

## 1 **Assessment of landscape-scale fluxes of carbon dioxide and methane in subtropical** 2 **coastal wetlands of South Florida**

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### 26 **Key Points:**

- 27 • Airborne eddy covariance measurements reveal heterogeneity in CH<sub>4</sub> and CO<sub>2</sub> fluxes  
28 across southern Florida.
- 29 • Variability in carbon fluxes were primarily driven by vegetation types, season, ecosystem  
30 productivity, and soil inundation.
- 31 • Southern Florida served as a net carbon sink during all flight periods, with CH<sub>4</sub> emissions  
32 offsetting CO<sub>2</sub> deposition by 11–91%.

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35

## 36 **Abstract**

37 Coastal wetlands play a significant role in the storage of 'blue carbon', indicating their  
38 importance in the carbon biogeochemistry in the coastal zone and in global climate change  
39 mitigation strategies. We present airborne eddy-covariance observations of CO<sub>2</sub> and CH<sub>4</sub> fluxes  
40 collected in southern Florida as part of the NASA BlueFlux mission during April 2022, October  
41 2022, February 2023, and April 2023. The flux data generated from this mission consists of over  
42 100 flight hours and more than 6000 km of horizontal distance over coastal saline and freshwater  
43 wetlands. We find that the spatial and temporal heterogeneity in CO<sub>2</sub> and CH<sub>4</sub> exchange is  
44 primarily influenced by season, vegetation type, ecosystem productivity, and soil inundation.  
45 The largest CO<sub>2</sub> uptake fluxes of more than 20 μmol m<sup>-2</sup> s<sup>-1</sup> were observed over mangroves  
46 during all deployments and over swamp forests during flights in April. The greatest CH<sub>4</sub> effluxes  
47 of more than 250 nmol m<sup>-2</sup> s<sup>-1</sup> were measured at the end of the wet season in October 2022 over  
48 freshwater marshes and swamp shrublands. Although the combined Everglades National Park  
49 and Big Cypress National Preserve region was a net sink for carbon, CH<sub>4</sub> emissions reduced the  
50 ecosystem carbon uptake capacity (net CO<sub>2</sub> exchange rates) by 11–91%. Average total net  
51 carbon exchange rates during the flight periods were -4 to -0.2 g CO<sub>2</sub>-eq m<sup>-2</sup> d<sup>-1</sup>. Our results  
52 highlight the importance of preserving mangrove forests and point to potential avenues of further  
53 research for greenhouse gas mitigation strategies.

## 54 **Plain Language Summary**

55 Coastal wetlands play a crucial role in trapping and storing carbon, aiding in climate change  
56 adaptation and mitigation efforts. Carbon dioxide (CO<sub>2</sub>) uptake and methane (CH<sub>4</sub>) emissions  
57 were measured from an aircraft over wetlands of southern Florida during different times of the  
58 year. Season, vegetation, ecosystem productivity, and water depth were found to have a large  
59 influence on carbon exchange. Mangroves with the largest canopy heights showed the highest  
60 CO<sub>2</sub> uptake, while CH<sub>4</sub> emissions peaked during the wet season over freshwater marshes where  
61 surface water depths were greatest. CH<sub>4</sub> emissions diminished the overall carbon uptake capacity  
62 of southern Florida. Results emphasize the importance of preserving coastal wetland ecosystems  
63 and suggest potential directions for further research aimed at mitigating greenhouse gas  
64 emissions.

## 65 **1 Introduction**

66 Vegetated ecosystems mitigate the impact of anthropogenic CO<sub>2</sub> emissions by serving as  
67 natural carbon stores (e.g., Barbier et al., 2011; Donato et al., 2011; Murdiyarso et al., 2015;  
68 Duarte 2017). The terrestrial biosphere is estimated to remove 10–40% of the CO<sub>2</sub> emitted from  
69 fossil fuels, and coastal aquatic vegetation removes a further 3–25% (Barbier et al., 2011;  
70 Donato et al., 2011; Murdiyarso et al., 2015; Duarte 2017; Friedlingstein et al., 2023). Coastal  
71 vegetated ecosystems have been of recent interest for “blue” carbon mitigation strategies because  
72 of their efficiency for short-term carbon storage in vegetation biomass (above- and belowground)  
73 and long-term carbon storage in soils and sediments (e.g., McLeod et al., 2011; Barbier et al.,  
74 2011; Donato et al., 2011; Murdiyarso et al., 2015; Macreadie et al., 2021; Poulter et al., 2023).  
75 “Blue” carbon refers to the carbon that is captured by oceans and coastal ecosystems. Although  
76 coastal vegetated ecosystems (i.e., mangroves, salt marshes, seagrasses) cover an area equivalent  
77 to ~2% of terrestrial forest area, their carbon burial rates are 50 times faster per unit area,  
78 making the net contributions of coastal wetlands to carbon sequestration comparable to that of

79 terrestrial forests (McLeod et al., 2011; Duarte et al., 2013). Yet these ecosystems are under  
80 continued threat due to hurricanes, land development, and sea level rise, which contribute to a  
81 global net loss of mangroves and salt marshes at a rate of 0.13–2% annually (McLeod et al.,  
82 2011; Goldberg et al., 2020; Campbell et al., 2022; Murray et al., 2022).

83 Understanding the climate change mitigation potential of these ecosystems requires  
84 accurate accounting of their carbon balance. Only a fraction of the CO<sub>2</sub> taken up by coastal  
85 vegetation is sequestered in sediments via long-term carbon burial. Much of this carbon is stored  
86 in shorter-term above- and belowground biomass, re-emitted to the atmosphere through soil  
87 respiration, or transported to the ocean as particulate organic carbon (POC), dissolved organic  
88 carbon (DOC), and dissolved inorganic carbon (DIC) (Rosentreter 2018b; Sanderman et al.,  
89 2018; Simard et al., 2019; Adame et al., 2021). In addition, anoxic soil conditions and  
90 methanogenic archaea in coastal vegetated ecosystems produce CH<sub>4</sub> (e.g. Bartlett et al., 1987;  
91 Rosentreter 2018c; Al-Haj and Fulweiler, 2020). Methane emissions have the potential to  
92 significantly offset the climate mitigation potential of coastal wetlands, as the global warming  
93 potential (GWP) of methane is 81.2 and 27.9 times greater than that of CO<sub>2</sub> on a 20- and 100-  
94 year scale, respectively (Forster et al., 2021). Estimates of global CH<sub>4</sub> emissions from coastal  
95 wetlands are poorly constrained, with uncertainties stemming from large regional differences,  
96 lack of direct measurements, and anthropogenic impacts on wetland disturbance and hydrology  
97 (Harrison et al., 2017; Kroeger et al., 2017; Saunois et al., 2020; Rosentreter et al., 2021).

98 Several methods exist for quantifying carbon exchange at landscape to global scales, each  
99 with their own benefits and limitations. Concentration measurements from aircraft, ground sites,  
100 and satellites can be coupled with inverse models to provide a “top-down” inference of  
101 atmosphere-biosphere CO<sub>2</sub> and CH<sub>4</sub> exchange (e.g. Wang et al., 2018; Saunois et al., 2020; Ma  
102 et al., 2021; Schiferl et al., 2022; Gaubert et al., 2023). However, these top-down approaches  
103 suffer from considerable uncertainties related to atmospheric transport and heavily rely on prior  
104 assumptions for source attribution. Satellite-based top-down approaches allow for the assessment  
105 of changes in CO<sub>2</sub> and CH<sub>4</sub> fluxes over multiple years with global coverage, which is particularly  
106 important in areas where direct ground-based and airborne measurements are limited (e.g.,  
107 Campbell et al., 2020). Satellite approaches are, however, further limited by additional  
108 uncertainties related to satellite retrievals. Bottom-up inferences of carbon atmosphere-biosphere  
109 exchange in wetlands utilize biophysical process models, inventories of biomass, and remotely  
110 sensed surface properties to indirectly calculate fluxes (Hayes et al., 2018; Saunois et al., 2020;  
111 Ma et al., 2021; Friedlingstein et al., 2022; Zhang et al., 2023). These models, however, rely on  
112 complicated parameterizations and assumptions of biological activity across a complex diversity  
113 of ecosystems and environmental conditions. This leads to large uncertainties, disagreements  
114 between different modeling approaches, and inconsistencies between top-down and bottom-up  
115 methods (Melton et al., 2013; Pandey et al., 2021; Saunois et al., 2020; Ma et al., 2021).

