

1 **Title**  
2 Resolving space and time variation of lake-atmosphere carbon dioxide fluxes using multiple  
3 methods  
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19 **Abstract**

20 Lakes emit globally significant amounts of carbon dioxide (CO<sub>2</sub>) to the atmosphere, but  
21 quantifying these rates for individual lakes is extremely challenging. The exchange of CO<sub>2</sub>  
22 across the air-water interface is driven by physical, chemical, and biological processes in both  
23 the lake and the atmosphere that vary at multiple spatial and temporal scales. None of the  
24 methods we use to estimate CO<sub>2</sub> flux fully capture this heterogeneous process. Here, we  
25 compared concurrent CO<sub>2</sub> flux estimates from a single lake based on commonly used methods.  
26 These include floating chambers (FC), eddy covariance (EC), and two concentration gradient  
27 based methods labelled fixed (F-*p*CO<sub>2</sub>) and spatial (S-*p*CO<sub>2</sub>). At the end of summer, cumulative  
28 carbon fluxes were similar between EC, F-*p*CO<sub>2</sub> and S-*p*CO<sub>2</sub> methods (-4, -4 and -9.5 gC), while  
29 methods diverged in directionality of fluxes during the fall turnover period (-50, 43 and 38 gC).  
30 Collectively these results highlight the discrepancies among methods and the need to  
31 acknowledge the uncertainty when using any of them to approximate this heterogeneous process.  
32

33 **Plain Language Summary**

34 Lakes comprise a small percentage of the landscape, but they are active and complex areas of  
35 carbon cycling. Lakes receive mixed carbon inputs from upstream sources, process this carbon  
36 internally, store it in sediments and biomass, and export it downstream. In addition, some  
37 fraction of the carbon in lakes exchanges into and out of the atmosphere, linking lakes with the  
38 global atmosphere. The exchange of carbon dioxide across lake surfaces has globally significant  
39 implications but quantifying these rates has yet to be fully resolved. Here, we compared four  
40 methods of estimating diffusive carbon dioxide exchange between the atmosphere and the lake  
41 surface. Flux rates generally agreed during the summer, but estimates diverged in the fall, a  
42 critical time period with elevated carbon cycling rates. These discrepancies among methods may  
43 arise because of the high degree of spatial and temporal variability in gas exchange and our  
44 ability to portray these processes accurately. In the future, we need to improve our observational  
45 resolution to better estimate carbon gas exchange between lakes and the atmosphere.  
46

47 **Key Points**

- 48 1) Lake-atmosphere CO<sub>2</sub> exchange estimate using four common methods  
49 2) CO<sub>2</sub> concentration gradient flux estimates agreed in direction and magnitude with floating  
50 chambers but disagreed with eddy covariance.  
51 3) Inconsistencies among methods highlight the spatial and temporal assumptions underlying  
52 methods and the need to acknowledge uncertainty  
53

54 **Author Contribution Statement**

55 DER and ARD collected eddy covariance data, LCL and EHS spatial gradient data, EHS and  
56 ARD collect fixed spatial gradient data, while HH and AKB collected chamber data. AKB led  
57 analysis with assistance from DER and LCL. AKB and DER drafted the manuscript, all authors  
58 helped edit the manuscript.  
59

60 **Competing Financial Interests**

61 The authors declare no competing financial interests.

## 62 **1. Introduction**

63           Lakes are a major component of the Earth's carbon cycle and an increasing focus has  
64 been placed on carbon dynamics within inland waters [*Biddanda, 2017; Tranvik et al., 2009;*  
65 *Williamson et al., 2009*]. A substantial fraction of the organic carbon that is delivered to or fixed  
66 within lakes is outgassed to the atmosphere as carbon dioxide (CO<sub>2</sub>) [*Cole et al., 2007; Cory et*  
67 *al., 2014*]. While there is consensus that collectively lakes and other inland waters emit  
68 meaningful amounts of CO<sub>2</sub> to the atmosphere, it remains extremely difficult to calculate  
69 spatially and temporally resolved emission rates for individual lakes. This difficulty is because  
70 the exchange of CO<sub>2</sub> across the air-water interface is driven by multiple physical, chemical, and  
71 biological processes in both the lake and the atmosphere that vary at multiple spatial and  
72 temporal scales. The scientific community lacks methods to fully capture the spatial and  
73 temporal heterogeneity in gas exchange between lakes and the atmosphere. Thus, every estimate  
74 of global CO<sub>2</sub> emissions from lakes has uncertainty.

75           The reason that lake-atmosphere fluxes are difficult to quantify is because they vary in  
76 magnitude [*Raymond et al., 2013*], in time [*Reed et al., 2018*], and across space [*Natchimuthu et*  
77 *al., 2016*]. In many temperate dimictic lakes, seasonal phenologies in ice-cover and stratification  
78 govern the direction and magnitude of CO<sub>2</sub> flux. Large off-gassing events occur during periods  
79 of vertical mixing such as ice-off and fall turnover [*Denfeld et al., 2016*]. Lakes with higher  
80 productivity show pronounced temporal variation in CO<sub>2</sub> flux [*S C Maberly et al., 2013*],  
81 characterized by influx during the summer periods coinciding with higher rates of primary  
82 production [*Reed et al., 2018*]. Thus, for even a single lake, flux estimation needs to be  
83 continuous and year-round in order to capture the temporal heterogeneity in gas exchange.  
84 Spatially, heterogeneity in metabolic processes, hydrology, and turbulence can have pronounced

85 impacts on CO<sub>2</sub> flux from the lake surface. Rivers flowing into lakes typically differ in a number  
86 of physical, chemical, and biological properties that can create contrasts in pCO<sub>2</sub> in habitats  
87 where they enter a lake [Chmiel *et al.*, 2019]. Further, spatial heterogeneity varies temporally [L  
88 C Loken *et al.*, 2019; Natchimuthu *et al.*, 2016] due to changes in river flow, lake mixing, and  
89 biological processes. Thus, to accurately measure CO<sub>2</sub> flux from a single lake we need to  
90 incorporate both spatial and temporal processes.

91 Any calculations of lake-atmosphere CO<sub>2</sub> flux are limited in either spatial or temporal  
92 extent. Perhaps the simplest and most cost-effective method for measuring gas efflux from lakes  
93 is using floating chambers (FC) [Bastviken *et al.*, 2015]. Chambers are placed atop the lake  
94 surface and the flux is derived from the gas accumulation rate within the chamber. However, flux  
95 chambers characterize only a small area of the lake for what is typically a short deployment, and  
96 can alter turbulence and thus gas exchange within the chamber environment [Vachon *et al.*,  
97 2010]. Historically, FC's for CO<sub>2</sub> required manual gas sampling followed by laboratory  
98 determination of gas concentrations, while newer FCs integrate continuous CO<sub>2</sub> sensors and  
99 automatic purging mechanisms that allow for longer deployments [Bastviken *et al.*, 2015;  
100 Jonsson *et al.*, 2008; Martinsen *et al.*, 2018]. While a single measurement is small in its spatial  
101 scale, multiple chambers have been used to quantify the spatial variability of gas emissions  
102 within and among lake habitats [Natchimuthu *et al.*, 2016; Tangen *et al.*, 2016]. Similarly,  
103 measuring temporal variability of fluxes using FCs is common but in both cases, characterizing  
104 spatial and/or temporal variability with this approach is time intensive. New automated chambers  
105 show promise in increasing the duration of continuous observation [Duc *et al.*, 2012].

106 A common alternative to FCs is modeling exchange rates using the concentration  
107 gradient or boundary layer method (F-pCO<sub>2</sub>) [Cole and Caraco, 1998] that is based on

108 differences in  $p\text{CO}_2$  between the lake surface and the atmosphere and an estimate of water  
109 turbulence or gas transfer velocity ( $k$ ). Spatial scales of  $p\text{CO}_2$  measurements within the water  
110 column are on the order of cubic centimeters and typically fixed in space. Estimation of  $k$  is  
111 typically based on empirically derived models using wind-speed, lake size, and/or water density  
112 gradients [Crusius and Wanninkhof, 2003; MacIntyre et al., 2010; Read et al., 2012].  $k$  can also  
113 change in response to environmental conditions [Natchimuthu et al., 2016; Vachon and Prairie,  
114 2013]; moreover, estimation of  $k$  can vary by multiple orders of magnitude simply due to model  
115 choice [Dugan et al., 2016]. Recent  $p\text{CO}_2$  studies have shown the scaling  $k$  from point  
116 measurements to the lake scale strongly underestimates emissions [Mammarella et al., 2015;  
117 Schubert et al., 2012]. New methods have been developed to quickly quantify spatial variation in  
118  $p\text{CO}_2$  and  $k$  [Bastviken et al., 2015; Crawford et al., 2015] and have revealed substantial spatial  
119 variations in  $p\text{CO}_2$  and fluxes within individual lakes and reservoirs [L C Loken et al., 2019;  
120 Natchimuthu et al., 2016; Paranaíba et al., 2018]. Boundary layer methods have provided the  
121 most frequent and comprehensive understanding of  $\text{CO}_2$  exchange between lakes and the  
122 atmosphere, yet most assume spatial homogeneity and are reliant on physical lake models that  
123 have large uncertainty.

