



1 **Carbon dioxide and methane fluxes from different surface types**
2 **in a created urban wetland**

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18



19 **Abstract.** Many wetlands have been drained due to urbanization, agriculture, forestry or other purposes, which has
20 resulted in losing their ecosystem services. To protect receiving waters and to achieve services such as flood control and
21 stormwater quality mitigation, new wetlands are created in urbanized areas. However, our knowledge of greenhouse gas
22 exchange in newly created wetlands in urban areas is currently limited. In this paper we present measurements carried
23 out at a created urban wetland in boreal climate.

24 We conducted measurements of ecosystem CO₂ flux (NEE) and CH₄ flux (F_{CH₄}) at the constructed stormwater wetland
25 Gateway in Nummela, Vihti, Southern Finland using eddy covariance (EC) technique. The measurements were
26 commenced the fourth year after construction and lasted for one full year and two subsequent growing seasons. Besides
27 ecosystem scale fluxes measured by EC tower, the diffusive CO₂ and CH₄ fluxes from the open-water area (F_{w_CO₂} and
28 F_{w_CH₄}, respectively) were modelled based on measurements of CO₂ and CH₄ concentration in the water. Fluxes from
29 vegetated area were estimated by applying a simple mixing model using above-mentioned fluxes and footprint-weighted
30 fractional area. The half-hourly footprint-weighted contribution of diffusive fluxes from open water ranged from 0 to 25.5
31 % in year 2013.

32 The annual NEE of the studied wetland was 8.0 g C-CO₂ m⁻² yr⁻¹ with the 95 % confidence interval between -18.9 and
33 34.9 g C-CO₂ m⁻² yr⁻¹ and F_{CH₄} was 3.9 g C-CH₄ m⁻² yr⁻¹ with the 95 % confidence interval between 3.75 and 4.07 g C-
34 CH₄ m⁻² yr⁻¹. The ecosystem sequestered CO₂ during summer months (June-August), while the rest of the year it was a
35 CO₂ source. CH₄ displayed strong seasonal dynamics, higher in summer and lower in winter, with a sporadic emission
36 episode in the end of May 2013. Both CH₄ and CO₂ fluxes, especially those obtained from vegetated area, exhibited strong
37 diurnal cycle during summer with synchronized peaks around noon. The annual F_{w_CO₂} was 297.5 g C-CO₂ m⁻² yr⁻¹ and
38 F_{w_CH₄} was 1.73 g C-CH₄ m⁻² yr⁻¹. The peak diffusive CH₄ flux was 137.6 nmol C-CH₄ m⁻² s⁻¹, which was synchronized
39 with the F_{CH₄}.

40 Overall, during the monitored time period, the established stormwater wetland had a climate warming effect with 0.263
41 kg CO₂-eq m⁻² yr⁻¹ of which 89 % was contributed by CH₄. The radiative forcing of the open-water exceeded the vegetation
42 area (1.194 kg CO₂-eq m⁻² yr⁻¹ and 0.111 kg CO₂-eq m⁻² yr⁻¹, respectively), which implies that, when considering solely
43 the climate impact of a created wetland over a 100-year horizon, it would be more beneficial to design and establish
44 wetlands with large patches of emergent vegetation, and to limit the areas of open-water to the minimum necessitated by
45 other desired ecosystem services.

46 1 Introduction

47 Wetlands provide many beneficial ecosystem services such as flood control and water quality mitigation, natural habitat
48 for flora and fauna and recreational opportunities (Mitsch and Gosselink, 2015). Many wetlands have been drained
49 globally for agriculture, forestry and other purposes including urbanization at the cost of losing wetland ecosystem
50 services (Vasander et al., 2003). Migration from rural area to cities will increase in even greater number in the near future,
51 and UN 2016 report has predicted that 75 % of the world population will be living in cities by 2030. There is an urgent
52 need for more sustainable urbanism and one effective measure is to create functional and connected wetland networks in
53 cities (Lucas et al., 2015; Mungasavalli and Viraraghavan, 2006).

54 Wetlands can take up carbon dioxide (CO₂) through emergent and submerged vegetation but they are also important
55 sources of methane (CH₄), a greenhouse gas more potent than CO₂ when considered over a 100-year horizon (Stocker et
56 al., 2014). The exchange of greenhouse gases (GHG) such as CO₂ and CH₄ between atmosphere and ecosystem have



57 direct influence on the atmospheric concentration of these gases, thus besides the ecosystem services that wetland provide,
58 the GHG budget of constructed wetlands should be accounted for according to international agreements such as the Kyoto
59 protocol.

60 Reports on boreal wetlands, such as peatlands, have shown that large carbon storage remains in the soil due to anaerobic
61 conditions limiting microbial decomposition, and thus offering a global cooling effect (Frolking et al., 2006). However,
62 in newly constructed urban wetlands on mineral soil the gas exchange may be very different from natural wetlands: 1)
63 the cooling effect of a wetland may be reduced or it becomes a source of carbon due to the early successional stage of the
64 wetland, 2) wetlands in close proximity to urban centers receive significant amount of nutrients and dissolved organic
65 carbon from runoff and 3) urban wetlands exhibit high spatial heterogeneity and hydrology where different processes of
66 the production and transportation of GHG are involved. At the areas with emergent vegetation, CO₂ is absorbed by
67 photosynthetic activity during daytime and growing season and is released through respirational processes. At open-water
68 surfaces, the net production of CO₂ is a result of photosynthesis by algae, cyanobacteria as well as submerged aquatic
69 plants, respiration of organic carbon and oxidation of CH₄ produced in the water. When the CO₂ concentration in the
70 water exceeds atmospheric equilibrium, the surface becomes a source of CO₂. CH₄ can be produced through anaerobic
71 metabolism in wetland soil and can be transported to the atmosphere by plant-mediated pathway through aerenchyma,
72 sediment ebullition and diffusive fluxes at water-atmosphere interface. In open water, the transport is dominated by
73 diffusion whereas in vegetated area the plant-mediated transport is most prominent.

74 Urban wetlands have received extensive attention globally and their societal and economical importance have been
75 evaluated (Salminen et al., 2013), whereas their climate impact is still largely overlooked except for only a few studies
76 (e.g. Morin et al., 2014a; Morin et al., 2014b). The only review of GHG emission in constructed wetlands for wastewater
77 treatment reported that the average CO₂ emission was 92.3 mg CO₂-C m⁻² h⁻¹ and that the CH₄ emission ranged from 1.6
78 to 27 mg CH₄-C m⁻² h⁻¹ from free water surface (Mander et al., 2014). All of the studies were based on static chamber
79 measurements during a short period so that the annual carbon balance of the ecosystem could not be assessed. In contrast
80 to static chamber measurements, eddy covariance (EC) method provides continuous measurements of GHG exchange at
81 ecosystem scale, presenting the net result of fluxes as exchange in different source area contributing simultaneously within
82 the footprint extent (Baldocchi, 2003). It is worth noticing that one of the assumptions of the EC method is surface
83 homogeneity, yet in many study sites the situation are far from ideal. The change of source area due to changes in wind
84 provides difficulties in estimating GHG emissions in spatially heterogeneous sites especially in short-term flux
85 measurements (Baldocchi et al., 2012). Therefore, for heterogeneous sites such as urban wetlands, accurate footprint
86 modelling and surface area map at high spatial resolution are important in identifying the source area, and a land-surface
87 specific analysis is vital to reveal the diel pattern, sink/source strength of the wetland.

88 The objective of this study is to investigate how CO₂ and CH₄ surface-atmosphere exchange vary with seasonality and
89 spatial heterogeneity and what the annual radiative forcing of these gases are in a constructed urban wetland near town
90 Nummela, Municipality of Vihti, Southern Finland. The studied Gateway wetland was designed and implemented to serve
91 the purposes of stormwater quality treatment, creating an urban park, as well as supporting biodiversity. Besides taking
92 advantage of ecosystem-scale EC measurements, we also parse the variability of gas exchange induced by surface
93 heterogeneity (open water and vegetated area) using diffusional flux modeling and footprint modelling overlapped on a
94 high-resolution surface map. To illustrate how the urban wetland functions as a source or a sink of GHG equivalents, we



95 calculate separately the sustained global warming potential (SGWP) of CO₂ and CH₄ over a hundred-year horizon in each
96 surface type.

97 **2 Materials and Methods**

98 **2.1 Site description**

99 Our study site is a created stormwater wetland Gateway, located by an eutrophicated Lake Enäjärvi in the District of
100 Nummela, Municipality of Vihti, Southern Finland (60.3272°N, 24.3369°E). Southern Finland experiences a climate with
101 a 30-year mean air temperature of 4.6 °C and an annual precipitation rate of 627 mm in the period of year 1981-2010
102 (Pirinen et al., 2012).