116 Alternatively, atmosphere-biosphere fluxes can also be measured directly on a variety of  
117 scales and can provide a more discerning understanding of wetland fluxes in space and time.  
118 Ground-based chamber measurements are important for quantifying process-level drivers of  
119 carbon exchange from soils, leaves, roots, and stems (e.g. Nahlik and Mitsch 2011; Marín-Muñiz  
120 et al., 2015; Troxler et al., 2015; Rosentreter et al., 2018a). Chamber water-atmosphere CH<sub>4</sub> and  
121 CO<sub>2</sub> fluxes coupled to measurements of water properties have identified factors controlling the  
122 cycling of carbon in mangrove-dominated Australian estuaries and tidal freshwater marshes in  
123 Veracruz, Mexico (Marín-Muñiz et al., 2015; Rosentreter et al., 2021a). These types of studies

124 are extremely useful for linking carbon fluxes to underlying processes, but measurements are  
125 typically only conducted for a short period of time at a limited number of sites, making upscaling  
126 these findings difficult and sensitive to statistical assumptions.

127 Eddy covariance flux towers provide localized representations of net ecosystem exchange  
128 (NEE) fluxes over longer periods (e.g., Barr et al., 2010; Beringer et al., 2013; Malone et al.,  
129 2015; Shoemaker et al., 2015; Alvarado-Barrientos 2020; Zhu et al., 2021). Such EC towers have  
130 been used to quantify the seasonality of net ecosystem CO<sub>2</sub> exchange in mangrove forests in the  
131 Florida Everglades (Barr et al., 2010; Barr et al., 2012), the Yucatan Peninsula (Alvarado-  
132 Barrientos 2020), and southeastern China (Zhu et al., 2021). But the degree to which  
133 measurements at one flux tower are representative of other sites varies, even within the same  
134 region, as environmental conditions (e.g. soil properties, inundation, leaf area, tidal influence,  
135 etc.) can vary from region to region and from site to site.

136 Airborne eddy covariance (EC) offers a viable approach to measure fluxes over larger  
137 areas, though with a more limited temporal resolution (e.g. Crawford et al., 1996; Sellers et al.,  
138 1997; Zulueta et al., 2013; Wolfe et al., 2015; Desjardins et al., 1982; Wolfe et al., 2018; Hannun  
139 et al., 2020). This technique has the advantage of elucidating heterogenous fluxes over a large  
140 region (15-100 km) at a relatively fine spatial scale (~1km). One recent application of the  
141 technique in a tropical Zambian wetland highlighted large discrepancies between land surface  
142 models and observations (Shaw et al., 2022). Zulueta et al. (2013) also utilized airborne EC to  
143 derive heterogenous CO<sub>2</sub> fluxes over distinct ocean, mangrove, and desert ecosystems in Baja  
144 California Sur, Mexico. This latter study also utilized tower flux measurements and vegetation  
145 indices to assess the representativeness of towers and provide a simple model for scaling up to  
146 regional CO<sub>2</sub> fluxes. However, these studies represent two of only very few that have used  
147 airborne EC to measure greenhouse gas exchange in subtropical to tropical wetlands.

148 Here we analyze extensive airborne CO<sub>2</sub> and CH<sub>4</sub> flux measurements acquired over  
149 southern Florida during the NASA BlueFlux mission. Combining flux measurements and flux  
150 footprint analysis with detailed information of land surface properties, we explore the patterns in  
151 flux variability across this diverse landscape. We also utilize long-term ground-based flux  
152 datasets to provide a valuable point of comparison and a means of upscaling to estimate the net  
153 regional carbon balance. Although the definition of “blue” carbon typically only includes tidal  
154 saltwater wetlands, we also investigate carbon exchange in the freshwater wetlands within the  
155 greater Everglades coastal watershed system. These regions are also extremely influential in the  
156 carbon cycle of this coastal zone. The primary objectives of this study are to 1) elucidate the  
157 heterogeneity of atmosphere-biosphere carbon fluxes in southern Florida, 2) identify the  
158 underlying sources of this variability, and 3) provide an estimate of the net carbon balance during  
159 the sampling periods from an atmospheric perspective. In addressing each of these objectives, we  
160 identify potential avenues for applying our unique data set to addressing “blue” carbon  
161 greenhouse mitigation strategies.

162

## 163 2 Materials and Methods

### 164 2.1 BlueFlux Field Campaign

165 BlueFlux is a NASA-sponsored effort to understand the dynamics of carbon exchange in  
166 coastal wetlands and develop long-term gridded flux estimates for science and policy  
167 applications. The BlueFlux field campaign was developed to provide comprehensive  
168 measurements of ecosystem carbon fluxes in southern Florida, with a special emphasis on  
169 mangroves. BlueFlux observations bridge multiple scales of biosphere-atmosphere exchange,  
170 including chamber measurements of soil and vegetation fluxes, ecosystem-scale fluxes from  
171 existing EC tower sites, airborne EC measurements across the south Florida region, and lateral  
172 aquatic carbon fluxes (Poulter et al., 2023). Primary study regions include Everglades National  
173 Park (ENP) and Big Cypress National Preserve (BCNP) (Fig. 1). The focus of this study is the  
174 airborne EC component of the project.

175 Southern Florida is characterized by a subtropical to tropical climate. The wet season  
176 occurs from May through October, during which conditions are hot ( $>30^{\circ}\text{C}$ ) and humid ( $>80\%$   
177 relative humidity) with frequent convective thunderstorms. The average annual rainfall is  
178 typically 1000–1700 mm, with 70% of precipitation occurring during the wet season (Florida  
179 Climate Center [https://climatecenter.fsu.edu/products-services/data/statewide-](https://climatecenter.fsu.edu/products-services/data/statewide-averages/precipitation)  
180 [averages/precipitation](https://climatecenter.fsu.edu/products-services/data/statewide-averages/precipitation)). The dry season (November to April) is typically warm ( $13\text{--}22^{\circ}\text{C}$ ) and  
181 dryer, with very rare winter frosts projected to decrease over time (Ross et al., 2009). Flights  
182 were performed during April 2022, October 2022, February 2023, and April 2023. April months  
183 are typically in the tail-end of the dry season and beginning of the wet season, while October is  
184 considered the tail-end of the wet season. Temperatures during study months were roughly  
185 average for the area and season, and ENP and BCNP experienced an average amount of rainfall  
186 during April 2022 and October 2022. However, Hurricane Ian made landfall north of the study  
187 region on September 28, 2022. There were higher observed water levels in the weeks following  
188 the hurricane at EC tower sites (EDEN, <https://sofia.usgs.gov/eden>). Conversely, February 2023  
189 and April 2023 experienced below average rainfall that was 26% and 73%, respectively, of  
190 normal levels (South Florida Water Management District [https://www.sfwmd.gov/weather-](https://www.sfwmd.gov/weather-radar/rainfall-historical/year-to-date)  
191 [radar/rainfall-historical/year-to-date](https://www.sfwmd.gov/weather-radar/rainfall-historical/year-to-date)). Atlantic basin hurricanes frequently pass over Southern  
192 Florida between August and November. Such hurricanes have resulted in significant alteration to  
193 the coastal wetlands of Southern Florida over the past 32 years (Taillie et al., 2020). The terrain  
194 is mostly flat, with some small hills (up to 6 m above mean sea level) in the northwest portion of  
195 BCNP.