124 A third approach for quantifying lake  $\text{CO}_2$  fluxes is eddy covariance (EC) [Morin et al.,  
125 2018; Reed et al., 2018]. In contrast to the water-based approaches, EC uses measurements of  
126 concentrations of gas in the atmosphere along with high-frequency measurements of wind speeds  
127 in 3 dimensions. While this top-down flux method seems like the silver bullet for quantifying  
128  $\text{CO}_2$  flux, EC has several assumptions built into estimation and is spatially limited. It relies on  
129 measurement during windy periods and includes uncertainty of footprint models that estimate the  
130 area over which fluxes are being measured (i.e., the footprint), with a single flux estimate

131 integrating over 100's of square meters. Turbulence and footprint issues can lead to upwards of  
132 80% of EC data being excluded [Reed *et al.*, 2018]. EC estimates represent the average flux from  
133 a portion of the lake surface, which bias observations toward near-shore areas [Morin *et al.*,  
134 2018] where most towers are located. Despite these limitations, EC offers a promising method  
135 for assessing carbon fluxes from lakes [Vesala *et al.*, 2012].

136         Because each technique for measuring carbon flux has its limitations, efforts have been  
137 made to compare these methods. However, these investigations have been limited to relatively  
138 short time periods [Erkkila *et al.*, 2018; Podgrajsek *et al.*, 2016; Schubert *et al.*, 2012]. These  
139 authors found discrepancies among methods for quantifying CO<sub>2</sub> flux in both space and time.  
140 While estimates of carbon fluxes are critical for global carbon cycling, how best to measure lake-  
141 atmosphere fluxes remains challenging and is an open question for the scientific community.

142         In order to compare methods of quantifying lake-atmosphere fluxes of CO<sub>2</sub>, we leveraged  
143 multiple concurrent datasets from a single north temperate lake (Lake Mendota, Wisconsin,  
144 USA). This lake has been subject to prior CO<sub>2</sub> flux investigations [L C Loken *et al.*, 2019; Reed  
145 *et al.*, 2018]. Here, we combined flux records based on measurements of *p*CO<sub>2</sub> at a moored buoy,  
146 measurements distributed across the entire lake surface, EC from a tower located at the end of a  
147 narrow peninsula, and FC. The overarching question of this work is: Are lake-atmosphere CO<sub>2</sub>  
148 flux estimates consistent among *p*CO<sub>2</sub>, FC, and EC methods? Due to multiple temporal and  
149 spatial scales which the independent observations are taken over, we seek to answer the question  
150 using 1) analysis of flux distribution over multiple seasons, 2) quantifying cumulative sums of  
151 carbon flux, 3) direction comparison of methods, and 4) spectral time-series analysis of fluxes.

152

## 153 **2. Methods**

154 **2.1 Site Description**

155 Lake Mendota is a well-studied lake located in Southern Wisconsin, USA (43.1° N, 89.4°  
156 W) and is part of the North Temperate Lakes Long-Term Ecological Research (NTL-LTER)  
157 program. It is dimictic and eutrophic, with a surface area of 39.9 km<sup>2</sup>, a maximum depth of 25.3  
158 m (mean 12.7 m). The majority of the lake's watershed is composed of agricultural and urban  
159 land uses, resulting in elevated nutrient concentrations and high productivity [*Carpenter et al.*,  
160 2007]. Thermal stratification typically occurs between May and October and ice cover from late  
161 December through March. We defined seasons using water column temperature gradients with  
162 spring and fall as periods in which the water column was isothermal, while in summer the lake  
163 was thermally stratified.

164  
165 **2.2 Flux Estimates**

166 **2.2.1 Fixed point concentration gradient method (F-*p*CO<sub>2</sub>)**

167 Since 2006, NTL-LTER has managed a monitoring buoy on Lake Mendota that is  
168 moored above the lake's deepest point (43.0995°N, 89.4045°W). The buoy is equipped with  
169 meteorological and limnological sensors and is deployed seasonally (~April through October),  
170 capturing the majority of the ice-free season. In 2015, a Turner Designs C-sense *p*CO<sub>2</sub> sensor  
171 (Turner Designs, San Jose, USA; 3% accuracy) was added to the buoy and installed at 0.5 m  
172 depth. For this study, we used wind speed, surface water temperature, and surface *p*CO<sub>2</sub>  
173 [*Magnuson et al.*, 2019]. Wind speed was measured at a height of 2.7 m above the lake surface  
174 using an anemometer (R. M. Young Marine Wind Monitor). Water temperature and *p*CO<sub>2</sub> were  
175 measured at a depth of 0.5m using a RBR concerto thermistor string and a Turner C-Sense CO<sub>2</sub>  
176 sonde, respectively. Wind speed and water temperature were measured every 30 minutes, while

177  $p\text{CO}_2$  was measured every 15 minutes.  $p\text{CO}_2$  in air was measured from an in situ spectroscopy  
178 gas analyzer (Picarro, inc. 4-Species Gas Analyzer) located at a nearby building,

179 Using data collected at the buoy, we calculated the diffusive efflux of  $\text{CO}_2$  from the lake  
180 surface to the atmosphere according to:

$$181 \quad \text{Flux} = k_{\text{gas}} \times kh \times (p\text{CO}_{2\text{water}} - p\text{CO}_{2\text{air}}) \quad (1)$$

182 This fixed-point boundary layer method (F- $p\text{CO}_2$ ) is based on the partial pressure gradient  
183 between the water ( $p\text{CO}_{2\text{water}}$ ) and the atmosphere ( $p\text{CO}_{2\text{air}}$ ). Multiplying this difference by the  
184 Henry's law constant ( $kh$ ) converts to molar units and by the gas transfer velocity ( $k_{\text{gas}}$ ) to  
185 generate diffusive flux estimates. We estimated  $k_{\text{gas}}$  using concurrent wind speed and water  
186 temperature recorded at the buoy, applying the  $k_{600}$  model and Schmidt model coefficients  
187 provided in *Raymond et al.* [2013]. The Henry's law constant ( $kh$ ) was calculated using  
188 atmospheric pressure and temperature-dependence models provided in *Plummer and Busenberg*  
189 [1982] See *L C Loken et al.* [2019] for further description of the  $p\text{CO}_2$  flux model.  $p\text{CO}_2$  flux  
190 estimates were computed at 30-minute intervals. To temporally match observations between  
191 methods, a subset of F- $p\text{CO}_2$  was used from 8 a.m.-12:00 p.m., the time period that overlapped  
192 with the majority (>90%) of the spatially-explicit  $p\text{CO}_2$  sampling times (described below).

193

### 194 **2.2.2 Spatial concentration gradient method (S- $p\text{CO}_2$ )**

195 In addition to the F- $p\text{CO}_2$ -based flux estimation at the buoy, we also compared flux  
196 estimates using  $p\text{CO}_2$  measurements from the entire lake surface (S- $p\text{CO}_2$ ) from *L C Loken et al.*  
197 [2019]. For the entire ice-free period of 2016, [*L C Loken et al.*, 2019] generated  $\text{CO}_2$  efflux  
198 estimates at 988 points distributed in a gridded pattern across the lake surface. Efflux  
199 measurements were based on measurements of  $p\text{CO}_2$  that were distributed across the entire lake

200 surface. Similar to the F- $p\text{CO}_2$  method, efflux was calculated using the difference in  $p\text{CO}_2$   
201 between the water and the air. *L C Loken et al.* [2019] used a spatially explicit  $k$  model [*Vachon*  
202 *and Prairie*, 2013], which takes into account wind speed and direction and allows  $k$  to vary  
203 across the lake surface.  $p\text{CO}_2$  measurements were collected over a ~3-hour window in the  
204 morning during each survey, and efflux was estimated at daily time scales. Two subsets of S-  
205  $p\text{CO}_2$  data was used to quantify spatial variability, 10 stratified random points from the entire  
206 lake and S- $p\text{CO}_2$  measurement locations from within the EC footprint.