103 The wetland was constructed in 2010 at the mouth of a 550 hectare largely urbanized (35 % impervious) watershed of
104 Stream Kilsoi. It was excavated over six weeks in early winter 2010 on an abandoned agricultural field growing meadow
105 vegetation. All of the old drainage ditches were blocked as amphibian habitats, which also ensured only one inlet route
106 receiving water from Stream Kilsoi and one outlet route discharging water to the nearby Lake Enäjärvi. Lake Enäjärvi is
107 a eutrophicated lake. The internal phosphorus load from human activities and the run-off from its catchments have resulted
108 in regular cyanobacterial blooms and fish kills in the lake.

109 The wetland park has a total area of 7 hectares within which - during mean water flow conditions - a 0.5 hectare inundated
110 wetland is located. This stormwater treatment wetland consists of an inlet stilling pond, a meandering shallow water area
111 with three habitat islands, and an outlet pond. The average water depth in the ponds is 1.5 m; within emergent vegetation
112 patches water depth ranges between 0.3 and 0.5 m. There are also submerged macrophytes in the open water as the water
113 is shallow, thus in the paper we refer the “vegetated area” to the area with emergent vegetation and “open water” to the
114 area covered by water in the absence of emergent macrophytes. The outlet bottom dam sets low water level (WL) to 50.04
115 m above the Baltic Sea level (N60+ coordinate system). Herbaceous vegetation has been allowed to fully self-establish
116 after the construction of the wetland. Annual monitoring of vegetation carried out in summers 2010, 2011 and 2012
117 indicated rapid self-establishment of vegetation which was rich in taxa and dominated by native species (Wahlroos et al.,
118 2015). At frequently-inundated area (elevation levels of 50-50.35 m), vegetation was arranged in dense patches with
119 different dominating wetland plant species: *Typha latifolia* L., *Iris pseudacorus* L., *Carex* spp. or *Juncus effuses* L. At the
120 major less-frequently inundated area (elevation levels of 50.35-50.45m), the wet meadow species *Filipendula ulmaria* L.
121 (Maxim.), *Lysimachia vulgaris* L., and *Lythrum salicaria* L. with the three species co-existing at 1:1:1 ratio formed the
122 plant community. Drier areas (elevation levels of 50.45-50.60 m) were mostly colonized by dry meadow species such as
123 *Poa* spp. and *Calamagrostis* spp., including patches dominated by *Cirsium* species (Fig. S1).

124 **2.2 Water and micrometeorological measurements**

125 Water monitoring stations were set up at the inlet (60.3283° N, 24.3356° E) and at the outlet (60.3281° N, 24.3377° E) of
126 the wetland. During the 2012-2013 and 2013-2014 monitoring periods, water temperature as well as water turbidity,
127 oxygen concentration, conductivity and pH were measured at the inlet and outlet monitoring station with the YSI-6000
128 series multiparameter sonde (YSI Inc., Yellow Springs, OH, USA). Measurements were conducted continuously with 10-
129 minute interval. Water level at the outlet was measured continuously with a pressure gauge (STS sensor, Sensor Technik
130 Sirmach AG, Switzerland). At the outlet monitoring station, the concentration of dissolved carbon dioxide ([CO₂]) and
131 dissolved methane ([CH₄]) were measured with Contros HydroC™ CO₂ and HydroC™ CH₄ sensors (CONTROS Systems



132 & Solutions GmbH, Germany). In 2014, the same sensors were also installed at the inlet monitoring station to measure
133 [CO₂] and [CH₄]. Dissolved CO₂ and CH₄ molecules diffuse from water column into the detection chamber through a
134 thin-film composite membrane where the concentration of CO₂ and CH₄ is determined by means of IR absorption
135 spectrometry and Tunable Diode Laser Absorption Spectroscopy, respectively.

136 Local weather conditions were recorded with a Vaisala WXT weather transmitter (WXT520, Vaisala Oyj, Finland) at the
137 inlet monitoring station. Rainfall, wind speed and direction, temperature and relative humidity were recorded
138 continuously at 10-minute interval. Photosynthetic photon flux density (PPFD) was measured with a PQS1 PAR quantum
139 sensor (Kipp & Zonen, the Netherland). Due to instrument failure we obtained PPFD data only from 26 Jan to 7 April
140 and from 22 July to 29 Dec 2013. The gaps were filled with PPFD data from another meteorological station nearby (60°38'
141 N, 23°58' E) in Lettosuo, Finland. The prevailing wind directions were southwest and northeast, and the average of half-
142 hourly average wind speed was 1.13 m s⁻¹ from January to December 2013 with higher wind speed in winter than in
143 summer. The average daily air temperature was 5.9 °C with the minimum and maximum daily temperatures of -24.4 °C
144 and 23.3 °C in year 2013. During the winter 2012-2013, there was ice coverage from the beginning of December 2012 to
145 the end of March 2013. In contrast, winter was mild and warm in 2014 and there was practically no snow cover during a
146 winter period (December 2013-March 2014).

147 **2.3 Greenhouse gas measurements by EC tower and gap-filling**

148 To understand the whole-ecosystem exchange of CO₂ and CH₄ in the wetland, a 2.9 m eddy covariance tower was
149 established in the autumn of 2012 on the southern side of the wetland. The operational period of the EC tower was the
150 entire calendar year of 2013 (from 1 January to 31 December 2013) and the peak growing season in 2014 (from 1 June to
151 31 August 2014). The EC set-up included a 3D-sonic anemometer (uSonic-3, Metek, Elmshorn, Germany) to measure
152 the three wind speed components and sonic temperature, a gas analyser (LI-7200, Li-Cor Inc., Lincoln, Nebraska, USA)
153 which measures CO₂ and H₂O mixing ratio and a TDL gas analyser (TGA100A, Campbell Scientific Inc., USA) to
154 measure CH₄ mixing ratio. Data from the analyzers were collected on a computer at the frequency of 10 Hz. The post-
155 processing of the EC flux data has been done with EddyUH post-processing software (Mammarella et al., 2016). The
156 fluxes were calculated as 30-min covariances between the vertical wind velocity and the gas mixing ratio using block
157 averaging. The raw data was despiked according to standard methods (Vickers and Mahrt, 1997). Coordinate rotations
158 were conducted by performing a two-step rotation to make the x-axis along the mean wind direction and the mean vertical
159 wind velocity zero within each 30-min block. The time lag between the anemometer and gas analyzer signals, resulting
160 from the transport through the inlet tube, were determined for each 30-min interval by maximizing the cross-correlation
161 function between vertical wind speed and the scalar (CO₂ and CH₄). The fluxes were corrected for high-frequency loss
162 due to the limited frequency response of the EC system and low-frequency loss due to the limited averaging time period
163 used for calculating the fluxes. Theoretically and experimentally determined co-spectral transfer functions at low and
164 high frequency were used in the correction (Mammarella et al., 2009).

165 After calculating the fluxes, data collected from periods when sonic anemometer showed sign of freezing (mean
166 temperature < 0.5 °C and standard deviation of temperature > 1.5 °C) were discarded. The data collected during weak
167 turbulence with friction velocity below 0.1 m s⁻¹ have been removed. The measurement points with flux stationarity
168 greater than 1 were omitted to ensure the quality of the co-variances. Fluxes were further filtered according to the wind
169 direction. Since the patchy forest to the southeast of the EC tower (from 100° to 200°) and the highway to the west (from
170 200° to 280°) could potentially lead to flow distortion and additional source of CO₂ and CH₄, only fluxes from 280° to



171 100° were accepted for further analysis. The percentage of 30-min fluxes excluded from this analysis was 72 % for CO₂
172 and 73 % for CH₄ in 2013, whereas in 2014 the percentage for data exclusion was 54 % for CO₂ and 68 % for CH₄.

173 We used an artificial neural network (ANN) technique to gap-fill half-hourly flux data using meteorological variables
174 (Moffat et al., 2007; Papale et al., 2006). Those variables included radiation, air temperature, water temperature, water
175 level, wind speed, relative humidity, time of the day, season, and dissolved CO₂ and CH₄ concentration in the water. We
176 tested the model performance with different ANN architectures, starting from the architecture with the most complexity,
177 then reduced the variables to find the simplest ANN architecture with good performance (more than 5 % loss in model
178 accuracy with additional variable reduction). For CO₂, water level and wind speed were found to have trivial contribution
179 to the ANN model thus they were removed from the model input, while for CH₄, only wind speed was removed for the
180 same reason. We found that dissolved gas concentration greatly improved the model prediction as they captured the
181 variation of diffusive fluxes from the water (Fig. S2). Ancillary meteorological variables in general had good data
182 coverage and short gaps (up to several hours) were gap-filled by linear interpolation. The only exception was dissolved
183 gas concentration, which had long measurement breakage in year 2013 (day of year 214-254). Fluxes were therefore gap-
184 filled with two separate ANNs, one with dissolved gas concentration and one without. During the above mentioned period
185 with long gaps, the ANN modeled without dissolved gas concentration were used to gap-fill.