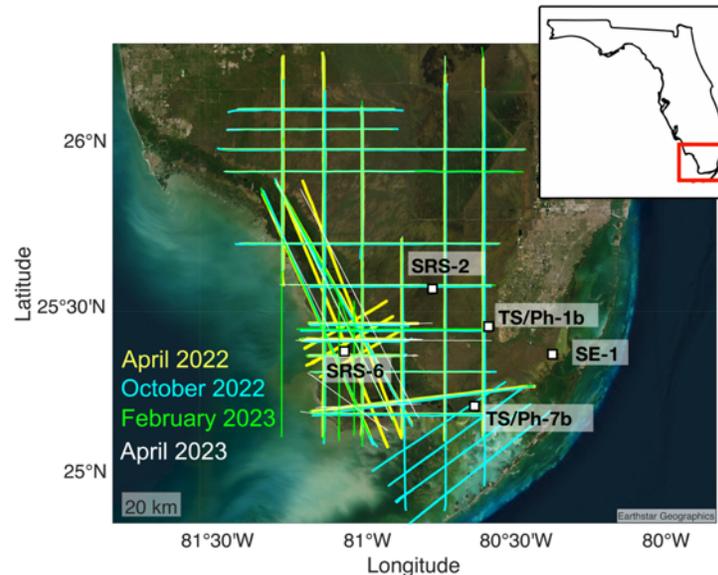
196

### 197 2.2 Airborne flux measurements

#### 198 2.2.1 Flight strategy

199 Airborne operations utilized a Beechcraft King Air A90 owned and operated by Dynamic  
200 Aviation. Deployments entailed four 2-week intensives, each consisting of 6–8 flights with  
201 durations of 2–4 hours each ( $\sim 25$  flight hours per deployment). A typical flight consisted of  
202 straight and level legs at an altitude of 90 m above mean sea level and a ground speed of 65–80

203  $\text{m s}^{-1}$ , along with occasional overlapping legs at higher altitudes (up to 300 m) to constrain flux  
 204 divergence. Vertical profiles were performed periodically (up to 3 km) to ascertain boundary  
 205 layer depth. Flux legs were typically oriented across the mean horizontal wind flow, spanned  
 206 lengths of 20–100 km, and concentrated on mangrove forests and regions of recent mangrove  
 207 dieback ('ghost' forests) (Fig. 1) that resulted from impacts of Hurricane Irma (September 2017)  
 208 (Lagomasino et al., 2021). Other considerations for flight design included overflight of existing  
 209 ground sites and avoidance of nesting bird colonies and Seminole and Miccosukee tribal lands.  
 210 In total, flux transects during all deployment periods comprise more than 6000 km of linear  
 211 distance.



**Figure 1:** Flux transects from all flights between April 2022 and April 2023. Flux legs from April 2022, October 2022, February 2023, and April 2023 are shown in yellow, cyan, green, and white, respectively. The locations of five ground sites with eddy covariance towers are indicated with square markers.

212

### 213 2.2.2 Instrument Payload

214 The BlueFlux study utilized an upgraded version of the NASA Carbon Airborne Flux  
 215 Experiment (CARAFE) platform, originally described by Wolfe et al. (2018). Ambient air was  
 216 sampled from a common inlet (1.3 cm stainless steel with fluoropel coating) located under right  
 217 wing. The sample tube was fluorinated ethylene propylene (FEP) with an inner diameter of 0.65  
 218 cm and a length of about 8 m. Gas measurements utilized two commercial Picarro closed-path  
 219 analyzers. A model g2311f (hereafter, PFlux) provided continuous measurements of  $\text{CH}_4$ ,  $\text{CO}_2$   
 220 and  $\text{H}_2\text{O}$  at 10 Hz, while a model g2401m (hereafter PConc) provided measurements of  $\text{CH}_4$ ,  
 221  $\text{CO}_2$ ,  $\text{H}_2\text{O}$ , and  $\text{CO}$  at 0.5 Hz. Agilent IDP3 scroll pumps maintained gas flows of  $\sim 5.5$  slm and  
 222  $\sim 1$  slm, respectively. The greater pressure stability of the PConc (which is designed for flight)  
 223 provides an accuracy standard, while the PFlux provided the fast measurements required for

224 eddy covariance. Supplement Section S1 describes comparisons and corrections for the two  
225 instruments. The PConc was calibrated in the lab before and after each mission (see Sect 2.2.3).

226 An Aventech Aircraft-Integrated Meteorological Measurement System (AIMMS-20)  
227 provided 20 Hz measurements of aircraft position and attitude, air temperature and pressure, and  
228 3-D wind velocities. The probe was mounted under the left wing and calibrated via the  
229 manufacturer-provided protocol at the start of each deployment.

### 230 2.2.3 Flux calculations

231 Data from the AIMMS-20 probe (20 Hz vertical wind speed,  $w$ , and potential  
232 temperature,  $\theta$ ), and the 10-Hz measurement of  $\text{H}_2\text{O}$ ,  $\text{CO}_2$ , and  $\text{CH}_4$  were time aligned to a  
233 common 10-Hz time base and combined to determine fluxes of  $\text{CH}_4$ ,  $\text{CO}_2$ , latent heat (LE) and  
234 sensible heat (H) using airborne eddy covariance with the continuous wavelet transform (CWT)  
235 method (Torrence and Compo, 1998; Wolfe et al., 2018). Flux legs were selected as flight  
236 segments greater than 15 km in linear distance with an aircraft roll not exceeding  $5^\circ$  and altitude  
237 variation within  $\pm 10$  m. Scalar time series were detrended by subtracting a 40-second ( $\sim 4$  km)  
238 moving average and time-shifted by 0 to 3 s based on lag correlation to the vertical wind (Fig.  
239 S4). This detrending length was selected to remove non-turbulent variability and maintain the  
240 largest eddies contributing to the flux (Moncrieff et al., 2006).

241 Following this pre-processing of the data, fluxes were calculated using CWT (Section  
242 S2). Flux data for analysis is filtered by the cone of influence (COI), i.e., the spectral region  
243 where additional errors and uncertainties may be present due to edge effects (Torrence and  
244 Compo, 1998). Data are excluded where the fraction of the cospectral power that resides within  
245 the COI is greater than 0.5. Fluxes are further filtered to exclude measurements where the  
246 friction velocity ( $u^*$ )—as determined from momentum fluxes at aircraft height (Section 2.4.3)—  
247 is less than  $0.2 \text{ m s}^{-1}$  over land, or less than  $0.1 \text{ m s}^{-1}$  over water. This criterion was selected to  
248 exclude periods with insufficient vertical mixing (e.g. Hogstrom 1988; Barr et al., 2010; Hayek  
249 et al., 2018). The selected  $u^*$  filtering limits are in accordance with EC towers (Barr et al., 2010)  
250 and were verified for flight data with the method of Hayek et al., 2018.

### 251 2.2.4 Uncertainties

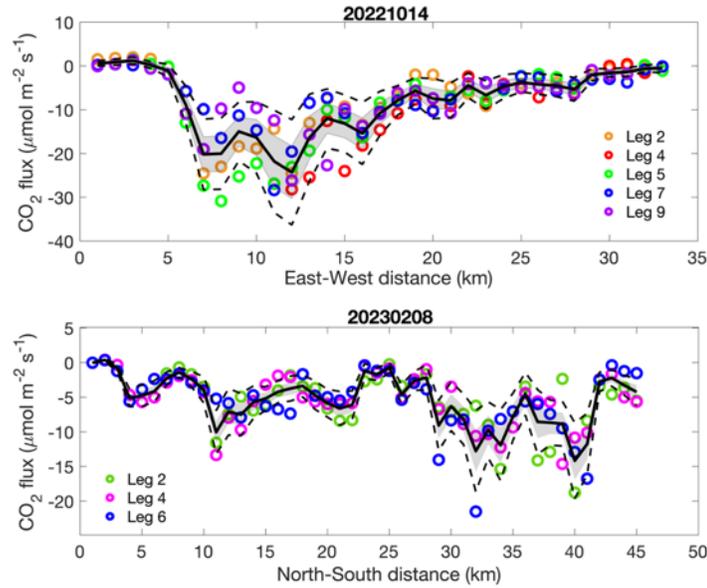
252 Systematic error contributions to flux uncertainties include those due to under-sampling  
253 of low frequencies ( $\text{SE}_{\text{turb}}$ ), the instrument response time which can limit detection of high-  
254 frequency signals ( $\text{SE}_{\text{RT}}$ ), and instrument accuracy ( $\text{SE}_{\text{acc}}$ ).  $\text{SE}_{\text{RT}}$  and  $\text{SE}_{\text{turb}}$  were calculated  
255 according to Wolfe et al., 2018. The e-folding response time used to calculate  $\text{SE}_{\text{RT}}$  was  
256 determined through laboratory tests to be  $90 \pm 10$  ms for the PFlux instrument—which translates  
257 to an effective cutoff frequency of 3.8 Hz.  $\text{SE}_{\text{acc}}$  for each scalar is based on measurement  
258 accuracy. Accuracy for  $\text{CH}_4$  and  $\text{CO}_2$  measurements are 0.05%, and 0.2%, respectively,  
259 determined through laboratory calibration with WMO-grade calibration NOAA cylinders (IDs  
260 CC746186 and CA03516). The PFlux stated  $\text{H}_2\text{O}$  accuracy is 0.8%. The AIMMS-20 probe has a  
261 stated accuracy of 10% for vertical wind speed.