207

### 208 **2.2.3 Flux Chamber Diffusion Method (FC)**

209 We conducted four FC campaigns between July 6, 2017 and April 24, 2018.  $\text{CO}_2$  sensors  
210 (Sensair K30) were installed inside floating chambers with a diameter of 0.3 m and a height of  
211 0.12 m. Flux rates were calculated using the chamber dimensions (surface area and volume) and  
212 continuous  $p\text{CO}_2$  measurements within the enclosed headspace. Each 24-hr sampling campaign  
213 consisted of 7 sampling trips spaced every 4 hours with the goal of measuring flux rates over a  
214 complete diel cycle. For each measurement, we placed two chambers on the lake surface in the  
215 middle of the lake (same location as the buoy) and let them drift for 5 minutes. We repeated the  
216 FC procedure 3 times per chamber and calculated the average of the 6 flux measurements.  $\text{CO}_2$   
217 flux was calculated as:

$$218 \quad \text{Flux} = \frac{\Delta p\text{CO}_2}{\Delta t} \times \frac{V}{SA} \quad (2)$$

219 where  $V$  is the chamber volume ( $0.03114 \text{ m}^3$ ),  $SA$  is the chamber bottom area ( $0.071 \text{ m}^2$ ), and  $t$   
220 is time (s). Prior to the first campaign, we calibrated all sensors using  $\text{N}_2$  gas and the “zero  
221 calibration” method per *Bastviken et al.* [2015]. For all subsequent campaigns we re-confirmed  
222 the zero  $\text{CO}_2$  readings using  $\text{N}_2$  gas.

223

#### 224 **2.2.4 Eddy Covariance Tower**

225 Eddy covariance flux observations (Ameriflux site: US-PnP, doi:  
226 10.17190/AMF/1433376) were collected from a tower at the end of a ~50 m wide peninsula on  
227 the shore of Lake Mendota (Figure 1) starting on June 20, 2016. These flux observations were  
228 made with a sonic anemometer (CSAT3, Campbell Scientific, Logan UT, USA) and open-path  
229 infrared gas analyzer for CO<sub>2</sub> and water vapor gas concentration (LI-7500A, Li-Cor, Lincoln,  
230 NE, USA) at a height of 12.4 m above the lake on a 0.95 m boom, along with measurements of  
231 air temperature and humidity (Vaisala, Inc. HMP45C). Measurements of incoming solar  
232 radiation and atmospheric pressure were collected from a nearby meteorological tower located  
233 on the roof of the Atmospheric, Oceanic, and Space Sciences building.

234 Eddy fluxes were calculated based on the covariance of vertical wind velocity and scalar  
235 concentrations following the approach of *Mauder and Foken* [2015], with quality control flags  
236 for stationarity, integral turbulence, and propagates estimates of random error. Using an eddy  
237 flux surface flux footprint model [*Kljun et al.*, 2015], we identified and removed non-lake data at  
238 30-minute time-scales, primarily when winds were from the forested portion of the peninsula.  
239 After footprint screening and quality control, 26% of data were remaining.

240

#### 241 **2.3 Comparison of methods**

242 Flux estimates varied in temporal and spatial coverage (Table 1). EC-based fluxes were  
243 collected continuously since 2016. Buoy-based F-*p*CO<sub>2</sub> estimates are also continuous since this  
244 time, with the exception of winter months. We only have S-*p*CO<sub>2</sub> rates for the ice-free period of  
245 2016, which were collected ~weekly and daily rates were modelled by interpolating *p*CO<sub>2</sub>

246 through time (see *L C Loken et al.* [2019] for details). Thus, these three data sources (*S-pCO<sub>2</sub>*, *F-*  
247 *pCO<sub>2</sub>*, and EC) overlapped throughout the ice-free period in 2016. We collected FC flux rates  
248 seasonally starting in summer 2017 (July 28-29, 2017, Oct 28-29, 2017, and April 23-24, 2018).  
249 Thus, there are three 24-hr intervals where FC, EC, and *F-pCO<sub>2</sub>* estimates overlapped in time. In  
250 addition to temporal overlap, we must also consider spatial coverage as sampling sites varied  
251 among methods. Both the FC- and *F-pCO<sub>2</sub>*-based rates were determined at the center of the lake.  
252 The EC rates reflect the area surrounding the tower along the lake's southern shoreline, and *S-*  
253 *pCO<sub>2</sub>* covered the entire lake surface (Figure 1).

254 Because of varying temporal resolution among datasets, we converted all datasets to daily  
255 averages, representing the coarsest temporal scale. Using the *S-pCO<sub>2</sub>* flux estimates, we  
256 generated two additional spatial datasets. First we randomly selected 10 stratified points from the  
257 entire lake to visualize spatial variability across the lake. Second, we subset the *S-pCO<sub>2</sub>* dataset  
258 by only including flux estimates from within the EC footprint for a comparison between these  
259 two methods that was not confounded by differences in sampling areas. Cumulative fluxes from  
260 2016 were calculated from *F-pCO<sub>2</sub>*, *S-pCO<sub>2</sub>*, and EC observations.

261 In addition to comparing similarity in seasonal pattern and magnitude, we also wanted to  
262 determine if the different methods exhibited similar temporal variance. To do so, we calculated  
263 Fourier power spectra of each daily time series. Data analysis was done in Matlab R2019a and  
264 IDL 8.6.0.

265

## 266 **3. Results**

### 267 **3.1 Patterns among methods**

268 Footprint modeling revealed that the EC footprint originated primarily from open water,  
269 with very little apparent input from the terrestrial peninsula (Figure 2a), with the distance of  
270 maximum contribution of fluxes on average being 40 m while the distance containing 80% of  
271 flux contribution was 410 m. Friction velocity ( $u^*$ ) values were high due to winds crossing the  
272 peninsula, showing increased turbulence due to the tree canopy (Figure 2b). While winds  
273 originated from all directions, wind speeds were lower over the peninsula as well (Figure 2c).  
274 These factors combined to limit the footprint along the narrow range of wind directions over the  
275 peninsula.

276 In all years, F- $p\text{CO}_2$  flux estimates followed a similar pattern of near-zero or slightly  
277 negative fluxes denoting  $\text{CO}_2$  movement from the atmosphere to the lake during spring and  
278 summer months before becoming strongly positive (net  $\text{CO}_2$  efflux from the lake to the  
279 atmosphere) in the fall (Figures 3, 4). Daily-averaged fluxes varied from  $-1.2$  to  $4.1 \mu\text{m}^{-2} \text{s}^{-1}$   
280 across all dates with a CV of 8.42. This same pattern was also demonstrated by the S- $p\text{CO}_2$   
281 method (Figures 3, 4), and flux estimates were similar in magnitude and direction as the F- $p\text{CO}_2$   
282 results in 2016 ( $-0.39$  to  $1.6 \mu\text{m}^{-2} \text{s}^{-1}$ , CV of 6.34). The limited set of FC deployments also  
283 followed the same general pattern of  $\text{CO}_2$  influx to the lake in spring, a weaker influx during  
284 summer, and efflux in the fall (Figure 3b-d). However, the range of FC flux values were  
285 narrower than for the two  $p\text{CO}_2$ -based methods ( $-1.6$  to  $2.1 \mu\text{m}^{-2} \text{s}^{-1}$ , CV of 8.72).

286 Fluxes derived from the EC method were characterized by higher variation, often shifting  
287 from negative to positive fluxes within a period of 1-3 days. Daily-averaged fluxes varied from -  
288  $22.5$  to  $18 \mu\text{m}^{-2} \text{s}^{-1}$ , and the coefficient of variation was 3.13. There were no clear seasonal  
289 patterns in terms of magnitude, direction, or variance, although large  $\text{CO}_2$  uptakes were recorded

290 prior to ice-on in both 2016 and 2017 and negative- and smaller positive fluxes were more  
291 common during ice-covered winter days.

292

### 293 **3.2 Comparisons among methods**

294 Differences among methods were clearly illustrated when flux data were expressed as  
295 cumulative flux (Figure 5). All methods indicated the lake was a slight CO<sub>2</sub> sink over the  
296 summer; however, estimates diverged substantially during fall. Both the *S-pCO<sub>2</sub>* and *F-pCO<sub>2</sub>*  
297 methods consistently indicated CO<sub>2</sub> flux into the lake all summer and substantial CO<sub>2</sub> flux out of  
298 the lake during fall. At the end of the year, the cumulative flux based on *F-pCO<sub>2</sub>* was 15% higher  
299 (43.4 vs 37.7 gC m<sup>-2</sup>) than flux based on *S-pCO<sub>2</sub>*, but both followed similar temporal trends. In  
300 contrast, the EC method suggested the lake fluctuated between CO<sub>2</sub> source and sink behavior  
301 with a high degree of variability on the weekly time-scale. At the end of summer (day ~268), the  
302 EC-based cumulative flux was comparable to the boundary layer-based rates. However, during  
303 fall, once mixing begins, the EC cumulative flux became progressively more negative,  
304 suggesting the lake became a more substantial CO<sub>2</sub> sink.

