*	0.003**
	DIN (µmol L ⁻¹)	square root	0.132	0.041*
	NH ₄ ⁺ (µmol L ⁻¹)	square root	0.481	0.265
	NO ₂ ⁻ (µmol L ⁻¹)	square root	0.125	0.009**
	NO ₃ ⁻ (µmol L ⁻¹)	square root	0.104	0.007**
	pCO ₂ (ppm)	square root	0.073	0.328
	CH ₄ (nmol L ⁻¹)	logarithmic	0.573	0.009**
Small surfaces (<1ha) vs High Surfaces (>1ha) (■)				
None	%N ₂ O (%)	logarithmic	0.03*	0.012*
	DIN (µmol L ⁻¹)	square root	0.0953	0.043*
	NH ₄ ⁺ (µmol L ⁻¹)	square root	0.11	0.567
	NO ₂ ⁻ (µmol L ⁻¹)	square root	0.103	0.037*
	NO ₃ ⁻ (µmol L ⁻¹)	square root	0.107	0.047*
	pCO ₂ (ppm)	square root	0.128	0.018*
	CH ₄ (nmol L ⁻¹)	logarithmic	0.873	0.011*

Table S7: Equation of quantiles regressions in listed figures and p-value (Wald test).

Figure	Database	Variable 1 (response)	Variable 2 (explicative)	Equation of quantile regression	p-value (Wald test)	Statistical significance		
Fig. 4	All	pCO ₂ (ppm)	%O ₂ (%)	$\sqrt{pCO_2} = 198.06 - 76.83 \times \log(\%O_2)$	0.0000	****		
			Temperature (°C)	$\sqrt{pCO_2} = 66.2 - 9.10 \times \log(T)$	0.5400	ns		
			Chl-a (µg L ⁻¹)	$\sqrt{pCO_2} = 64.14 - 2.18 \times \sqrt{Chla}$	0.1095	ns		
			DIN (µmol L ⁻¹)	$\sqrt{pCO_2} = 45.42 + 1.40 \times \sqrt{DIN}$	0.1624	ns		
			SRP (µmol L ⁻¹)	$\sqrt{pCO_2} = 24.37 + 22.60 \times \sqrt{SRP}$	0.0004	***		
		P _{tot} (mg L ⁻¹)	$\sqrt{pCO_2} = 47.86 + 12.21 \times \sqrt{P_{tot}}$	0.4448	ns			
		CH ₄ (nmol L ⁻¹)	%O ₂ (%)	$\log(CH_4) = 4.60 - 0.80 \times \log(\%O_2)$	0.0124	*		
			Temperature (°C)	$\log(CH_4) = 2.22 + 0.91 \times \log(T)$	0.0000	****		
			Chl-a (µg L ⁻¹)	$\log(CH_4) = 3.26 - 0.04 \times \sqrt{Chla}$	0.0296	*		
			DIN (µmol L ⁻¹)	$\log(CH_4) = 3.52 - 0.04 \times \sqrt{DIN}$	0.0069	**		
			SRP (µmol L ⁻¹)	$\log(CH_4) = 3.06 + 0.02 \times \sqrt{SRP}$	0.8903	ns		
		P _{tot} (mg L ⁻¹)	$\log(CH_4) = 2.67 + 0.63 \times \sqrt{P_{tot}}$	0.0000	****			
		Fig. 5		%N ₂ O (%)	%O ₂ (%)	$\log(\%N_2O) = 2.78 - 0.33 \times \log(\%O_2)$	0.0488	*
					Temperature (°C)	$\log(\%N_2O) = 2.36 - 0.21 \times \log(T)$	0.0356	*
					Chl-a (µg L ⁻¹)	$\log(\%N_2O) = 2.13 + 0.01 \times \sqrt{Chla}$	0.6087	ns
DIN (µmol L ⁻¹)	$\log(\%N_2O) = 1.88 + 0.04 \times \sqrt{DIN}$				0.0000	****		
SRP (µmol L ⁻¹)	$\log(\%N_2O) = 1.96 + 0.15 \times \sqrt{SRP}$				0.0956	ns		
P _{tot} (mg L ⁻¹)	$\log(\%N_2O) = 2.21 - 0.07 \times \sqrt{P_{tot}}$			0.2153	ns			
NH ₄ ⁺ (µmol L ⁻¹)	$\log(\%N_2O) = 2.06 + 0.03 \times \sqrt{NH_4^+}$			0.0436	*			
NO ₃ ⁻ (µmol L ⁻¹)	$\log(\%N_2O) = 1.91 + 0.04 \times \sqrt{NO_3^-}$			0.0000	****			