262 Random errors in fluxes include contributions from uncorrelated instrument noise and  
263 turbulent variability. Methods have been developed for traditional ensemble-averaged EC to

264 represent the individual contributions of these two sources of uncertainty, as well as to  
 265 empirically calculate the total random error based on the cross- and auto- covariance of scalar  $s$   
 266 and  $w$  at different time lags (e.g. Leschow et al., 1994, Finkelstein and Simms, 2001). It is not  
 267 immediately obvious how these approaches are best applied to time-resolved CWT analysis.  
 268 Here we propose a new method for quantifying random flux errors (RE) for CWT based upon  
 269 Langford et al. (2015). In this approach, the root mean squared deviation from zero of the cross-  
 270 covariance between  $s$  and  $w$  is used to represent the random flux error (RE):

$$271 \quad RE = \sqrt{N} \sqrt{\left(0.5 \left( (\sigma_{w's'[-\Gamma]})^2 + (\overline{f_{w's'[-\Gamma]}})^2 + (\sigma_{f_{w's'[\Gamma]}})^2 + (\overline{f_{w's'[\Gamma]}})^2 \right)\right)^2} \quad (1)$$

272 Here  $N$  is the number of data points per second,  $\sigma_{w's'}$  is the standard deviation of the covariance  
 273 ( $f_{w's'}$ ) and  $\overline{f_{w's'}}$  is the average cross-covariance over a time lag range of  $-\Gamma$  or  $+\Gamma$ . Primes



**Figure 2:** CO<sub>2</sub> fluxes for repeated legs over the same flight path on October, 14, 2022 (top) and February 8, 2023 (bottom). Colored markers represent 1 km average fluxes for the different legs. Solid black lines, shading, and dotted lines represent the mean flux, 1 $\sigma$  random error, and 2 $\sigma$  random error, respectively.

274 denote deviations from the mean of  $w$  and  $s$ . We define  $\Gamma$  over a time lag range from one to 100  
 275 data points. Here, 100 was chosen as the maximum lag considered for  $\Gamma$  to be representative of  
 276 the integral time scale. This representation considers the variability in the cross-covariance of  $s$   
 277 and  $w$ , as well as the offset from zero related to non-turbulent trends in the data.

278 Random flux errors vary along flux legs due to variations in turbulence and tend to be  
 279 larger in magnitude for larger magnitude fluxes. For 1-Hz-averaged flux measurements, the  
 280 median limit of detection (LOD), defined as twice the median random flux error, is 2.8  $\mu\text{mol m}^{-2}$   
 281  $\text{s}^{-1}$  and 18.3  $\text{nmol m}^{-2} \text{s}^{-1}$  for CO<sub>2</sub> and CH<sub>4</sub> fluxes, respectively. Average fluxes at 1 km resolution  
 282 have a median LOD of 0.9  $\mu\text{mol m}^{-2} \text{s}^{-1}$  and 5.8  $\text{nmol m}^{-2} \text{s}^{-1}$  for CO<sub>2</sub> and CH<sub>4</sub> fluxes,

283 respectively. This error is approximately a factor of three lower than that derived using the  
284 approach of Wolfe et al. (2018), which was calculated from the sum (rather than the standard  
285 deviation and mean) of the cross covariance between  $s$  and  $w$  in analogy to Finkelstein and Sims  
286 (2001). Random fluxes were also estimated experimentally over a leg on April 19, 2022, by  
287 overflowing the inlet with calibration gas. The standard deviation of the 10-Hz CO<sub>2</sub> “flux” was  
288  $0.7 \mu\text{mol m}^{-2} \text{s}^{-1}$  (mean  $-0.0045 \mu\text{mol m}^{-2} \text{s}^{-1}$ ), providing an estimation of combined instrument  
289 noise and turbulence random errors over this leg. We found the approach of Wolfe et al. (2018)  
290 to result in an unreasonably large random flux error over the same flight track of  $9.0 \mu\text{mol m}^{-2} \text{s}^{-1}$   
291 <sup>1</sup>, compared with our new parameterization ( $3.3 \mu\text{mol m}^{-2} \text{s}^{-1}$ ) (Figure S11).

292 During each flight we performed repeat flux legs over the same ground track above a  
293 mangrove forest to provide an additional test of the variability in flux measurements due to  
294 random error. Figure 2 shows 1km-averaged CO<sub>2</sub> fluxes from repeat legs during two different  
295 flight patterns flown in October 2022 and February 2023. Leg-to-leg variability typically falls  
296 within that expected based on random errors estimated via Eqn. (1), providing further validation  
297 of the calculated random flux errors.

298 We include an additional uncertainty estimate from the vertical divergence of fluxes (see  
299 Supplementary Information Section S5). During each flight we performed vertically stacked legs  
300 to estimate the change in flux with altitude and allow for extrapolation of fluxes to the surface. In  
301 most cases the differences in calculated surface fluxes and fluxes measured at the aircraft altitude  
302 ( $< 100$  m) were not statistically significant ( $\alpha = 0.05$ , two-sided t-test). This is not surprising, as  
303 the aircraft altitude was typically in the lowest 10 % of the boundary layer. We therefore assume  
304 that the surface fluxes are equal to the aircraft altitude fluxes and include the difference between  
305 the extrapolated surface flux and flux measured at the aircraft altitude as an additional systematic  
306 error. We do not correct the reported fluxes for the calculated surface flux divergence because  
307 this correction uncertainty is typically much larger in magnitude than the correction itself, which  
308 would thus introduce even greater uncertainty. The magnitudes of all contributing flux errors are  
309 shown in Figure S12. The largest sources of systematic uncertainty are divergence effects (IQR  
310 3-30%) and  $SE_{acc}$  (10%). The effect of RE is small when averaged over a flux leg (IQR 1–5%  
311 uncertainty), but large for 1-Hz fluxes (IQR 30–60 % uncertainty)

### 312 2.3 Flux towers

313 Several flux towers located in the Everglades regions of southern Florida measure half-  
314 hourly fluxes of CO<sub>2</sub>, CH<sub>4</sub>, sensible heat (H), and latent energy (LE) using the eddy covariance  
315 method. These towers are part of the Florida Coastal Everglades Long-Term Ecological  
316 Research (FCE LTER) and the AmeriFlux tower networks. Towers are located along the Shark  
317 River Slough (SRS) and the Taylor Slough/Panhandle (TS/Ph) (Fig. 1) hydrologic gradients  
318 (Barr et al., 2010; Malone et al., 2015) and are representative of freshwater marsh (SRS-2),  
319 freshwater marsh prairies (TS/Ph-1), mangrove forests (SRS-6) and mangrove scrub (TS/Ph-7).  
320 These EC towers measure vertical wind speed and virtual temperature with 3D sonic  
321 anemometers (SRS-6: model RS-50, Gill Co., Lymington, England; SRS-2, TS/Ph-7, and TS/Ph-

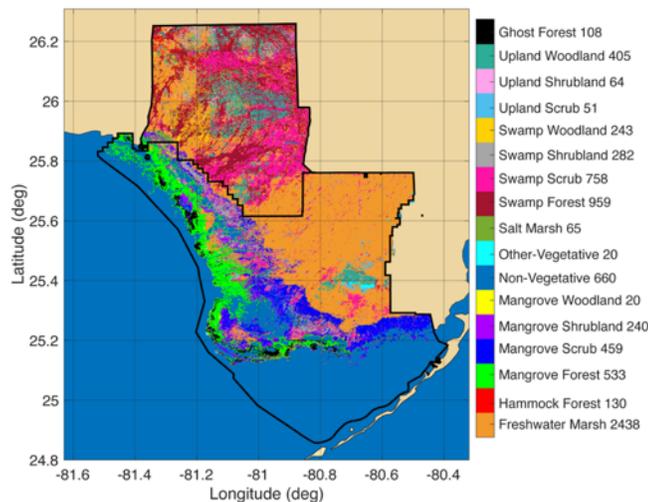
322 1: CSAT 3B, Campbell Scientific Inc., Logan, Utah). CO<sub>2</sub>/H<sub>2</sub>O (LI-7500) and CH<sub>4</sub> (LI-7700) are  
 323 measured at 20 Hz with open path infrared gas analyzers (LI-COR, Inc., Lincoln, Nebraska).