305 CO<sub>2</sub> fluxes based on FC (flux chamber) agreed in magnitude and direction with the *F-*  
306 *pCO<sub>2</sub>* during spring, summer, and fall (Figure 3b-d). Comparing FC with EC, the two methods  
307 disagreed in flux magnitude during summer and direction during fall.

308 The discrepancy between methods could be caused by the temporal or spatial resolution  
309 of observations. The day-time EC data more closely aligned with the *F-pCO<sub>2</sub>* and *S-pCO<sub>2</sub>* results  
310 during the summer. These methods agreed that the daytime flux of CO<sub>2</sub> during the summer was  
311 consistently into the lake. During the fall, the daytime EC fluxes remained negative, suggesting a  
312 consistent flux of CO<sub>2</sub> into the lake. Spatially, the *S-pCO<sub>2</sub>* results within the EC footprint were

313 consistent with the majority of the S-*p*CO<sub>2</sub> data. This suggests the lake was relatively  
314 homogeneous in regard to flux rates, with subset S-*p*CO<sub>2</sub> locations showing ~20% variability in  
315 accumulated fluxes at the end of the year. Temporal subsets of EC data show differences during  
316 the summer with the full-day EC data, but ultimately small differences in accumulated fluxes at  
317 the end of the year. Average EC error was 38.9%, with larger accumulated errors during the fall.

318         Directly comparing estimates using linear regression models further demonstrate the  
319 dissimilarity among methods. The two concentration gradient methods, F-*p*CO<sub>2</sub> and S-*p*CO<sub>2</sub>,  
320 agreed in magnitude and direction ( $R^2 = 0.55$ ,  $p$  value  $< 0.001$ ; Figure 6a). When flux estimates  
321 were categorized by season, data from the summer were tightly clustered, while data from the  
322 fall were more scattered. Comparing EC to S-*p*CO<sub>2</sub> (Figure 6b), there was poor agreement ( $R^2 =$   
323  $0.07$ ;  $p = 0.03$ ), and the regression model had a negative slope. Thus, daily flux rates using EC  
324 disagreed in direction with the concentration-based methods.

325         Fourier power spectral decomposition (Figure 7) of daily flux from EC, F-*p*CO<sub>2</sub> and S-  
326 *p*CO<sub>2</sub> data all had similar patterns over the quantifiable frequencies, with highest spectral power  
327 is seen in EC time-series, and S-*p*CO<sub>2</sub>, and finally F-*p*CO<sub>2</sub>. Seasonal and synoptic (3-10 day)  
328 variability dominate all three, though the EC tower also shows a sub-monthly (~20 day) mode of  
329 variability not seen in the other two.

330

#### 331 **4. Discussion**

332         Few studies have used multiple measurements of long-term lake-atmosphere fluxes to  
333 address systemic biases in methods. Using concurrent long-term records from a single lake, we  
334 showed divergent behavior among flux estimates, particularly during the fall turnover period.  
335 EC-based calculations had large and opposing sign CO<sub>2</sub> flux estimates compared to FC and

336 concentration gradient-based methods (F- $p\text{CO}_2$  and S- $p\text{CO}_2$ ). FC-based methods agreed in  
337 direction and magnitude as  $p\text{CO}_2$ -based methods, however we lack sufficient FC coverage to  
338 interrogate the validity of this agreement. Together, these results suggest that at least for this lake  
339 and these estimates, EC and concentration gradient methods for estimating  $\text{CO}_2$  flux differ  
340 dramatically.

341         The spatial and buoy-based concentration gradient estimates closely agreed. Both  
342 estimates followed similar seasonal patterns, indicating that the lake was taking in  $\text{CO}_2$  from the  
343 atmosphere during the summer, and emitted a substantial amount during the fall. The buoy-based  
344 data showed this seasonal phenology in three consecutive years [Reed *et al.*, 2018], aligning with  
345 other studies of productive lakes [S Maberly, 1996] and the perception that productive lakes  
346 behave as  $\text{CO}_2$  sinks during the summer [Balmer and Downing, 2011]. The agreement between  
347 the spatial and buoy-based concentration data suggests low spatial heterogeneity in  $\text{CO}_2$  fluxes  
348 across the surface of Lake Mendota. On average most of the lake surface was within a within a  
349  $0.2 \mu\text{mol m}^{-2} \text{s}^{-1}$  range in  $\text{CO}_2$  flux [L C Loken *et al.*, 2019]. Thus, spatial variability is small  
350 compared to the seasonal variability from all our  $\text{CO}_2$  flux methods (Figures 3 and 5). However,  
351 spatial heterogeneity increased during fall turnover, making the buoy location less representative  
352 of the whole lake during this period [L C Loken *et al.*, 2019]. During periods of chaotic water  
353 mixing, the representativeness of a single location decreases [Erkkila *et al.*, 2018]. Thus, we  
354 suspect the discrepancies among methodologies during the summer season are not due to spatial  
355 heterogeneity in gas exchange across the lake surface.

356         With a limited number of FC observations, FC data approximately matched F- $p\text{CO}_2$  and  
357 S- $p\text{CO}_2$  during the spring and summer. Comparing FC and F- $p\text{CO}_2$  methods, López Bellido *et al.*  
358 [2009] found that FC were systematically higher than F- $p\text{CO}_2$ , due to site-specific and time-

359 specific gas transfer velocities. They used daily concentration measurements and hence were not  
360 able to access daily patterns. *Podgrajsek et al.* [2014] found that FC and EC fluxes generally  
361 agreed, except when  $p\text{CO}_2$  varied within the EC footprint. This is expanded on to show higher  
362 eddy covariance  $\text{CO}_2$  fluxes at night relative to F- $p\text{CO}_2$  and also that F- $p\text{CO}_2$  methods need to  
363 account for convection within the water column [*Podgrajsek et al.*, 2016]. *Erkkila et al.* [2018]  
364 found F- $p\text{CO}_2$ -based estimates were lower than EC while those based on FC were higher than  
365 EC estimates. Together, there does not appear to be an emerging trend among results, other than  
366 EC fluxes can be typically higher at night.

367  $k$  may be responsible for the discrepancy among flux estimates. The  $k$  model underlying  
368 our concentration gradient based methods may have not adequately portrayed turbulence at the  
369 lake surface. Convective mixing within the water column introduce error into F- $p\text{CO}_2$  methods  
370 [*Podgrajsek et al.*, 2016]. These models base  $k$  on wind speed, but the effects of individual wind  
371 events on lakes are highly variable. For example, two days with similar wind speed and direction  
372 likely do not have the identical patterns of surface turbulence, compounded with spatial  
373 variability of  $k$  across fine scales [*L C Loken et al.*, 2019]. Ultimately, concentration gradient-  
374 based estimates rely on measurements that have a spatial scale on the order of one liter, and  
375 while  $k$  parameter incorporates wind speed and convection at a broader scale, our  $k$  model does  
376 not account for processes at the finer scales. While only using short periods (1-3 days), *Eugster*  
377 *et al.* [2003] used eddy covariance and chambers from Alaska and Switzerland to show the  
378 importance of convective mixing due to lake-atmosphere fluxes, with significant differences  
379 between methods during periods of stratification and with deep, penetrative convection.

380 Another possible explanation is potential biases in EC measurements during periods of  
381 low turbulence, complex turbulence, or advection. *Morin et al.* [2018] noted in a model study the

382 role of tower height and lake-land circulations in driving eddy transport that would be bias  
383 traditional flux calculation based on half-hourly Reynold's decomposition. As the surface cools,  
384 enhanced low-level atmospheric stability may suppress turbulence, leading to larger than typical  
385 storage or advective contribution to surface fluxes [Lee *et al.*, 2004]. As noted by Xu *et al.*  
386 [2019], below-sensor storage flux calculation can be critical to correcting tower-measured flux to  
387 represent surface flux, especially periods around sunrise and sunset. However, while we lack  
388 storage flux observations at this site or models of local circulation and turbulence on the  
389 peninsula, and there is no evidence in the data of a preferential circulation during fall or other  
390 periods of stable conditions. Further work on data quality filtering of EC over lakes is necessary  
391 to build confidence in its use over lakes.