NO ₂ ⁻ (µmol L ⁻¹)	$\log(\%N_2O) = 1.84 + 0.36 \times \sqrt{NO_2^-}$			0.0000	****			
O ₂ :DIN (mol:mol)	$\log(\%N_2O) = 2.41 - 0.35 \times \log(O_2:DIN)$			0.0001	****			
O ₂ :NH ₄ ⁺ (mol:mol)	$\log(\%N_2O) = 2.27 - 0.07 \times \log(O_2:NH_4^+)$	0.1007	ns					
O ₂ :NO ₃ ⁻ (mol:mol)	$\log(\%N_2O) = 2.46 - 0.31 \times \log(O_2:NO_3^-)$	0.0001	****					
O ₂ :NO ₂ ⁻ (mol:mol)	$\log(\%N_2O) = 2.94 - 0.32 \times \log(O_2:NO_2^-)$	0.0001	****					
Fig. 6	Average for each pond	pCO ₂ (ppm)	Surface (ha)	$\sqrt{pCO_2} = 68.64 - 6.74 \times Surface$	0.0020	**		
			Distance (km)	$\sqrt{pCO_2} = 57.64 - 0.23 \times Distance$	0.9203	ns		
		CH ₄ (nmol L ⁻¹)	Surface (ha)	$\log(CH_4) = 3.18 + 0.13 \times Surface$	0.0168	*		
			Distance (km)	$\log(CH_4) = 2.92 + 0.08 \times Distance$	0.0036	**		
		%N ₂ O (%)	Surface (ha)	$\log(\%N_2O) = 2.36 - 0.12 \times Surface$	0.0479	*		
Distance (km)	$\log(\%N_2O) = 2.73 - 0.09 \times Distance$	0.0081	**					
Fig. S2	Fall	CH ₄ (nmol L ⁻¹)	P _{tot} (mg L ⁻¹)	$\log(CH_4) = 1.99 + 1.35 \times \sqrt{P_{tot}}$	0.0273	*		
			Winter	$\log(CH_4) = 2.87 - 0.09 \times \sqrt{P_{tot}}$	0.8880	ns		
			Spring	$\log(CH_4) = 2.29 + 0.46 \times \sqrt{P_{tot}}$	0.0123	*		
			Summer	$\log(CH_4) = 2.55 + 0.38 \times \sqrt{P_{tot}}$	0.0038	**		
			Fall	$\log(CH_4) = 3.07 + -0.01 \times \sqrt{DIN}$	0.8728	ns		
	Winter	CH ₄ (nmol L ⁻¹)	Spring	$\log(CH_4) = 2.92 - 0.01 \times \sqrt{DIN}$	0.6101	ns		
			Summer	$\log(CH_4) = 3.86 - 0.07 \times \sqrt{DIN}$	0.0080	**		
			DIN (µmol L ⁻¹)	$\log(CH_4) = 3.43 + 0.01 \times \sqrt{DIN}$	0.9405	ns		
			Fall	$\log(\%N_2O) = 1.78 + 0.05 \times \sqrt{DIN}$	0.0005	***		
			Winter	$\log(\%N_2O) = 1.90 + 0.03 \times \sqrt{DIN}$	0.0000	****		
Spring	%N ₂ O (%)	Summer	$\log(\%N_2O) = 1.86 + 0.06 \times \sqrt{DIN}$	0.0071	**			
		Fall	$\log(\%N_2O) = 1.77 + 0.08 \times \sqrt{DIN}$	0.0317	*			
		Summer	DIN (µmol L ⁻¹)	$\sqrt{DIN} = 13.25 - 0.98 \times Distance$	0.0294	*		
			NH ₄ ⁺ (µmol L ⁻¹)	$\sqrt{NH_4^+} = 2.27 + 0.33 \times Distance$	0.3210	ns		
			NO ₃ ⁻ (µmol L ⁻¹)	$\sqrt{NO_3^-} = 15.22 - 1.45 \times Distance$	0.0015	**		
NO ₂ ⁻ (µmol L ⁻¹)	$\sqrt{NO_2^-} = 2.01 - 0.16 \times Distance$		0.0311	*				
atm NO ₂ ⁻ (µg m ⁻³)	$atm NO_2^- = 37.56 - 3.35 \times Distance$		0.0003	***				
%N ₂ O (%)	atm NO ₂ ⁻ (µg m ⁻³)	$\log(\%N_2O) = 1.66 + 0.04 \times atm NO_2^-$	0.0043	**				
Fig. S4		CHl-a (µg L ⁻¹)	Distance (km)	$\sqrt{Chla} = 1.61 - 0.05 \times Distance$	0.3410	ns		