186 Levenberg-Marquardt algorithm was used in the learning process of ANN. The optimized number of neurons in the hidden
187 layer were determined by training the network 100 times with varying number of neurons (from 3 to 15), and 10 neurons
188 was considered to be sufficient after evaluating the performance of the network using root-mean-square-error (RMSE)
189 (data not shown). The entire dataset was divided into three parts, 2/3 of the data was used to train the networks, 1/6 for
190 testing the networks and the remaining 1/6 was used for validating the networks. Since the training of the networks can
191 be biased towards periods with greater data coverage (e.g. daytime conditions), the environmental variables were first
192 divided into five natural clusters using a k-mean clustering algorithm in Matlab (MATLAB 2015a, The MathWorks, Inc.,
193 Natick, Massachusetts, United States), and then the data used for training, testing and validation was proportionally
194 extracted from each cluster. After each data extraction, the network was reinitialized for 10 times to avoid local minima
195 and the initialization with the lowest RMSE was selected and resulting network was saved. We repeated the whole process
196 of data extraction and initialization for 20 times, and we used the median of these 20 predictions to gap-fill the missing
197 flux values. The uncertainty of the ANN gap-filling procedure was presented using a 95 % confidence interval of the 20
198 ANN predictions.

199 The gap-filled net ecosystem exchange (NEE) can be further partitioned into two components gross ecosystem production
200 (GEP) and ecosystem respiration (R_{eco}) according to the following equation:

$$201 \quad NEE = GEP + R_{eco}, \quad (1)$$

202 where positive R_{eco} represents a net carbon flux from the ecosystem to the atmosphere and negative GEP represents a net
203 carbon input from the atmosphere to the ecosystem. Thus the negative NEE indicates that the ecosystem is a carbon sink
204 and the positive NEE means the ecosystem is a carbon source. R_{eco} was estimated using a model describing the temperature
205 dependence of R_{eco}

$$206 \quad R_{eco} = R_0 e^{[E \left(\frac{1}{T_0} - \frac{1}{T_{air} + T_1} \right)]} \quad (2)$$

207 where $E = 346.37$ K is an activation-energy-related physiological parameter, T_{air} is the air temperature, $T_0 = 56.02$ K and
208 $T_1 = 227.13$ K (Lloyd and Taylor, 1994; Aurela et al., 2009). R_0 is the rate of ecosystem respiration at 10 °C. We first fitted



209 the model with nighttime NEE (which represents the nighttime ecosystem respiration since photosynthesis is assumed to
210 be zero at night) and determined E . We then calculated R_0 for each of the bi-weekly periods (Aurela et al., 2009). This
211 model was then extrapolated to daytime periods so that R_{eco} in the daytime was obtained. GEP was estimated as the
212 difference between NEE and R_{eco} .

213 2.4 Diffusive gas exchange

214 We calculated diffusive gas exchange F from open water according to the boundary layer model

$$215 F = k (c_{aq} - c_{eq}), \quad (3)$$

216 where k is the gas transfer velocity (cm h^{-1}), c_{aq} is the gas concentration in surface water (mol m^{-3}) and c_{eq} is the gas
217 concentration that surface water would have when it reaches equilibrium with the air (mol m^{-3}). c_{aq} and c_{eq} can be obtained
218 according to the solubility of the gas

$$219 c_{aq} = 10^{-3} k_H p \chi_{water} \quad (4)$$

$$220 c_{eq} = 10^{-3} k_H p \chi_{air} \quad (5)$$

221 where k_H is Henry's law constant for the respective gas ($\text{mol L}^{-1} \text{atm}^{-1}$), p is air pressure (atm), χ_{water} is the gas mixing ratio
222 in surface water (ppm) and χ_{air} is the gas mixing ratio in the air (ppm). In this study, χ_{water} was obtained from outlet
223 monitoring station as it was located most time in the flux footprint area and it had longer data coverage than from inlet
224 monitoring station. The gas transfer velocity k can be calculated as the formula below (Cole and Caraco, 1998):

$$225 k = (2.07 + 0.215 U_{10}^{1.7}) \left(\frac{S_c}{600}\right)^{-0.5}, \quad (6)$$

226 where U_{10} is the horizontal wind speed extrapolated to 10 m using the theoretical log wind profile equation (m s^{-1} ,
227 approximately $U_{10} = 1.15 U$ where U is the measured wind speed at 2.9 m height in the study site) and S_c is the
228 temperature-dependent Schmidt number of the respective gas. When gas concentration measurement was not available,
229 linear interpolation was applied to obtain monthly and annual diffusive GHG fluxes from the open water.

230 Although the above-mentioned Cole-Caraco (CC) method is the most simple and most often used model for gas transfer
231 velocity, the limitation of CC method is that it considers wind as the sole factor to cause the water turbulence and to drive
232 the gas exchange. More complicated models were suggested to include the effect of buoyancy flux driven turbulence
233 (Heiskanen et al., 2014; Tedford et al., 2014). It is important to note that we should apply with caution the model
234 parameterization concluded from other sites with different meteorological and environmental condition. In the present
235 study, the open water is connected shallow open-water pools with a maximum depth of 2 m while other studies are for
236 deeper waters. Meanwhile, recent study showed good agreement between the diffusive fluxes calculated using CC
237 method and measurements based on floating chamber (Cole et al., 2010).

238 2.5 Estimating zone fluxes and radiative forcing

239 By combining EC tower and diffusive flux from the open-water, the following model can be derived

$$240 F_{EC} = F_{water} \times f_{water} + F_{water} \times f_{water} \quad (7)$$



241 where F_{EC} is the flux measured by EC tower, F_{water} and F_{veg} stands for the flux from open-water and vegetated area,
242 respectively, f_{water} and f_{veg} are the footprint-weighted spatial fraction of open-water and vegetated area. In this study,
243 ebullition was neither measured nor calculated, so the flux from water was only represented by the diffusive flux.

244 Specifically, we first modelled the half-hourly flux footprint with a parameterization of a three-dimensional backward
245 Lagrangian footprint model (Kljun et al., 2015) in Matlab (MATLAB 2015a, The MathWorks, Inc., Natick, Massachusetts,
246 United States). Periods in which the wind came from the patchy forest to the southeast of the EC tower (between 100°
247 and 200°) and the highway to the west (between 200° and 280°) were eliminated in the footprint analysis. Secondly, a
248 land cover classification map of vegetated and open-water zones was delineated manually using a high-resolution aerial
249 image acquired from National Land Survey of Finland during the growing season of 2013 (open source:
250 <https://www.maanmittauslaitos.fi/>) with an image manipulation software (Gimp 2.10.6). Thirdly, the flux footprints were
251 aligned and combined with the land cover classification map to calculate half-hourly f_{veg} and f_{water} within 90 % footprint
252 contour lines. Specifically, we assigned each footprint pixel within the 90 % footprint area to either open-water or
253 vegetated area on the land cover classification map while the footprint of the pixels outside 90 % footprint area were
254 regarded as zero. f_{water} was calculated as the sum of footprint within open-water area to the total footprints while f_{veg}
255 was calculated as the sum of footprint within vegetated area to the total footprints. Noting that none of the 90 % footprint
256 contour lines exceeded the map area, the sum of f_{veg} and f_{water} equaled to 1. In order to obtain the long-term aggregated
257 footprint of carbon fluxes, we calculated also the monthly and annual aggregated footprint climatology during the study
258 period.

259 To better understand the influence of greenhouse gas fluxes in this urban wetland, we calculated the sustained global
260 warming potential (SGWP) for CO₂ and CH₄ over a hundred-year horizon in each surface type. The difference between
261 SGWP and global warming potential (GWP) is that SGWP accounts for the effect of GHG remains in the atmosphere
262 during the period. Since CH₄ is a more potent greenhouse gas, we multiply the emission of CH₄ by a factor of 45 to
263 convert it to kg CO₂-eq m⁻² yr⁻¹ (Neubauer and Megonigal, 2015).