392 EC may have other benefits, even when subject to potential systematic bias. Here, when  
393 examining the spectral density of the multiple observations, the EC observations show a 20-30  
394 day frequency not observed by the other methods, including the similarly high frequency buoy  
395 measurements. Eugster *et al.* [2003] also concludes that EC methods should be used in order to  
396 collect process-scale data from the full season. Similarly, Podgrajsek *et al.* [2016] suggests the  
397 high temporal resolution of EC is crucial to resolve diel changes in flux, combined with  
398 measurements within the water column with high (30 minute) frequency. Reed *et al.* [2018] used  
399 a different EC observation dataset on Lake Mendota, not used here due to a large amount of gaps  
400 from that tower's location during the study period, but showed high degrees of coherence  
401 between CO<sub>2</sub> flux and air temperature at a similar sub-monthly (20-30 day) timescale. An  
402 emerging trend in aquatic flux literature is this monthly timescale of variation where Liu *et al.*  
403 [2011]; Liu *et al.* [2016] connect synoptic weather patterns to mixing, and Shao *et al.* [2015] and

404 *Ouyang et al.* [2017] show monthly correlation between CO<sub>2</sub> flux and chlorophyll and algal  
405 blooms.

406         There are ways to capture this 20-30 day timescale without high temporal coverage.  
407 Previously *Natchimuthu et al.* [2016] used a long-term FC dataset and then sub-sampling the  
408 observations following the methods of *Wik et al.* [2016]. They concluded that only  $\geq 8$   
409 measurement days, distributed over multiple seasons, and high enough spatial coverage ( $\geq 8$   
410 locations during summer,  $\leq 5$  during spring and fall) are key for representative ( $\pm 20\%$ ) flux  
411 estimates at the annual timescale. However, they note that the flux estimates would be biased if  
412 observations excluded episodic events such as lake circulation patterns, diel or seasonal  
413 variation, or high flux areas from a lake. Given the mismatch between what the EC literature is  
414 concluding about needing high temporal resolution observations and the FC literature about only  
415 needing  $\geq 8$  days for CO<sub>2</sub> [*Natchimuthu et al.*, 2016], with a lower average flux [*Wik et al.*, 2016],  
416 we argue that while it is possible to estimate annual fluxes from a small number of sample days,  
417 functionally we think it would be difficult to observe only 8 days of FC fluxes and have a high  
418 degree of confidence that we have captured the temporal processes needed. Ultimately, we do  
419 judge the flux signal found at the 20-30 day frequency as important and the best way to capture  
420 appears to be EC methods.

421

## 422 **5. Conclusions**

423         While major advances have been made, quantifying lake-atmosphere fluxes from  
424 individual lakes over multiple spatial and temporal scales, remains a challenge. We are becoming  
425 more aware of the importance of lakes in global and local carbon cycles. Accurately accounting

426 for temporally and spatially heterogeneity in the flux of carbon across lake surfaces is vital for  
427 incorporation and constraining process-based predictions within lake models.

428 Overall, there is a need for increased spatiotemporal resolution in studies of CO<sub>2</sub>  
429 exchange between lakes and the atmosphere. Long term temporal data collection is essential to  
430 capture, diel, 20-30 day, and seasonal patterns. Spatially, there is still an open question as to  
431 which method is capturing flux magnitude correctly, as each method integrates different  
432 processes into the observation. This is done most explicitly when choosing between multiple k  
433 models but is also implicated when screening EC data. There is no emerging trend in magnitude  
434 or direction between methods and additional work is needed to bridge spatiotemporal scales.

435  
436

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443 FC 2017-18 campaigns.

444

445 Data for eddy covariance (US-PnP) can be found at Ameriflux:  
446 <http://dx.doi.org/10.17190/AMF/1433376>. CO<sub>2</sub> concentrations used in the spatial (dataset 337,  
447 doi: doi:10.6073/pasta/fe9c5437f67254f521bf5f7e0308bf93) and temporal concentration  
448 gradient (dataset ID 129, doi:10.6073/pasta/9bced2f6ff81aa30f0f573766c0a410b) can be found  
449 at the NTL-LTER database at <https://lter.limnology.wisc.edu/data> and are indexed in the  
450 Environmental Data Initiative. Floating chamber data have been deposited into the  
451 Environmental Data Initiative database at  
452 <https://doi.org/10.6073/pasta/f6a915989753aba6f18b6b095e7a52d0>.

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589

590 **Tables**

591 Table 1. Temporal duration, water/gas sampling frequency, and spatial extent and resolution for the four methods used to estimate  
 592 CO<sub>2</sub> fluxes in Lake Mendota between 2016 and 2018, along with data availability information.

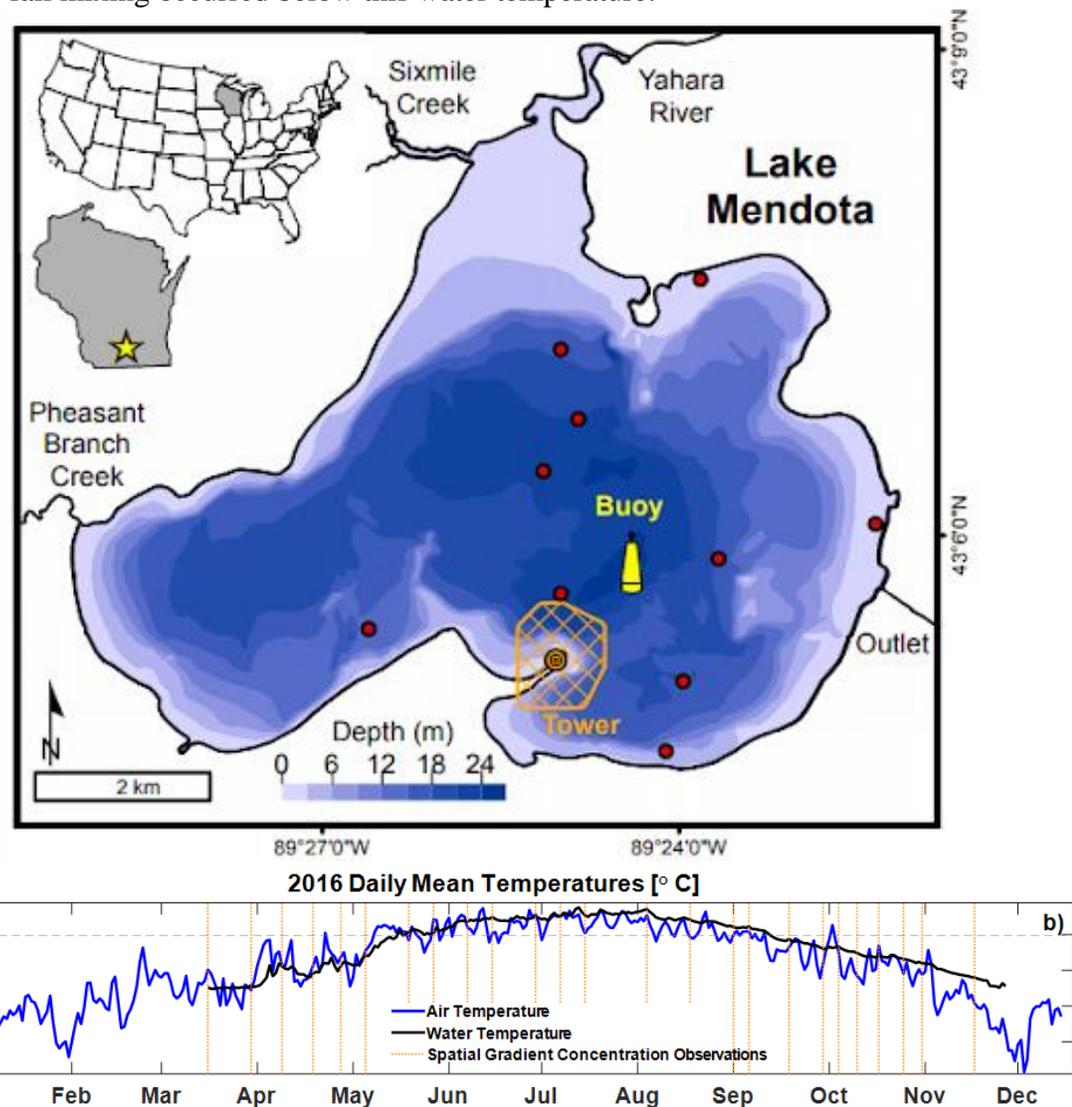
Method	Measurement period	Sampling frequency	Spatial extent	Spatial resolution	Citation
Fixed point concentration gradient (F- <i>p</i> CO <sub>2</sub> )	Open Water Seasons (approx. April-Oct) 2016	15 min	Single point	10 cm <sup>3</sup>	<i>Magnuson et al.</i> [2019]
Spatial concentration gradient (S- <i>p</i> CO <sub>2</sub> )	Mar-Dec 2016	14 d	whole lake	200 m <sup>2</sup>	<i>L Loken et al.</i> [2019]
Flux chamber diffusion (FC)	4 measurement campaigns, Jul 2017 – Apr 2018	5 min sampling, every 4 hours for 24 hours	Single point	0.28 m <sup>2</sup>	<i>A R Desai</i> [2019]
Eddy covariance (EC)	June 2016-August 2018	30 min	1 km <sup>2</sup>	1 km <sup>2</sup>	<i>A Desai</i> [2018]

593

594 **Figures**

595 Figure 1

596 Panel (a) Lake Mendota. Buoy (yellow) is deployed in the deepest part of the lake and is the  
597 location for the F-pCO<sub>2</sub> and FC flux estimates. The red circles are a stratified selection of data  
598 points from the S-pCO<sub>2</sub> method used in Figure 5. Grid section (orange with a center circle) of EC  
599 tower location and 1 km<sup>2</sup> footprint. Panel (b) 2016 average daily air (blue) and surface water  
600 (black) temperatures. Spatial gradient concentration measurements were taken on the 2016 days  
601 of year indicated by the (25 orange) vertical lines. Dashed line (gray) at 20°C used to symbolize  
602 phenology. Summer stratification is generally when surface waters were above 20°C, while  
603 spring and fall mixing occurred below this water temperature.