## 324 2.4 Flux decomposition by land classification

325 Southern Florida is a heterogenous landscape with a wide range of vegetation types.  
 326 Vegetation phenology, quantity, and productivity modulate CO<sub>2</sub> uptake. Other features like  
 327 salinity, water levels, surface water extent, tides, inundation period, and soil moisture can lead to  
 328 changes in CH<sub>4</sub> emission fluxes and biological CO<sub>2</sub> respiration. We consider several  
 329 geographical data sets to identify some of the causes of the observed variability in GHG fluxes  
 330 across the flight domain.

### 331 2.4.1 Vegetation coverage

332 Land cover and vegetation information for ENP and BCNP was obtained from Ruiz et al.  
 333 (2019, 2021) and Whelan et al. (2020). Land classifications based on these data sets included 16  
 334 different classes (Fig. 3) at 50 m spatial resolution. Dominant land classifications sampled during  
 335 BlueFlux over ENP and BCNP were freshwater marsh (21%), mangrove forest (17%), mangrove  
 336 scrub (10%), mangrove shrubland (6.6%), salt marsh (2.2%), swamp forest (6.9%), swamp scrub  
 337 (7.5%), swamp shrubland (5.7%), and upland forest (3.5%). The “Ghost Forest” land class was  
 338 added to identify where mangrove forests experienced extensive die-offs by drowning following  
 339 Hurricane Irma in 2017 (Lagomasino et al., 2021). Ghost forests constituted 2.2% of all land  
 340 classes sampled, and 13% of the mangrove forest sampled. Dominant vegetation species found in  
 341 each class are listed in Supplementary Information Section S3. Some flight tracks are outside of  
 342 the ENP and BCNP boundaries and are therefore not included in the vegetation analysis.



**Figure 3:** Map of vegetation coverage for the Everglades National Park (ENP) and Big Cypress National Preserve (BCNP) regions. Figure adapted from Ruiz et al. (2019, 2021) and Whelan et al., (2020). The ghost forest area was adapted from Lagomasino et al. (2021). Numbers following the vegetation types in the figure legend denote the area of each region in

units of km<sup>2</sup>. Black solid lines denote the boundaries of ENP in the south and BCNP in the north.

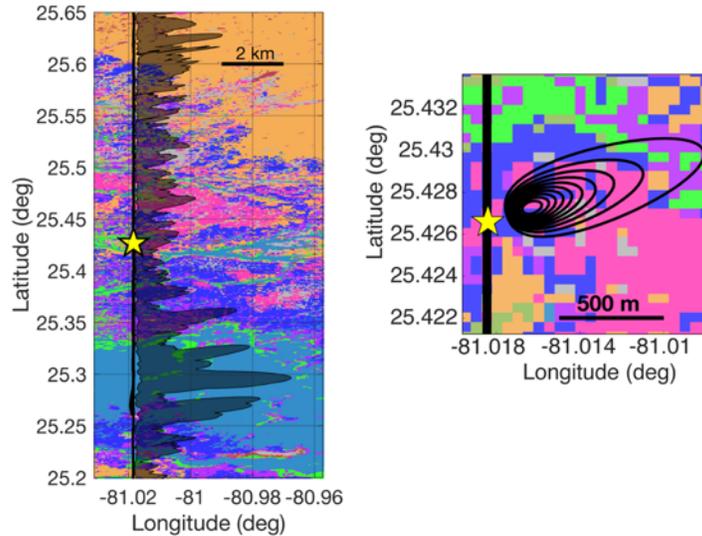
#### 343 2.4.2 Other surface characteristics

344 Remotely sensed satellite products of enhanced vegetation index (EVI), normalized  
345 difference vegetation index (NDVI), leaf area index (LAI), the fraction of photosynthetically  
346 active radiation (400-700 nm) absorbed by green vegetation (FPAR), and soil moisture were  
347 obtained over South Florida as an average for each flight month. The vegetation indices were  
348 obtained from the Moderate Resolution Imaging Spectroradiometer (MODIS) on the Terra  
349 satellite at a resolution of 250 m (Didan et al., 2015). LAI and FPAR were acquired from the  
350 combined MODIS Terra + Aqua land data products at 500 m resolution (Myneni et al., 2015).  
351 The Soil Moisture Active Passive mission (SMAP) provided a remotely sensed soil moisture  
352 product at 9 km resolution (O'Neil et al., 2023). Above ground biomass density (AGBM) and  
353 canopy heights for the study region were estimated from the Global Ecosystem Dynamics  
354 Investigation (GEDI) Lidar 2021 data products at 1 km resolution (Dubayah et al., 2021, 2023).

355 The Everglades Depth Estimation Network (EDEN, <https://sofia.usgs.gov/eden>) provides a  
356 long-term daily estimate of surface water-level. This data set consists of surface water depth  
357 estimates at 400 m resolution obtained from a model that interpolates measurements from a

358 dense network of water gauges through the Everglades and water management areas of South  
 359 Florida (Haider et al., 2020).

### 360 2.4.3 Footprint analysis



**Figure 4:** (Left) a single flux transect at 90 m altitude from the flight on April 19, 2023 superimposed on the map of vegetation cover. The shaded area shows the cumulative footprint containing the area contributing to 90% of the flux signal. (Right) A single footprint at the measurement location marked with the star. Contours depict the weighted contributions to the observed flux from 10% to 90% in 10% increments. Background colors denote land classification (Fig. 2).

361 Along each flux segment below 100 m altitude we computed a 2D flux footprint that  
 362 expresses the relative contribution of each upwind surface element to the observed flux. This  
 363 footprint analysis is detailed in Hannun et al. (2020). Briefly, to compute the flux footprint we  
 364 used the parameterization of Kljun et al. (2015) based on a Lagrangian stochastic particle  
 365 dispersion model (Kljun et al., 2002). Inputs to the 2D footprint calculation include the  
 366 measurement height, mean horizontal wind speed ( $U$ ), planetary boundary layer (PBL) height,  
 367 the Obukhov length ( $L_{OB}$ ), standard deviation of the lateral velocity fluctuations ( $\sigma_v$ ), and the  
 368 friction velocity ( $u^*$ ). For these calculations we used the PBL height obtained from the High-  
 369 Resolution Rapid Refresh (HRRR) 3 km product interpolated along our flight track. HRRR PBL  
 370 height was validated against single-point determinations of actual PBL height based on  
 371 observations of trace gas vertical profiles during each flight. We calculated  $u^*$  from the  
 372 momentum fluxes of the horizontal winds  $u$  and  $v$  (also determined with the CWT) and validated  
 373  $u^*$  with measurements from the Everglades network of EC towers (Fig. S13 and S14). The flux  
 374 footprint was then rotated into the mean wind direction and translated to geographical  
 375 coordinates. Example flux footprints along a flight track are shown in Figure 4. For segments  
 376 below 100 m altitude, 90 % of the flux signal is contained within a region 1–2 km up wind of the

377 measurement location over land. Footprints are typically larger over water, with 90% of the flux  
378 signal contained within 5 km of the measurement point.