264 2.6 Statistical analysis

265 The Pearson correlations (r) were determined between fluxes and environmental variables. Differences in the fluxes and
266 environmental variables between the two peak growing seasons (summer 2013 and 2014) were evaluated using the t -test.
267 Cumulative annual GHG fluxes measured by EC tower are reported as the median of the 20 ANN predictions and
268 uncertainty are presented as 95 % confidence interval of the 20 ANN predictions. As diffusive GHG fluxes were
269 calculated from gas concentration meteorological parameters, no standard error is reported for the cumulative annual fluxes
270 from the open water. All statistical analysis were performed in Matlab (MATLAB 2015a, The MathWorks, Inc., Natick,
271 Massachusetts, United States).

272 3 Results

273 3.1 Ecosystem seasonality and environmental variables

274 Daily average PPFD ranged from 0.9 to 691.5 $\mu\text{mol m}^{-2} \text{s}^{-1}$ in year 2013 with the highest value appeared in July. June had
275 the highest monthly average PPFD with 486.1 $\mu\text{mol m}^{-2} \text{s}^{-1}$ followed by July and August with 470.2 and 430.6 $\mu\text{mol m}^{-2}$
276 s^{-1} , respectively. The PPFD during the peak growing season in year 2014 was on average 361.8 $\mu\text{mol m}^{-2} \text{s}^{-1}$, lower than
277 that during the same period in 2013 (Fig. 1a).



278 Mean daily water temperature (T_{water}) ranged from 0 °C in March to 23.7 °C in June with an annual average of 7.9 °C in
279 2013 and from 0 °C in February to 21.4 °C in July in 2014. Mean daily air temperature (T_{air}) had more fluctuation and
280 ranged from -15.6 °C in January to 23.3 °C in June 2013 and from -19.0 °C in January to 23.4 °C in July 2014 (Fig. 1b).
281 The open-water area experienced an ice-covered period between 1 January and 31 March 2013, while the winter 2013-
282 2014 was so mild and warm that there was practically no snow cover during December 2013 – March 2014. Comparing
283 the temperature between the two peak growing seasons, both T_{water} and T_{air} were higher in June 2013 while T_{air} was lower
284 in July 2013 than in 2014. In August, there was no significant temperature difference between the two years. Four seasons
285 were classified for the ecosystem based on the trend in T_{air} and T_{water} . In spring (April and May), the daily temperature
286 started to increase, the vegetation showed a sign of early growing season and the warm temperature unfroze the lake. In
287 summer, the peak growing season (June - August), vegetation exhibited the maximum growth which was reflected in the
288 large negative GEP value, and the temperatures reached the annual maxima. In autumn (September and October), daily
289 temperatures began to drop and the vegetation showed signs of early senescence. In winter (January to March and
290 November, December), temperatures reached the annual minima and vegetation was inactive in carbon sequestration.
291 Precipitation was higher in August 2014 than in the preceding August, almost twice as high as that of 2013.

292 WL was higher in the winter and lower in the summer in 2013. The daily average of WL varied between 50.06 m in July
293 2013 and 50.4 m in April 2013. There was a spring peak in 2013 when the highest WL was observed due to snow melt
294 while in 2014 no such event appeared due to the mild winter 2013-2014 without ice-covered period (Fig. 1c). The average
295 daily WL from January to August was similar (50.13 cm and 50.15 cm for 2013 and 2014, respectively). However, during
296 peak growing season, it was on average 5.7 cm higher in 2014 than in 2013.

297 The annual rainfall in 2013 (snowfall not included) was 363.6 mm which happened mostly during summer and autumn
298 (Fig. 1d). The maximum daily-averaged rainfall was in August (26.7 mm day⁻¹) while monthly-averaged rainfall was
299 highest in November with 73.8 mm month⁻¹ followed by August with 68.3 mm month⁻¹. In 2014, exceptionally high
300 amount of rainfall was observed in August (125.7 mm month⁻¹), while the amount of rainfall in the other months were
301 similar to 2013.

302 The daily-averaged CO₂ concentration in the water ([CO₂]) in 2013 had large variation with the maximum (9324 ppm)
303 and the minimum (353 ppm) both happening in October (Fig. 1e). [CO₂] was higher in summer months (5457 ppm) and
304 lower in winter months (3345 ppm). [CO₂] was higher in 2014 with an average of 4924 ppm from January to August than
305 in 2013 with an average of 3781 ppm. It also exhibited seasonal variation with high concentration in summer (8084 ppm)
306 and low concentration in winter (3513 ppm). The [CO₂] measured in the inflow was generally lower than that in the
307 outflow and they were well correlated ($r=0.84$). [CH₄] in the outflow was on average five times higher in 2014 than in
308 2013. The average annual concentration was 0.81 μmol L⁻¹ in year 2013 and 2.25 μmol L⁻¹ in 2014. There were peak
309 [CH₄] episodes in the outflow in May 2013 with a maximum of 5.43 μmol L⁻¹. During the summer months in 2014 there
310 were even higher outflow [CH₄] peaks with a maximum of 16.83 μmol L⁻¹. The [CH₄] had a mean of 0.42 μmol L⁻¹ in the
311 inflow which was lower than that in the outflow, and there was no prominent [CH₄] peaks observed in the inflow. [CH₄]
312 in the inflow and outflow were weakly correlated ($r = 0.2$) (Fig. 1f).

313 3.2 Flux footprint mapping

314 A footprint distribution was modeled for each half hour when an eddy flux measurement was collected at the EC tower.
315 The open-water area accounted for 10 % to 16 % of the total wetland area within the footprint while the rest was comprised



316 of wetland vegetation. When weighted with footprint distribution, f_{water} ranged from 0 to 25.5 % and f_{veg} from 74.5 % to
317 100 %. The 1st quantile, median and 3rd quantile of f_{water} and f_{veg} were 0.09 %, 14.1 %, 17.9 % and 82.1 %, 85.9 %, 91.3
318 %, respectively.

319 The monthly cumulative footprint was slightly different for CO₂ and for CH₄ due to the different missing flux values.
320 However, the difference on average was so small (7 %) and the footprint of CO₂ was used in further analysis. The flux
321 footprints were shown to be northeast to the EC mast due to the wind direction filtering meaning only half-hourly data
322 with wind directions from the wetland area were considered in the analysis (Fig. S3). The monthly-average of the 90 %
323 footprint area covered a minimum of 0.69 ha to a maximum of 2.28 ha with a mean of 1.3 ha. The mean extent of the 90
324 % flux footprints was 128 m. After applying flux footprint function, the monthly-average of the footprint-weighted spatial
325 fraction of open water showed lower value in summer and higher value in winter ranging from 11.3 % to 21.4 % with a
326 mean of 13.3 % in 2013. In 2014 during the peak growing season, on average 13.8 % of the wetland area was comprised
327 of open water and the mean f_{water} was 10 %.

328 3.3 CO₂ and CH₄ fluxes

329 3.3.1 Ecosystem CO₂ and CH₄ fluxes

330 Ecosystem CO₂ and CH₄ fluxes measured by EC tower showed the ecosystem was nearly CO₂ neutral and it was a small
331 CH₄ source in year 2013.

332 Daily average of NEE was near zero during winter time (January to March, on average 0.37 $\mu\text{mol C-CO}_2 \text{ m}^{-2} \text{ s}^{-1}$), slightly
333 positive in spring and it became negative from the end of May till the end of August indicating the ecosystem was a CO₂
334 sink during this period, with a maximum negative value of -5.14 $\mu\text{mol C-CO}_2 \text{ m}^{-2} \text{ s}^{-1}$ in June. Daily-average NEE was
335 highest in September with a maximum of 3.29 $\mu\text{mol C-CO}_2 \text{ m}^{-2} \text{ s}^{-1}$, possibly due to the suppressed GEP and high R_{eco} . In
336 October, November and December, NEE remained low but still positive (on average 0.77 $\mu\text{mol C-CO}_2 \text{ m}^{-2} \text{ s}^{-1}$),
337 demonstrating the milder winter between 2013 and 2014 (Fig. 2 and Fig. S4). NEE, GEP and R_{eco} exhibited strong
338 seasonality in 2013 NEE was negative during June, July and August meaning the ecosystem was a CO₂ sink while the
339 rest of year it was a CO₂ source. NEE was lowest in June and highest in September. Both GEP and R_{eco} achieved their
340 highest values in July (Fig. S4). The cumulative NEE in 2013 was 8 g C-CO₂ $\text{m}^{-2} \text{ yr}^{-1}$ with the 95% confidence interval
341 between -18.9 and 34.9 g C-CO₂ $\text{m}^{-2} \text{ yr}^{-1}$ (Fig. 2).