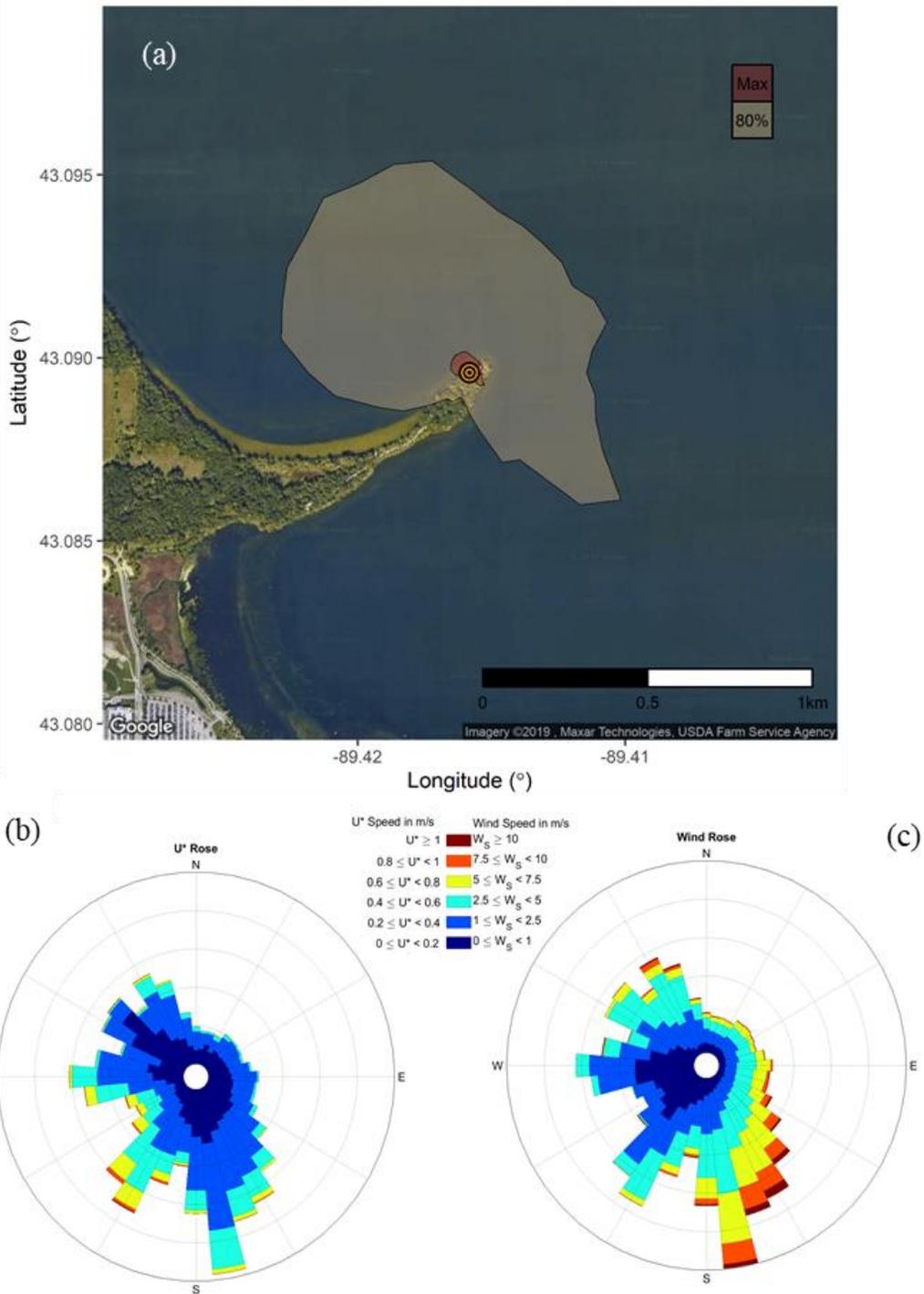


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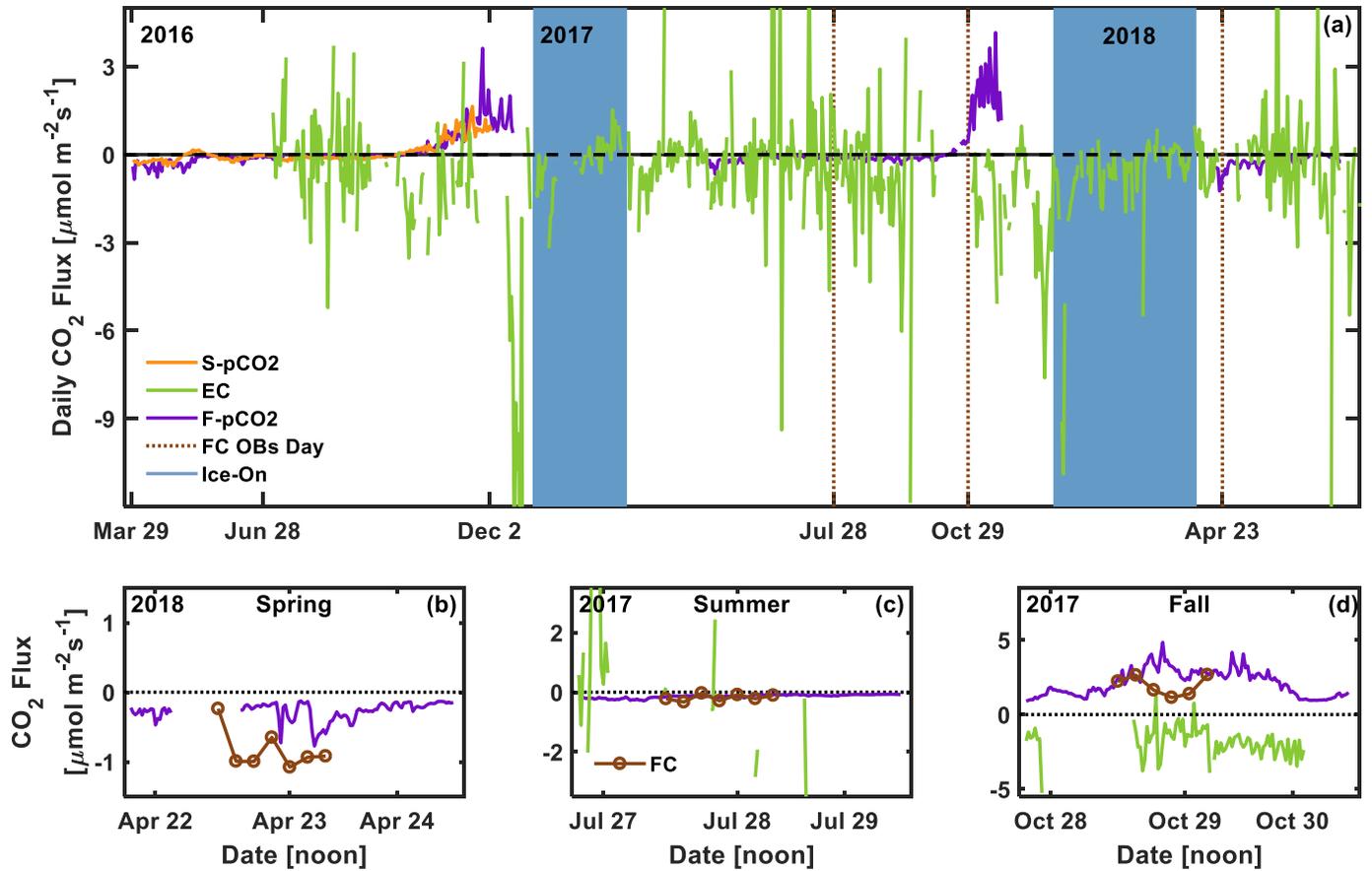
606

607 Figure 2  
 608 Panel (a) Map of picnic point EC tower and contributing footprint showing the distance of  
 609 maximum flux and distance of 80% of the footprint. Average friction velocity ( $u^*$ , panel b) and  
 610 wind speed (panel c) measured from the eddy covariance tower, shown in  $10^\circ$  bins.