#### 379 2.4.4 Flux disaggregation

380 The observed flux contains contributions from the fluxes of different land classes  
381 contained within the flux footprint. To derive the mean flux for each vegetation type over a set of  
382 flux observations (e.g., flux leg, single flight, or period of deployment), we used the method  
383 described in Hannun et al. (2020). This method utilizes the Disaggregation combining Footprint  
384 analysis and Multivariate Regression (DFMR) technique of Hutjes et al. (2010). Here the  
385 observed flux is treated as a linear combination of component fluxes from each land class within  
386 the footprint, such that:

$$387 \quad F_{obs} = \sum_{k=1}^n C_k F_k \quad (6)$$

388  $C_k$  is the fractional contribution of the  $k^{\text{th}}$  land class to the flux footprint, and  $F_k$  is the average  
389 flux from the corresponding land class over the observation period.  $C_k$  for each 1 Hz flux  
390 observation was determined by overlaying the footprint function onto a gridded map of land  
391 cover (Fig 3, Fig 4) and weighting by the contribution of each grid cell to the observed flux  
392 (areas closer to the measurement point contribute more heavily to the measurement).  $F_k$  was  
393 calculated via multilinear regression of  $F_{obs}$  versus  $C_k$  for each land class that constituted more  
394 than 25% of the flux footprint over more than 10 linear km of cumulative (but not necessarily  
395 consecutive) observations. This criterion was selected to ensure sufficient sampling of each land  
396 class during the observation period. The regions that met this criterion were mangrove forests,  
397 mangrove scrub, mangrove shrubland, ghost forest, salt marsh, freshwater marsh, swamp shrub,  
398 swamp scrub, swamp forest, and upland forest.

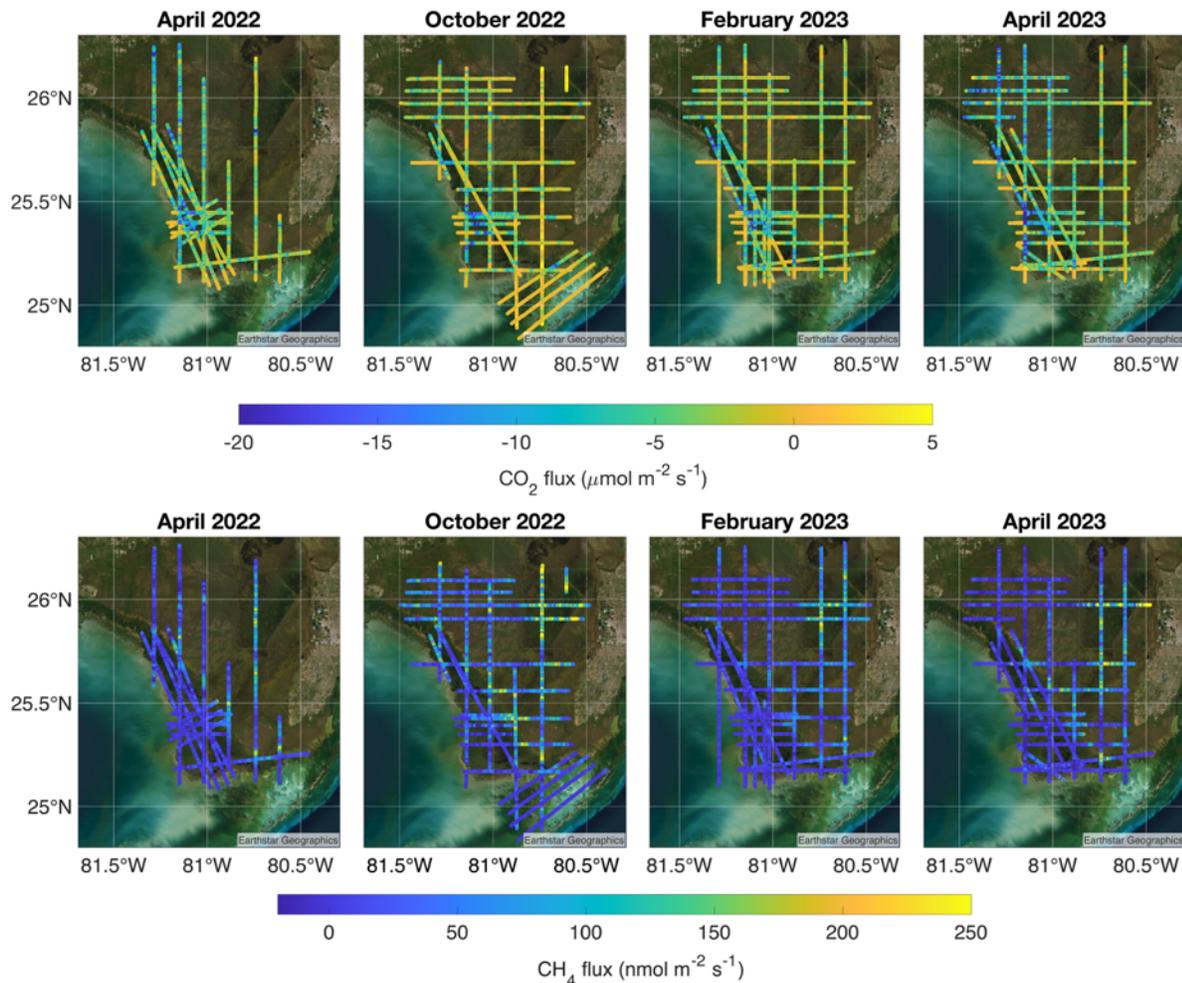
399 Uncertainty in mean fluxes for each land class was calculated as the statistical uncertainty  
400 in the regression. Random and systematic errors, as well as the calculated divergence correction  
401 (summed in quadrature to yield the total uncertainty) for each flux observation were also  
402 propagated through the regression analysis. We do not include uncertainties in the land surface  
403 data or the footprint analysis, as we expect these to be comparatively small (Hannun et al., 2020).

404 Disaggregation of fluxes from additional categorical land data were computed using the  
405 same method as for vegetation data. Continuous numerical land cover data, such as NDVI, do  
406 not require multivariate regression to disaggregate fluxes. After superimposing the footprint  
407 function onto the geographical data set, the footprint weighted average of the land cover data can

408 simply be calculated at each 1-Hz observation in the same manner as for fractional land class  
 409 contributions (e.g.,  $C_k$ ).

### 410 3 Results and Discussion

#### 411 3.1 Heterogenous CWT-derived Fluxes



**Figure 5:** Spatial distribution of (top)  $\text{CO}_2$  and (bottom)  $\text{CH}_4$  1 km averaged fluxes over flight legs below 100 m altitude for all seasons. Here negative fluxes denote uptake and positive fluxes represent emission. Flux rates are color coded to scale below maps. Larger  $\text{CO}_2$  uptake fluxes appear as darker colors, while larger  $\text{CH}_4$  emissions appear as brighter colors

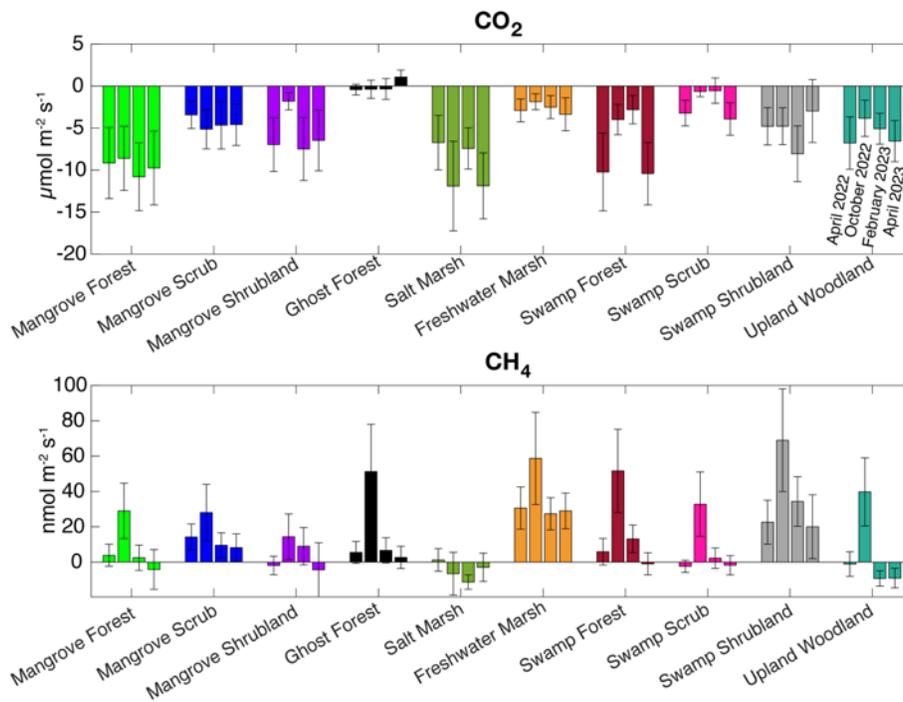
412 The airborne CWT derived fluxes measured below 100 m flight altitude are shown in  
 413 Figure 5. Negative and positive fluxes represent uptake and emission by the surface,  
 414 respectively. These fluxes provide a measurement of the net ecosystem exchange of  $\text{CO}_2$  and  
 415  $\text{CH}_4$ , as the aircraft samples the net canopy exchange (photosynthetic uptake, respiration, and  
 416 storage) of carbon. Downward (i.e., uptake)  $\text{CO}_2$  fluxes are largest (less than  $-15 \mu\text{mol m}^{-2} \text{s}^{-1}$ )