342 Daily-averaged CH₄ was low but not negligible from January to April (on average 5.1 nmol C-CH₄ $\text{m}^{-2} \text{ s}^{-1}$), with a sudden
343 rise in the end of May reaching a maximum of 48.9 nmol C-CH₄ $\text{m}^{-2} \text{ s}^{-1}$. During summer months the ecosystem exhibited
344 relatively high CH₄ emission (on average 15.4 nmol C-CH₄ $\text{m}^{-2} \text{ s}^{-1}$), not comparable with the emission episode in May
345 but higher than winter months. In autumn (September and October) the daily-average CH₄ was 8.8 nmol C-CH₄ $\text{m}^{-2} \text{ s}^{-1}$
346 and after that it gradually decreased throughout the rest of the year with an average of 5.5 nmol C-CH₄ $\text{m}^{-2} \text{ s}^{-1}$. The
347 cumulative CH₄ for 2013 was 3.9 g C-CH₄ $\text{m}^{-2} \text{ yr}^{-1}$ with the 95% confidence interval between 3.75 and 4.07 g C-CH₄ m^{-2}
348 yr^{-1} (Fig. 3).

349 Comparing the peak growing season between 2013 and 2014, the 30 mins NEE ranged from -20.0 $\mu\text{mol C-CO}_2 \text{ m}^{-2} \text{ s}^{-1}$ in
350 June to 18.5 $\mu\text{mol C-CO}_2 \text{ m}^{-2} \text{ s}^{-1}$ in September 2013. GEP reached maximum negative value in July 2013 with -30.5 μmol
351 $\text{C-CO}_2 \text{ m}^{-2} \text{ s}^{-1}$ and R_{eco} in June with 13.9 $\mu\text{mol C-CO}_2 \text{ m}^{-2} \text{ s}^{-1}$. During the peak growing season 2014, NEE had lowest
352 value -22.6 $\mu\text{mol C-CO}_2 \text{ m}^{-2} \text{ s}^{-1}$ in June, GEP -28.6 $\mu\text{mol C-CO}_2 \text{ m}^{-2} \text{ s}^{-1}$ and R_{eco} had its maximum in the beginning of
353 August 2014 with 11.3 $\mu\text{mol C-CO}_2 \text{ m}^{-2} \text{ s}^{-1}$. The monthly NEEs of peak growing season were -84.1, -76.1 and -22.2 g C-



354 $\text{CO}_2 \text{ m}^{-2} \text{ month}^{-1}$ in June, July and August 2013, and -97.6, -47.5 and -19.6 $\text{g C-CO}_2 \text{ m}^{-2} \text{ month}^{-1}$ in 2014. In both years,
355 daily-averaged GEP had its maximum negative value in July (-13.4 and -12.8 $\text{g C-CO}_2 \text{ m}^{-2} \text{ d}^{-1}$). Daily-averaged R_{eco} was
356 highest in June 2013 with 12.1 $\text{g C-CO}_2 \text{ m}^{-2} \text{ d}^{-1}$ while in 2014 R_{eco} was low in June and the peak was in the end of July
357 with 10.5 $\text{g C-CO}_2 \text{ m}^{-2} \text{ d}^{-1}$ (Fig. S5a). The average CH_4 emission in June, July and August were 24.4, 10.8 and 11 nmol
358 $\text{m}^{-2} \text{ s}^{-1}$ in 2013, and 15.5, 21.3 and 21.3 $\text{nmol m}^{-2} \text{ s}^{-1}$ in 2014, respectively (Fig. S5b).

359 3.3.2 Diffusive CO_2 and CH_4 fluxes from open-water area

360 Diffusive CO_2 and CH_4 fluxes from the open water were estimated based on wind speed, $[\text{CO}_2]$ and $[\text{CH}_4]$ (See Sect. 2.4).
361 The variation of diffusive fluxes demonstrated a pattern driven by both wind speed in short term and gas concentration
362 dynamics in the water in long term. Diffusive CO_2 fluxes ranged from -0.07 to 4.09 $\mu\text{mol CO}_2 \text{ m}^{-2} \text{ s}^{-1}$ with a mean of 1.04
363 $\mu\text{mol CO}_2 \text{ m}^{-2} \text{ s}^{-1}$ in 2013 indicating CO_2 oversaturation in the water. From June to September the averaged flux (1.27
364 $\mu\text{mol CO}_2 \text{ m}^{-2} \text{ s}^{-1}$) was higher than that of the other months (Fig. 4a), corresponding to the higher $[\text{CO}_2]$ in the water
365 during summer months (Fig. 1d). The monthly-averaged diffusive CO_2 flux during peak growing season in 2014 was
366 2.34, 2.71 and 1.99 $\mu\text{mol CO}_2 \text{ m}^{-2} \text{ s}^{-1}$ for June, July and August, significantly higher than during the same period in 2013
367 due to the high $[\text{CO}_2]$ in the open water (Fig. 1d).

368 The average diffusive CH_4 emissions in 2013 was 4.9 $\text{nmol C-CH}_4 \text{ m}^{-2} \text{ s}^{-1}$, where a peak emission appeared in late May
369 with the highest flux of 137.6 $\text{nmol C-CH}_4 \text{ m}^{-2} \text{ s}^{-1}$. Monthly-averaged CH_4 diffusive fluxes showed an increasing trend
370 towards the end of the year with large variation in May due to the peak concentration episode. This phenomenon was
371 mainly driven by the increasing dissolved CH_4 concentration in the outflow in 2013. The monthly-averaged diffusive CH_4
372 flux during peak growing season in 2014 was 20.9, 18.9 and 13.5 $\text{nmol CH}_4 \text{ m}^{-2} \text{ s}^{-1}$ for June, July and August, respectively
373 and they were significantly higher than the same period in 2013 due to the high $[\text{CH}_4]$ in the open water (Fig. 1e).

374 3.3.3 Diel patterns in CO_2 and CH_4 fluxes

375 Only non-gapfilled data were used for determination of diel patterns in both gas fluxes. CO_2 and CH_4 fluxes from vegetated
376 area (F_{veg}) was calculated for each 30-min interval according to formula (5). As expected, CO_2 flux showed strong diel
377 pattern in summer with CO_2 uptake during daytime and release in the night, which was controlled by photosynthetic
378 activity (Fig. 5a). The summer peak CO_2 uptake reached 11.5 $\mu\text{mol m}^{-2} \text{ s}^{-1}$ for the whole constructed wetland ecosystem
379 and 15.2 $\mu\text{mol m}^{-2} \text{ s}^{-1}$ for the vegetated area. The CO_2 flux from the vegetated area had higher maximum uptake than the
380 EC measurements carried out over the whole constructed wetland. In the winter, the CO_2 fluxes from both tower and
381 vegetation were similar, being on average 0.46 and 0.55 $\mu\text{mol m}^{-2} \text{ s}^{-1}$ respectively (Fig. 5b).

382 CH_4 flux also showed diel patterns in the summer with much larger variability than those from CO_2 flux. CH_4 emission in
383 general was higher in daytime than in nighttime. In the daytime in summer, CH_4 flux from the vegetated area was higher
384 than the flux measured from the tower while there was no difference during the nighttime (Fig. 5c, 5d). The summer peak
385 daytime flux from tower (18.9 $\text{nmol m}^{-2} \text{ s}^{-1}$) and vegetated area (24.7 $\text{nmol m}^{-2} \text{ s}^{-1}$) was 2.4 times and 3.3 times higher
386 than the nighttime flux (7.5 $\text{nmol m}^{-2} \text{ s}^{-1}$), respectively. This can be understood as daytime CH_4 flux is linked with
387 photosynthesis while nighttime CH_4 flux is controlled by other processes like diffusion, ebullition and convection between
388 the soil, water and atmosphere. In winter there was small (on average 4.6 $\text{nmol m}^{-2} \text{ s}^{-1}$) but constantly positive CH_4 flux
389 without obvious diel pattern.

390 3.4 Environmental variables with fluxes



391 Only non-gapfilled flux data were used in the Pearson correlation analysis between environmental variables and flux
392 pairs. Radiation, T_{air} and T_{water} all had high negative correlation coefficient (r) with NEE and high positive r with CH_4
393 flux in 2013, corresponding to the results of ANN model parameter selection. Radiation was best correlated with NEE
394 and T_{water} was best correlated with CH_4 (Table 1). The correlations were rather weak (small r or even the opposite sign of
395 r) during 2014 due to the short measuring period and narrow ranges of the variables. Water level was positively correlated
396 with NEE and negatively correlated with CH_4 , which was counter intuition, possibly because it was masked by
397 temperature variation as the water level was in general higher in winter and lower in summer. $[\text{CO}_2]$ and $[\text{CH}_4]$ were not
398 correlated with either NEE or CH_4 although they were shown to be important parameters in ANN model selection.