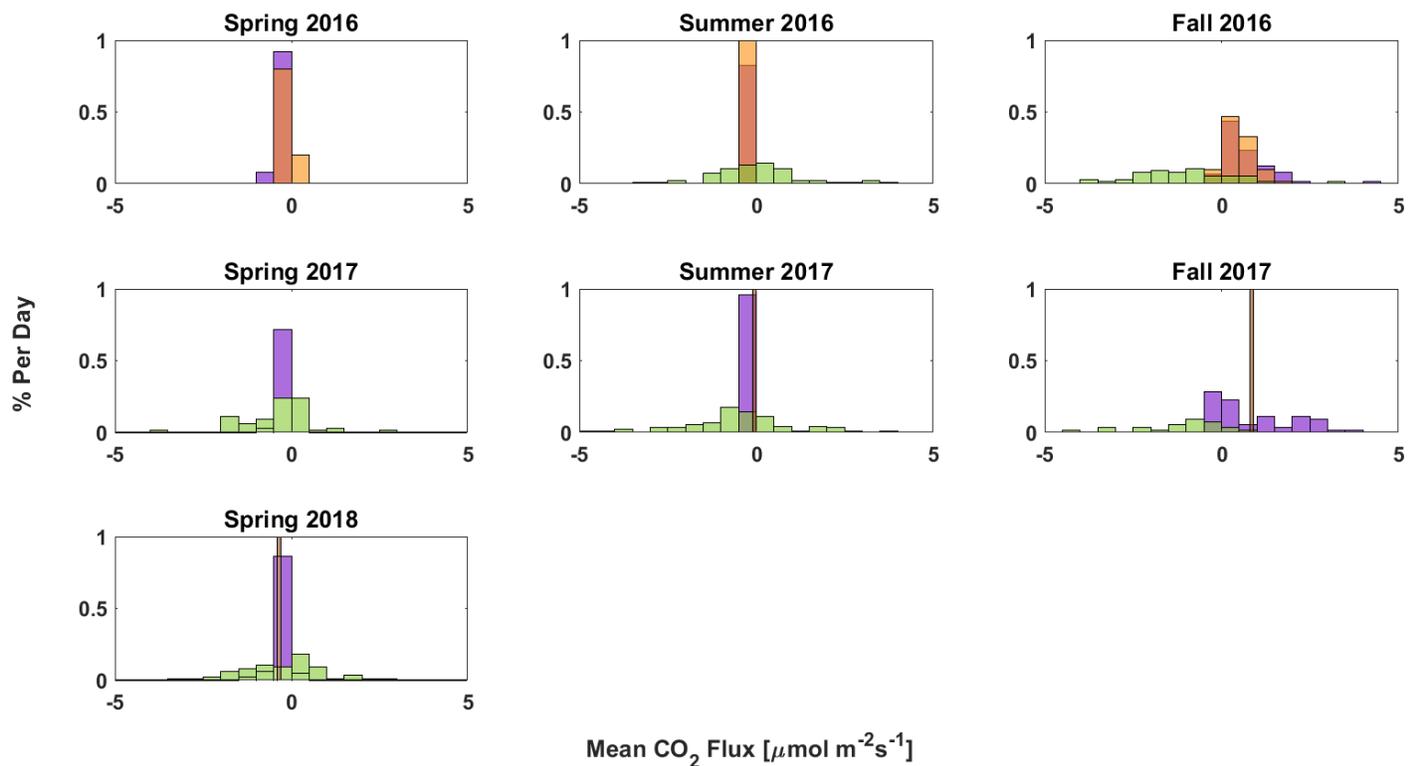
611

612 Figure 3  
 613 Panel A: Multi-year time series of mean daily CO<sub>2</sub> flux. F-*p*CO<sub>2</sub>, fixed gradient concentration  
 614 method, recorded from a stationary buoy (purple), S-*p*CO<sub>2</sub>, spatial gradient concentration  
 615 method, recorded by a moving boat (orange), and eddy covariance (green). Dates of flux  
 616 chamber measurements shown as brown dotted vertical line. Panels B-D: Hourly three day  
 617 subsets from spring, summer, and fall, centered on when FC data was collected. F-*p*CO<sub>2</sub> (purple),  
 618 and EC (green) being 30-minute data, and FC (brown) are every 4 hours for a diel cycle.  
 619



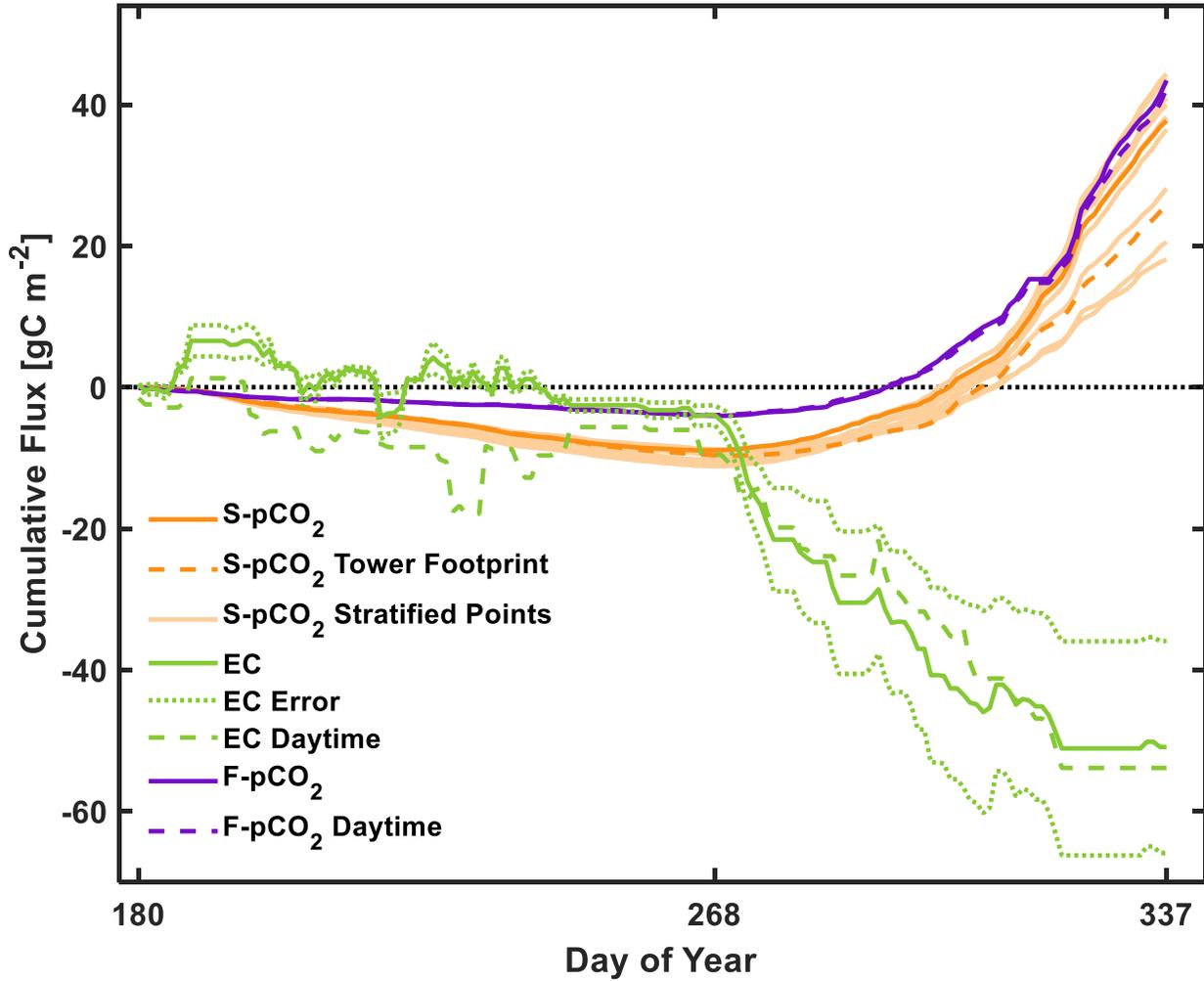
620

621 Figure 4  
622 Histograms of seasonal daily CO<sub>2</sub> gas fluxes. Spatial S-pCO<sub>2</sub> fluxes (orange), fixed F-pCO<sub>2</sub>  
623 fluxes (purple), EC fluxes (green) for spring 2016 to spring 2018 and three seasons of FC mean  
624 fluxes in 2016 (brown).  
625