417 during all flight periods over mangrove forests in the southwest portion of the flight domain, near  
 418 25°30' N and 81° W. During both April deployments, high rates of CO<sub>2</sub> uptake were also  
 419 observed in the northwest quadrant over the swamp and upland forests of BCNP.

420 Methane fluxes also demonstrate significant spatial heterogeneity during all deployment  
 421 periods. The largest methane fluxes (greater than 200 nmol m<sup>-2</sup> s<sup>-1</sup>) occur in the northeast portion  
 422 of the flight domain over freshwater marshes. High CH<sub>4</sub> emissions also appear in a band just  
 423 inland of the west coast in the transition region between mangroves and marshlands (Fig 3, Fig  
 424 5).

### 425 3.2 Drivers of CO<sub>2</sub> uptake and CH<sub>4</sub> emission

#### 426 3.2.1 Vegetation class



**Figure 6:** CO<sub>2</sub> and CH<sub>4</sub> fluxes disaggregated by land class for all flights during April 2022, October 2022, February 2023, and April 2023. Fluxes are grouped by vegetation coverage. Error bars represent  $\pm 1\sigma$  uncertainty in the component flux, which includes systematic, random, and divergence flux uncertainties propagated through the regression analysis, in addition to the regression residuals, which also reflect the variability in carbon fluxes for each vegetation regime.

427 Figure 6 shows that CO<sub>2</sub> and CH<sub>4</sub> fluxes clearly and consistently vary according to the  
 428 underlying vegetation type. In some cases, there is also an apparent seasonality in the fluxes.  
 429 Disaggregated fluxes demonstrate that the largest fluxes of CO<sub>2</sub> uptake were observed for  
 430 mangrove forests, salt marshes and swamp forests. The greatest CH<sub>4</sub> emissions were consistently

431 measured over freshwater marshes and swamp shrublands. Below we discuss in detail the  
432 patterns in observed carbon exchange for mangrove, salt marsh, freshwater wetlands, and upland  
433 ecosystems.

434

#### 435 *Mangroves*

436 Mangrove forest CO<sub>2</sub> fluxes did not exhibit much variation over the different sampling  
437 months, with the largest average uptake rate of  $-11 \pm 4 \mu\text{mol m}^{-2} \text{s}^{-1}$  observed during February  
438 2023. These fluxes are consistent with the range of  $-5$  to  $-18 \mu\text{mol m}^{-2} \text{s}^{-1}$  uptake observed in the  
439 Yucatan during 2017–2018 from an eddy-flux tower (Alvarado-Barrientos et al., 2021). Large  
440 peak daily fluxes of  $-13$  to  $-20 \mu\text{mol m}^{-2} \text{s}^{-1}$  and  $-17.1$  to  $-19.9 \mu\text{mol m}^{-2} \text{s}^{-1}$  have been observed in  
441 coastal southeastern China (Zhu et al., 2021) and near Hong Kong (Liu and Lai 2019),  
442 respectively. Weak seasonality of CO<sub>2</sub> fluxes was also observed in both other regions. We  
443 regularly observed fluxes of  $-17$  to  $-22 \mu\text{mol m}^{-2} \text{s}^{-1}$  near the SRS-6 tower site in southwestern  
444 ENP, where previous studies have reported the highest mangrove above-ground biomass and  
445 productivity (Castañeda-Moya et al., 2013; Danielson et al., 2017; Rivera-Monroy et al., 2019)  
446 (Fig. 5, Fig S17, Fig S18). Barr et al. (2010) reported peak uptake values between  $-15$  and  $-25$   
447  $\mu\text{mol m}^{-2} \text{s}^{-1}$  at the SRS-6 tower site prior to Hurricane Wilma in 2005, suggesting the strong  
448 carbon sink capacity of these riverine mangroves. Our smaller average midday CO<sub>2</sub> fluxes  
449 compared with other regions of mangroves may in part be due to lasting effects of past  
450 hurricanes. Hurricane Andrew in 1992 had devastating impacts on mangrove forests (Smith et  
451 al., 2005). Although trees recovered, canopy heights are still lower in some areas than pre-  
452 Andrew levels. High winds and large storm surges from Hurricane Irma in 2017 also created  
453 ghost forests and caused additional canopy height loss (Lagomasino et al., 2021).

454 Average mangrove shrub CO<sub>2</sub> fluxes were very similar to the  $-8.11 \mu\text{mol m}^{-2} \text{s}^{-1}$  fluxes  
455 measured for mangroves of similar heights and speciation in Baja California Sur, Mexico  
456 (Zulueta et al., 2013). Lesser average uptake of  $-3$  to  $-5 \mu\text{mol m}^{-2} \text{s}^{-1}$  was measured for mangrove  
457 scrub systems, which have a lower average canopy height ( $< 2$  m) compared to mangrove  
458 shrublands (2–5 m) and forests ( $> 5$  m). The mangrove shrublands exhibit greatly reduced CO<sub>2</sub>  
459 uptake during October 2022. The cause of this reduction in productivity is unclear but may be  
460 due to either increased freshwater inundation during the wet season, or differences in flight paths  
461 and conditions that led to more limited sampling of mangrove shrublands during October 2022.  
462 If the first explanation was the cause, we would likely have also seen an effect in mangrove  
463 scrub fluxes. Mangrove shrublands contributed half as much to the total cumulative flux  
464 footprints during October 2022 as the other deployment periods, making it difficult to rule out  
465 sampling bias. A single flight on October 17, 2022, comprised half of the sampled mangrove  
466 shrublands (Table S1). This particular flight resulted in a uniquely low estimate for the CO<sub>2</sub>  
467 fluxes for mangrove shrublands (Table S2). If this flight were omitted from the October 2022  
468 mangrove shrubland disaggregation calculation, the average CO<sub>2</sub> for this vegetation class during  
469 October 2022 would be  $-8 \pm 4 \mu\text{mol m}^{-2} \text{s}^{-1}$ , similar to the estimates for the other sampling  
470 periods.