399 3.5 Estimating radiative forcing from different zones

400 To obtain the climate forcings from each land surface type, we calculated the half-hourly and annual gas fluxes from the
401 vegetated area based on eq. (7) using footprint-weighted spatial fraction, ecosystem fluxes and diffusive fluxes from the
402 open water (See Sect. 2.5). The annual median value of footprint-weighted spatial extent was used to calculate the annual
403 fluxes, which showed open-water area was a CO_2 source ($297.5 \text{ g C-CO}_2 \text{ m}^{-2} \text{ yr}^{-1}$) and vegetated area was a CO_2 sink ($-$
404 $39.5 \text{ C-CO}_2 \text{ m}^{-2} \text{ yr}^{-1}$). Both open-water and vegetated area were CH_4 sources but the CH_4 emission from vegetated area
405 was higher than open-water area, being 4.26 and $1.73 \text{ g C-CO}_2 \text{ m}^{-2} \text{ yr}^{-1}$, respectively (Table 2).

406 Open water has contributed large amount of CO_2 emission into the atmosphere through diffusion ($1.09 \text{ kg CO}_2\text{-eq m}^{-2} \text{ yr}^{-1}$
407 $^{-1}$) whereas the CH_4 emission was relatively small ($0.104 \text{ kg CO}_2\text{-eq m}^{-2} \text{ yr}^{-1}$). Vegetated area was a small sink of CO_2 but
408 the cooling effect of vegetation by CO_2 uptake was relatively small ($-0.145 \text{ kg CO}_2\text{-eq m}^{-2} \text{ yr}^{-1}$) compared to its CH_4
409 emission ($0.256 \text{ kg CO}_2\text{-eq m}^{-2} \text{ yr}^{-1}$). Overall, the ecosystem had a small warming effect with $0.263 \text{ kg CO}_2\text{-eq m}^{-2} \text{ yr}^{-1}$
410 of which 89% was contributed by CH_4 (Table 2).

411 4 Discussion

412 4.1 The GHG fluxes from an urban stormwater wetland ecosystem

413 The studied urban wetland ecosystem was a small carbon source over the full-year studied period in year 2013. Due to
414 the scarcity of studies on urban wetlands using the EC method, we compare our results to restored wetlands which can be
415 considered to be proxy ecosystems to urban wetlands with both including rewetting practice in an ecosystem which has
416 been drained previously. The annual CO_2 balance of $8 \text{ g C-CO}_2 \text{ m}^{-2} \text{ yr}^{-1}$ from the ecosystem, or $-39.5 \text{ g C-CO}_2 \text{ m}^{-2} \text{ yr}^{-1}$
417 from the vegetated area (Table 2), were small compared to a restored wetland in western Denmark where the annual CO_2
418 balance ranged from -286 to $-53 \text{ g C-CO}_2 \text{ m}^{-2} \text{ yr}^{-1}$ (Herbst et al., 2013), and the annual CH_4 balance of $3.9 \text{ g C-CH}_4 \text{ m}^{-2}$
419 yr^{-1} was less than half of the annual CH_4 emission (between 9 and $13 \text{ g C-CH}_4 \text{ m}^{-2} \text{ yr}^{-1}$) in that study. Over a network of
420 restored freshwater wetlands in the California, the CO_2 sequestration can be up to nearly $700 \text{ g C m}^{-2} \text{ yr}^{-1}$ and CH_4 emission
421 up to $63 \text{ g C m}^{-2} \text{ yr}^{-1}$ (Hemes et al., 2018). It is not surprising that the studied ecosystem appeared to CO_2 neutral as it was
422 recently constructed. The herbaceous vegetation has been allowed to fully self-establish without human intervention and
423 at the early successional stage, plant diversity and biomass were still increasing each year (Wahlroos, 2019). With the
424 vegetation being more developed, a greater CO_2 uptake from the vegetated area can be expected in the following years.
425 The low CH_4 emission observed in this study may be due to the depletion of organic matters in the bottom soil from
426 agricultural uses thus it provided little substrate for anaerobic microbial activity to produce CH_4 . With the accumulation
427 of organic matters in the anoxic wetland sediment, CH_4 production may increase in the future. Certain chemical



428 compounds like Fe in mineral soils can also inhibit CH₄ production leading to much lower ecosystem-scale CH₄ flux
429 (Chamberlain et al., 2018). In the meanwhile, methane-oxidizing bacteria (methanotroph) regulates CH₄ consumption at
430 the soil-water interface. With the ecosystem being used previously as cropland, the physical disturbance of soil may have
431 greatly reduced the methanotroph communities so that the CH₄ oxidation may also be low in the soil (Smith et al.,
432 2000;Saggar et al., 2008). Furthermore, after the initial establishing phase, the ecosystem productivity can also be reduced
433 due to the standing litter that inhibits the generation of new vegetation growth. It was shown that in a restored freshwater
434 wetland the ecosystem was a net CO₂ sink (-804 ± 131 g C-CO₂ m⁻² yr⁻¹) in 2002-2003, six years after the restoration but
435 near CO₂ neutral in 2010-2011 due to the reduced photosynthetic plants (Anderson et al., 2016). Thus, given the urban
436 wetland is sustained for a sufficiently long period, it is still unclear whether the CO₂ uptake from vegetated zone would
437 compensate its CH₄ emission, not considering the large GHG emission from the open-water zone. Thus, similar studies
438 as the present one should be conducted at a later stage after the construction of the wetland to fully reveal the GHG
439 balance of the ecosystem along time.

440 Overall, the ecosystem CO₂ and CH₄ fluxes measured by EC tower ranged from -5.33 to 3.4 g C-CO₂ m⁻² day⁻¹ and from
441 1.0 to 55.2 mg C-CH₄ m⁻² day⁻¹, respectively. They are consistent with the flux ranges provided by other studies on GHG
442 fluxes in restored wetlands (Anderson et al., 2016;Knox et al., 2015;Matthes et al., 2014;Morin et al., 2014b;Herbst et al.,
443 2013), although for both gases they tend to be on the lower end. NEE, GEP and R_{eco} exhibited seasonal variation so that
444 the ecosystem was a CO₂ sink between June and August. The highest NEE appeared in September possibly because GEP
445 has greatly reduced due to plant senescent while R_{eco} remained relatively high because of the warm temperature (Fig. S4).
446 Previous studies have found good agreement between CH₄ emission and GEP as plants provide substrates for
447 methanogenesis (Rinne et al., 2018), which was not observed in the daily-average of gas fluxes in this study (Figs 2 and
448 3) as the peak CH₄ flux appeared in May and peak GEP appeared in July. Nonetheless, both CH₄ and CO₂ fluxes, especially
449 those obtained from vegetated area, exhibited strong diurnal cycle during summer with synchronized peaks around noon
450 (Fig. 5a, 5c). This finding reflects that short-term CH₄ emission from vegetation is linked with photosynthesis by
451 providing labile carbon from root exudate and by gas transport through aerenchyma and open stomata while long-term
452 CH₄ emission may be determined by complex processes related to environmental variables e.g. temperature and redox
453 potential (Linden et al., 2014).

454 4.2 Parsing GHG fluxes from heterogeneous land surfaces

455 We found the open-water area was constantly a source of CO₂ and CH₄ to the atmosphere during the studied period as the
456 [CO₂] and [CH₄] in the water generally exceeded the atmosphere equilibrium except the ice-covered period (Fig. 4). The
457 annual average of [CO₂] in the surface water in 2013 was 0.3% in our study, comparable to 0.4% in another temperate
458 restored wetland (McNicol et al., 2017), while the seasonal pattern (higher in summer and fall) was the opposite as they
459 have found. We also found that both [CO₂] and [CH₄] were higher in 2014 than 2013 (Fig. 1d, 1e). The O₂ concentration
460 ([O₂]) and O₂ balance ($[O_2]_{outlet} - [O_2]_{inlet}$) measured by another study on the same wetland (Wahlroos, 2019) could partially
461 explain the observed phenomenon. The relatively high water temperature and oxic conditions in the water in fall 2013
462 have allowed high decomposition of detritus leading to high [CO₂] (Wahlroos, 2019). The long period of hypoxia during
463 summer 2014 could explain the three-fold increase in [CH₄] as the condition was more favorable for CH₄ production. The
464 negative O₂ balance in summer 2014 indicated strong O₂ consumption by microbial decomposition producing CO₂ in the
465 water. As the long-term diffusive fluxes (daily and monthly) was mainly driven by gas concentration in the water, it was
466 straight forward to understand high diffusive CO₂ and CH₄ fluxes in 2014 comparing to 2013. Interestingly, the ecosystem



467 CH₄ emission in 2013 was well synchronized with the diffusive CH₄ flux by capturing sporadic emission episodes from
468 the water (Fig. S6a, S6c) while they were not synchronized in summer 2014 although several stronger diffusive peaks
469 happened (Fig. S6b, S6d). When footprint-weighted contribution was accounted for, it clearly revealed that the
470 synchronization of CH₄ emission from ecosystem and water was closely related to the flux footprint distribution. When
471 there was high flux contribution from the open water (20-25 %), high diffusive CH₄ was also reflected in ecosystem flux
472 measured by EC. This has further proved the application of footprint analysis is essential in explaining gas exchange from
473 heterogeneous surfaces using EC data.