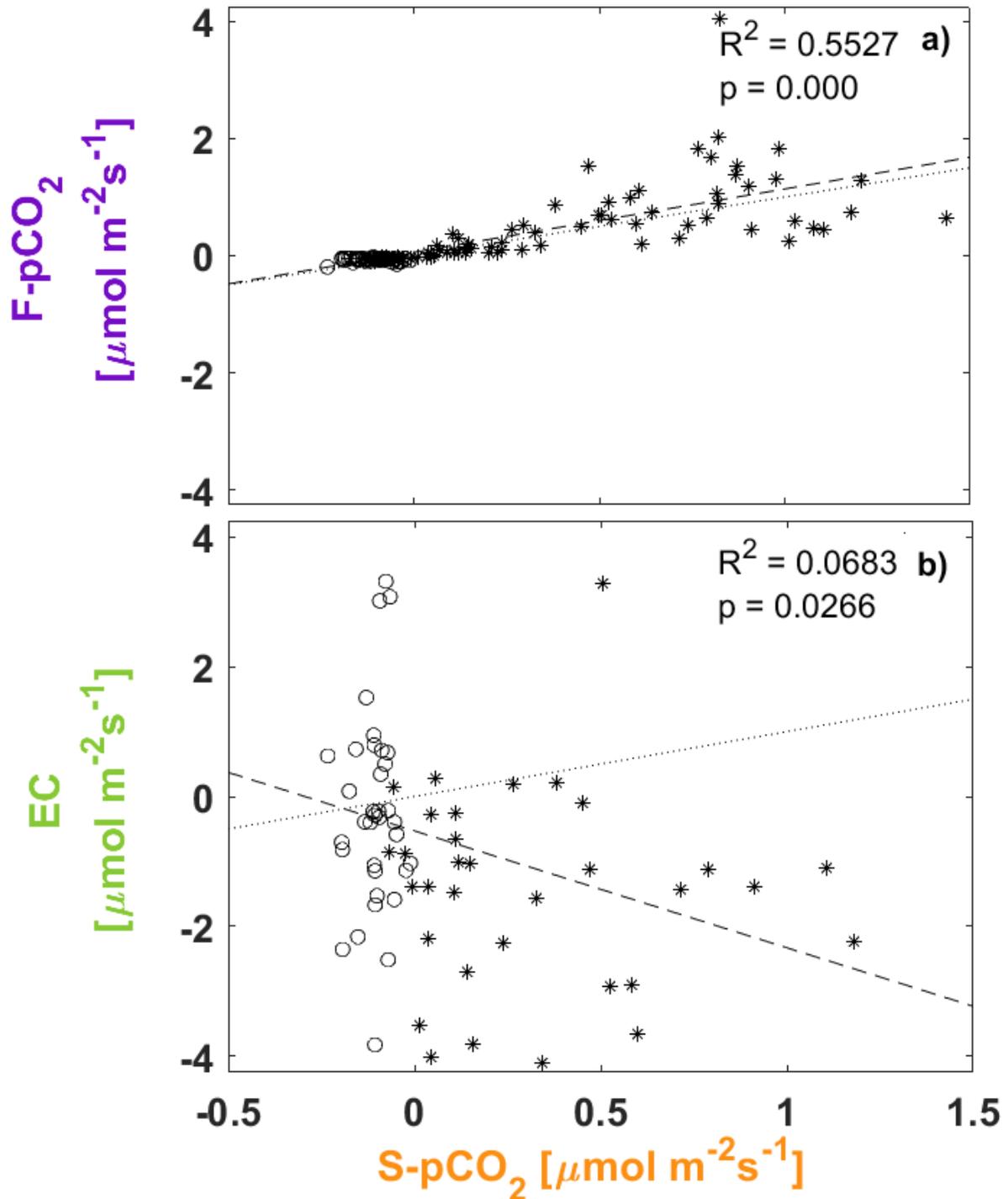
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628 Figure 5  
 629 Cumulative summation of lake-atmosphere CO<sub>2</sub> fluxes. Flux estimates using the S-pCO<sub>2</sub> method  
 630 (bold orange), from ten random points across the lake (orange), and within the tower footprint  
 631 (orange dashed line), EC (green) and EC only during day (8AM-12PM, green dashed line), and  
 632 the F-pCO<sub>2</sub> method (purple) and fixed boundary layer method during the day (8AM-12PM,  
 633 purple dashed line).



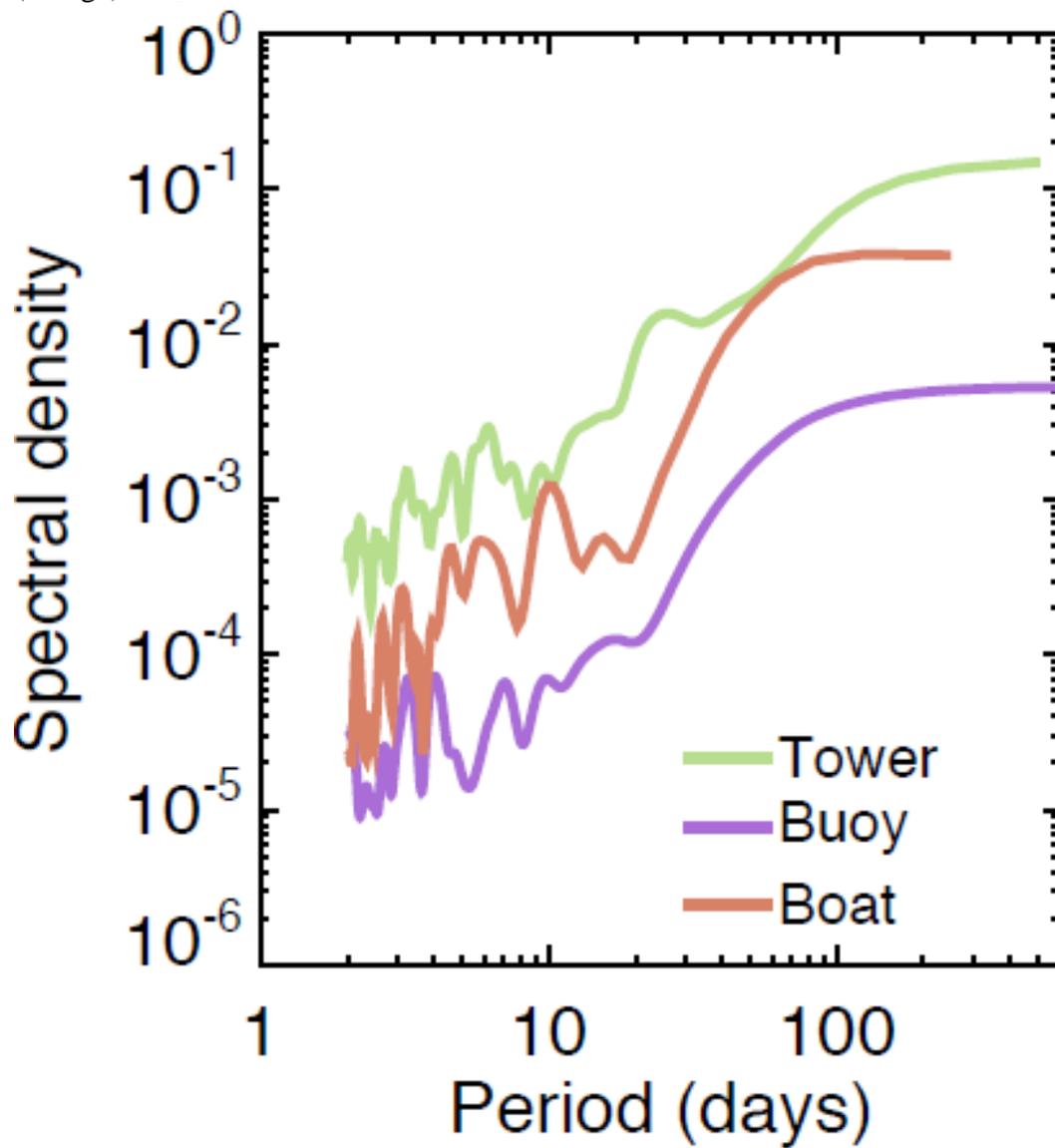
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 635

636 Figure 6 Daily mean S-pCO fluxes versus F-pCO<sub>2</sub> (panel a) and EC (panel b). Summer data are  
637 plotted as open circles, fall data as \*. Linear regression line (dashed) and one-to-one line  
638 (dotted). Statistics (*p* and R<sup>2</sup>) for linear regression included.  
639



640

641 Figure 7 Fourier power spectral decomposition of daily EC (green), F- $p\text{CO}_2$  (purple) and S- $p\text{CO}_2$   
642 (orange)  $\text{CO}_2$  flux.



643