474 It is worth noticing that in our study we only classified the surface landscapes into “open water” and “vegetation” but
475 neglected the difference in sink/source strength from different plant types within the vegetation zone (Fig. S1). We did
476 not account for the dissimilarity between vegetation types because the characteristics in gas exchange are much more
477 distinct between open water and vegetation, which was the focus of this study. For the same reason, ebullition was not
478 considered in this study neither, as ebullition was shown to have only minor significance in a restored wetland accounting
479 for less than 0.1% of ecosystem CO₂ flux and 4.1% of ecosystem CH₄ flux (McNicol et al., 2017). However, for a proper
480 downscaling analysis of EC data, the subareas of different plant types and ebullition should also be taken into account.

481 **4.3 Climate impact of urban wetland and implications for management**

482 In the present study, the urban boreal wetland had an overall SGWP of 0.263 kg CO₂-eq m⁻² yr⁻¹ which was comparable
483 or higher than other restored wetlands in boreal region (Herbst et al., 2013), and within the range of inter-annual variation
484 or lower than restored wetlands in temperate zone (McNicol et al., 2017; Anderson et al., 2016). Different from other
485 studies, the urban wetland was CO₂ neutral and a CH₄ source. It is worth noting that the paramount contribution of CH₄
486 in ecosystem SGWP was mainly driven by the large footprint-weighted spatial area of vegetation (See Sect. 3.2). In fact,
487 The SGWP of GHG emission from open water (1.194 kg CO₂-eq m⁻² yr⁻¹) was 10 times as large as that from vegetation
488 (0.111 kg CO₂-eq m⁻² yr⁻¹) (Table 2). The implication of this result is that during wetland restoration, it would be more
489 beneficial to have large patches of emergent vegetation area at least from the GHG emission point of view. Similar results
490 have been obtained by other studies as well that open water has more climate-warming impact than emergent vegetation
491 due to the large diffusive fluxes from open water (Stefanik and Mitsch, 2014; McNicol et al., 2017). The climate impact
492 of natural wetland depends on the net balance between the cooling effect of CO₂ uptake by vegetation and the warming
493 effect of other GHG emissions, mainly CH₄ (Bridgman et al., 2013). In wetlands constructed in urban area, the large
494 fraction of open water which is a significant emitter of CO₂, should also be taken into consideration when evaluating the
495 role of urban wetland in global climate change.

496 **5 Conclusions**

497 Urban wetlands have received global attention as a nature-based urban runoff management solution for sustainable cities,
498 as they provide cost efficient flood control and water quality mitigation as well as many ecological and cultural services.
499 In the meantime, the climate impact of urban wetlands should also be considered. Wetting a landscape may enhance the
500 CO₂ sequestration in the ecosystem, whereas CH₄ can be emitted due to the anaerobic conditions in the soil after wetting.
501 Furthermore, heterogeneity induced in newly created urban wetland may contribute differently to the overall climate
502 impact.



503 In the present study, for the first time a full annual carbon balance of an urban stormwater wetland in the boreal region
504 was evaluated and the radiative forcing from heterogeneous landscapes were presented. We found that, during the
505 monitored period at the study wetland, both the open water area and the vegetated area within the created wetland were
506 carbon sources, and thus the urban wetland had a net climate warming effect, the monitored fourth year after the wetland
507 establishment. The radiative forcing effect of the open-water area exceeded the vegetated area, which indicated that
508 limiting open-water surfaces and setting a design preference for areas of emergent vegetation in the establishment of
509 urban wetlands can be a beneficial practice when considering only the climate impact of a created urban wetland. In the
510 meanwhile, we also emphasize that the value of urban wetlands should not be determined solely by GHG radiative forcing.
511 The values of urban wetlands in other areas e.g. flood control, pollutant removal, biodiversity, recreation and education
512 are as well of paramount importance to human society.

513

514 **Data availability**

515 Eddy covariance, gas concentration and meteorological data are available from the DRYAD database at
516 https://datadryad.org/stash/share/WrtTNnpIt6FgLoMSZ_Wlr0IK22IcxqjGZAStuuKdHLS

517 **Author contribution**

518 IM, OW, HV, AO and TV designed the field study. SH, IM and JP carried out eddy covariance measurements, automatic
519 gas concentration measurements in the open water and manual field measurements. XL and IM participated in eddy
520 covariance data processing and analysis. XL analysed the results and prepared the manuscript with contributions from all
521 co-authors.

522 **Competing interests**

523 The authors declare that they have no conflict of interest.

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657 **Tables**

658 Table 1. Pearson correlation coefficient (r) between the daily averages of environmental variables and fluxes in year
659 2013 and 2014. NEE – net ecosystem exchange; T_{air} – air temperature; T_{water} – water temperature; PPFD –
660 photosynthetic photon flux density; WL – water level; $[\text{CO}_2]$ and $[\text{CH}_4]$ – CO_2 and CH_4 concentration measured in the
661 outlet; * indicates only peak growing season (June, July and August) are included in the analysis.

Flux	Year	T_{air}	T_{water}	PPFD	WL	$[\text{CO}_2]$	$[\text{CH}_4]$
CO_2	2013	-0.45	-0.61	-0.62	0.46	-0.34	0.18
	2014	0.43	0.54	-0.12	0.12	-0.12	-0.05
CH_4	2013	0.61	0.65	0.56	-0.3	0.17	-0.09
	2014*	0.37	0.26	0.27	-0.24	0.28	0.25

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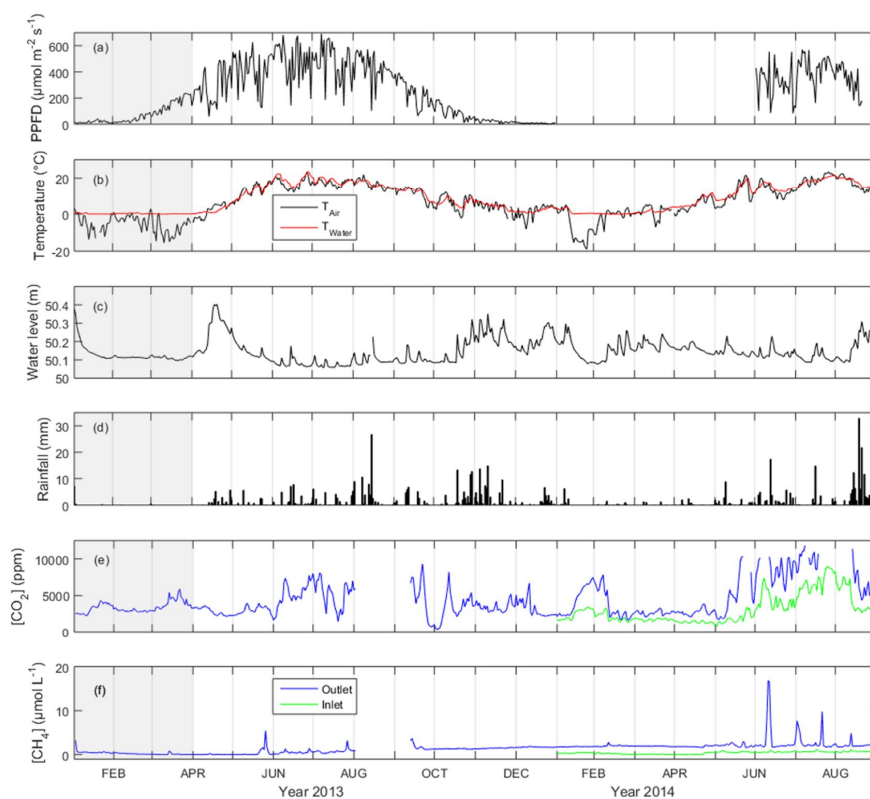
664 Table 2. Annual CO₂ and CH₄ exchange from different surface zones and their sustained global warming potential
665 (SGWP). Ecosystem, water and vegetation represent flux and SGWP measured or calculated from the ecosystem by EC
666 tower, from open water and from vegetated area. The numbers in the square bracket represent the 95% confidence interval
667 of the average. No error bounds are reported for flux and SGWP from water as they are modelled using gas concentration
668 in the water and meteorological measurements.

		Ecosystem	Water	Vegetation
Flux (g C m ⁻²)	CO ₂	8 [-18.9, 34.9]	297.5	-39.5 [-70.8, -8.1]
	CH ₄	3.9 [3.8, 4.1]	1.7	4.3 [4.1, 4.5]
SGWP (kg CO ₂ -eq m ⁻²)	CO ₂	0.029 [-0.069, 0.128]	1.090	-0.145 [-0.260, -0.030]
	CH ₄	0.234 [0.225, 0.244]	0.104	0.256 [0.246, 0.268]

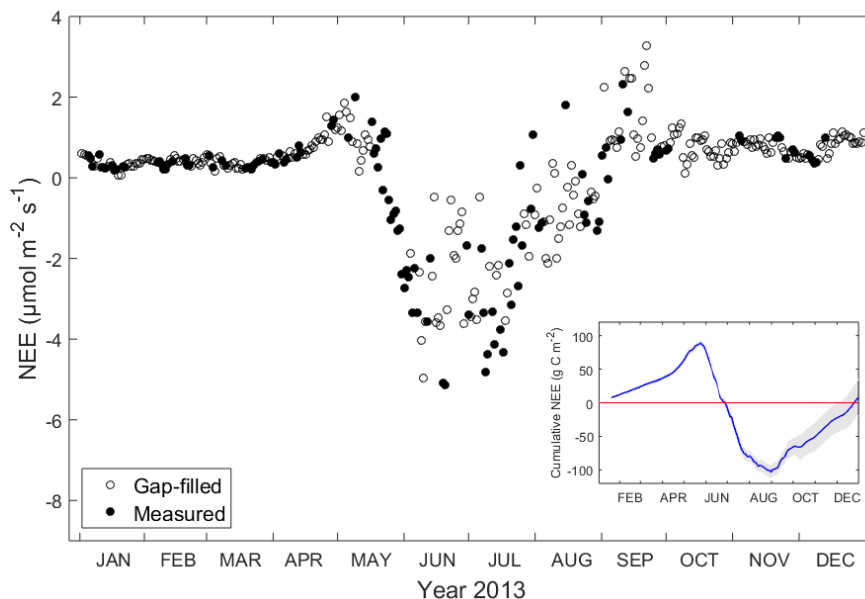
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670 **Figures**



671
672 Figure 1: The daily-average of (a) photosynthetic photon flux density (PPFD), (b) air and water temperature, (c) water
673 level, (d) rainfall, (e) CO₂ concentration and (f) CH₄ concentration from inlet and outlet of Nummela wetland from
674 January 2013 to August 2014. The grey zone indicates the ice-covered period.



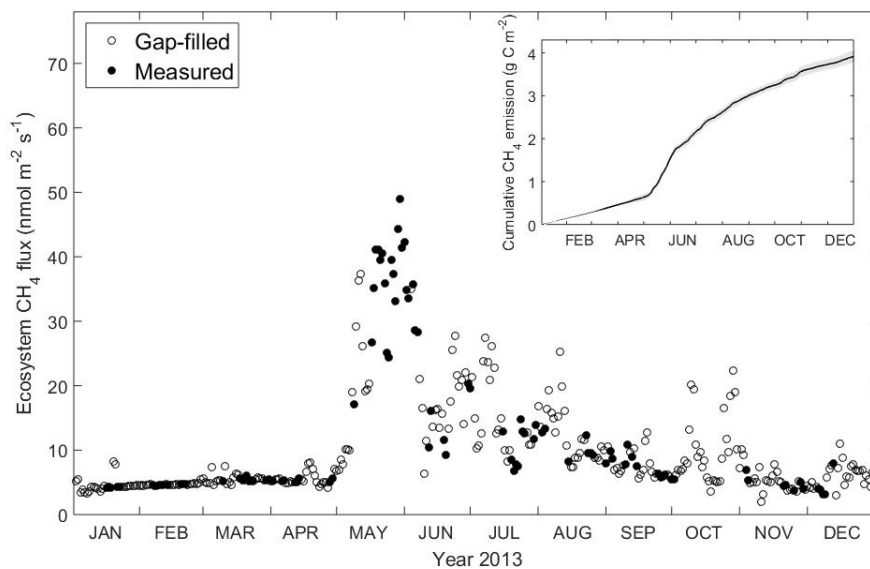
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676 Figure 2: Daily average of net ecosystem exchange of CO₂ (NEE, $\mu\text{mol m}^{-2} \text{s}^{-1}$) in year 2013. Filled dots indicate
677 measurement (when available half-hourly measurement data ≥ 10) and circles indicate gap-filled data (when available half-
678 hourly measurement data < 10). The insert shows cumulative NEE (g C m^{-2}) in the ecosystem and the red line indicates
679 the zero reference line. Error bounds (marked in grey) on cumulative NEE reflect the 95 % confidence interval for the
680 gap-filling procedure.

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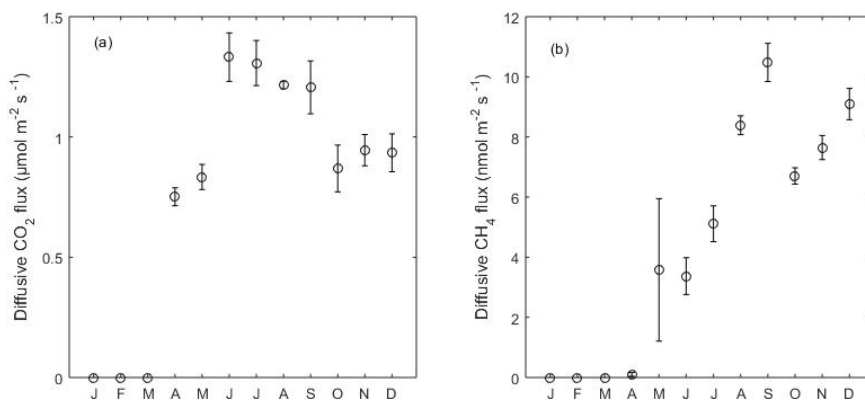


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685 Figure 3: Daily average of ecosystem CH₄ flux measured by EC tower and cumulative CH₄ emission in year 2013. Filled
 686 dots indicate measurement (when available half-hourly measurement data ≥ 10) and circles indicate gap-filled data (when
 687 available half-hourly measurement data < 10). The insert shows cumulative CH₄ emission with the error bounds in grey
 688 reflecting the 95 % confidence interval for the gap-filling procedure.

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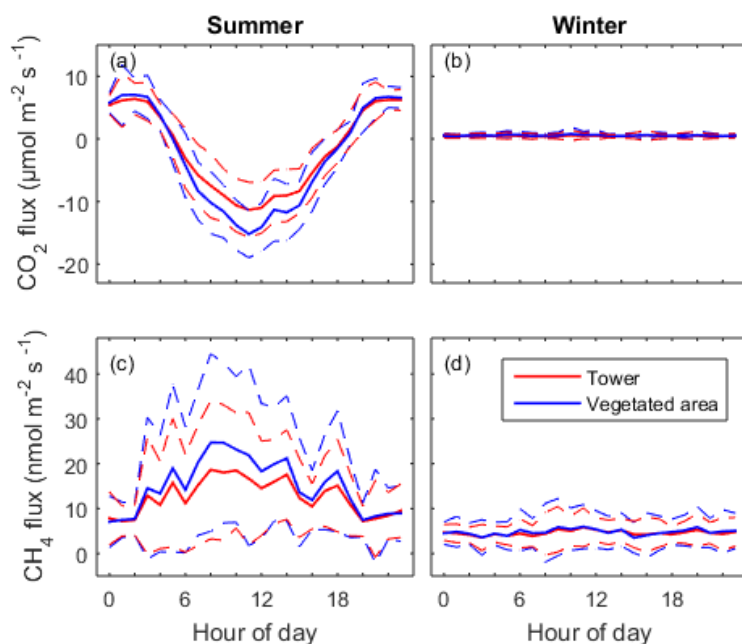
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692 Figure 4: Monthly-average of (a) diffusive CO₂ and (b) CH₄ flux from the open-water in year 2013. Error bar indicates
 693 the standard error of the mean. From January to March there was ice-covered period.

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695

696 Figure 5: Mean diel pattern of the half-hourly net CO₂ and CH₄ fluxes in summer ((a) and (c)) and in winter ((b) and
697 (d)). The dashed lines represent the standard deviation. Red lines indicate measurement from EC tower and the blue
698 lines show the fluxes modelled for vegetated area.

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