471 Measured CH<sub>4</sub> fluxes likely integrate contributions of water-atmosphere fluxes from  
472 mangrove tidal creeks and sediment-atmosphere fluxes. Insignificant CH<sub>4</sub> emissions were

473 measured for mangrove forests except during the October 2022 (end of wet season) period of  
474 high inundation, when average CH<sub>4</sub> fluxes were  $29 \pm 16 \text{ nmol m}^{-2} \text{ s}^{-1}$  (Fig. 6, lower panel).  
475 Fluxes from mangrove scrubs and shrublands were also higher in October 2022, though these  
476 areas tended to have larger dry-season CH<sub>4</sub> fluxes, ranging from 8 to 14  $\text{nmol m}^{-2} \text{ s}^{-1}$  among  
477 mangrove scrub and from -4 to 9  $\text{nmol m}^{-2} \text{ s}^{-1}$  for mangrove shrublands. Rosentreter et al.  
478 (2018a) observed a similar range of 0.5—12  $\text{nmol m}^{-2} \text{ s}^{-1}$  (40—1000  $\mu\text{mol m}^{-2} \text{ d}^{-1}$ ) water-  
479 atmosphere exchange from flux chambers at three sites in Australia, with the highest fluxes  
480 during the wet season. Much larger fluxes from mangrove soils of  $110 \pm 180 \text{ nmol m}^{-2} \text{ s}^{-1}$  ( $150 \pm$   
481  $250 \text{ mg m}^{-2} \text{ d}^{-1}$ ) were observed during the wet season in India (Jha et al., 2014), and CH<sub>4</sub> fluxes  
482 from soils ranged from 0.02 to 88  $\text{nmol m}^{-2} \text{ s}^{-1}$  at four mangrove sites in Taiwan (Lin et al.,  
483 2020). There is a large variability of CH<sub>4</sub> emissions from mangrove waters and soils that have  
484 been reported, with an estimated global average of  $3.9 \pm 1.2 \text{ nmol m}^{-2} \text{ s}^{-1}$  ( $339 \pm 106 \mu\text{mol m}^{-2} \text{ d}^{-1}$ )  
485 (Rosentreter et al., 2021).

486 Mangrove ghost forests predictably did not take up CO<sub>2</sub>. In these areas there had been a  
487 high tree mortality rate and massive defoliation post-Irma, without signs of recovery three years  
488 post-storm (Xiong et al., 2022). CO<sub>2</sub> exchange was not statistically different from zero during the  
489 first three deployment periods, but ghost forests served as a small source of CO<sub>2</sub> ( $1.1 \pm 0.8 \mu\text{mol}$   
490  $\text{m}^{-2} \text{ s}^{-1}$ ) during April 2023. It should be noted that during this deployment period we more heavily  
491 targeted ghost forests, particularly during the flight on April 18, 2023 (Table S1). Ghost forests  
492 were a methane source across all deployment months, particularly during October 2022 when we  
493 observed an average emission rate of  $51 \pm 27 \text{ nmol m}^{-2} \text{ s}^{-1}$ . During this period ghost forests  
494 emitted more CH<sub>4</sub> than any of the intact mangrove areas. Higher CH<sub>4</sub> emissions and eliminated  
495 CO<sub>2</sub> uptake from ghost forests highlights the importance of mangrove preservation for mitigation  
496 of carbon emissions, and the potential for additional GHG emissions as hurricanes and coastal  
497 development continue to threaten mangrove communities globally. This is particularly  
498 significant in south Florida mangrove communities, given the high tropical storm recurrence  
499 frequency in this region and the significant impacts of past hurricanes on forest structure and  
500 productivity (Danielson et al., 2017; Rivera-Monroy et al., 2019; Lagomasino et al., 2021; Xiong  
501 et al., 2022; Chavez et al., 2023).

## 502 *Saltwater Marshes*

503 Average daily NEE for saltwater marshes ranged from -6 to -12  $\mu\text{mol m}^{-2} \text{ s}^{-1}$  with no  
504 statistically significant seasonality. Similar fluxes of  $-6.7 \pm 5.5 \mu\text{mol m}^{-2} \text{ s}^{-1}$  (winter—spring) and  
505  $-7.9 \pm 6.4 \mu\text{mol m}^{-2} \text{ s}^{-1}$  were observed from an eddy-covariance tower in a tidal salt marsh in  
506 Brazil (Souza et al., 2022). An NEE range of -5 to -15  $\mu\text{mol m}^{-2} \text{ s}^{-1}$  was also recorded at a  
507 subtropical estuarine marsh in Taiwan (Lee et al., 2015).

508 CH<sub>4</sub> fluxes from saltwater marshes were insignificant, except during February 2023,  
509 when CH<sub>4</sub> fluxes were  $-11 \pm 4 \text{ nmol m}^{-2} \text{ s}^{-1}$ . Saline marshes typically emit less CH<sub>4</sub> than  
510 freshwater marshes because sulfate reduction dominates over methanogenesis during  
511 decomposition of organic matter (Bartlett et al., 1987). Low methane emission fluxes of  $0.08 \pm$   
512  $0.02 \text{ nmol m}^{-2} \text{ s}^{-1}$  ( $0.04 \pm 0.01 \text{ g m}^{-2} \text{ yr}^{-1}$ ) have also been observed in a tropical region of  
513 northwest Australia (Iram et al., 2021). In general, a large range of methane emissions from salt

514 marshes has been observed globally ( $-1$  to  $1090 \text{ nmol m}^{-2} \text{ s}^{-1}$ ,  $-92$  to  $94,000 \text{ } \mu\text{mol m}^{-2} \text{ d}^{-1}$ ), with an  
515 estimated average of  $2.6 \text{ nmol m}^{-2} \text{ s}^{-1}$  ( $224 \text{ } \mu\text{mol m}^{-2} \text{ d}^{-1}$ ) (Al-Haj and Fulweiler, 2020).

516 The reason for the different net  $\text{CH}_4$  and  $\text{CO}_2$  fluxes during each deployment is not  
517 immediately obvious. Methane and  $\text{CO}_2$  soil respiration fluxes in salt marshes are known to be  
518 influenced by tidal cycles (Kristensen et al., 2008; Rosentreter et al., 2018c; Iram et al., 2021)  
519 and it is possible that our flight data were skewed by sampling different tidal regimes. Methane  
520 uptake due to increased oxidation by methanotrophic bacteria has also been observed during the  
521 dry season of a coastal wetland in China (e.g. Hao et al., 2020). It is possible that abnormally low  
522 rainfall during February 2023 in southwest Florida contributed to the more significantly negative  
523 methane fluxes during this period.

#### 524 *Freshwater Marshes and Swamplands*

525 Freshwater marshes, swamp forests, swamp scrub, and swamp shrublands are all  
526 considered freshwater wetlands. Their different classifications reflect differences in vegetation  
527 composition and distribution, with a higher percentage of tall tree cover for swamp forests and a  
528 higher percentage of grasses in freshwater marshes (Section S3). The Shark River Slough, Taylor  
529 Slough, and several other sloughs that flow through the Big Cypress Swamp connect these areas  
530 with the saltwater tidal wetlands.  $\text{CO}_2$  uptake fluxes were relatively low over freshwater marshes  
531 ( $-2.9$  to  $-3.6 \text{ } \mu\text{mol m}^{-2} \text{ s}^{-1}$ ), with a weak seasonality. Similar daily peak fluxes for two freshwater  
532 marsh sites in ENP were observed using chamber measurements in 2008–2009 (Schedlbauer et  
533 al., 2012). The relatively low  $\text{CO}_2$  fluxes for freshwater marshes are likely due to the lower LAI  
534 and biomass for grasses than for regions containing larger shrubs and trees (Fig. 6, Fig S17, Fig  
535 S18). Inundation also plays a significant role in these systems, causing a decline in  
536 photosynthesis with increasing length of flooding (Zhao et al., 2021).  $\text{CO}_2$  fluxes in swamp  
537 shrublands were larger than in freshwater marshes, with a weak seasonality. These areas consist  
538 of a variety of evergreen tree species in a matrix of grasses (Supplementary Information Section  
539 S3). Average fluxes to swamp shrublands were more uncertain in April 2023, likely due to less  
540 area sampled than during other deployment periods (Table S1).

541 Many of the freshwater wetland regions of the Everglades, particularly freshwater  
542 marshes and swamp shrublands, contain periphyton mats in the water. These periphyton mats  
543 grow during the wet season and during periods of inundation, when they are active in fixing  $\text{CO}_2$   
544 from the atmosphere as calcium carbonate (Schedlbauer et al., 2012). The balance of  $\text{CO}_2$  uptake  
545 from plant and periphyton communities,  $\text{CO}_2$  emission from soils and waters, and the effect of  
546 inundation on these processes likely drives observed temporal changes in  $\text{CO}_2$  exchange.

547  $\text{CO}_2$  fluxes from swamp forests were largest during April 2022 and April 2023 ( $-10 \pm 5$   
548  $\text{ } \mu\text{mol m}^{-2} \text{ s}^{-1}$  and  $-10 \pm 4 \text{ } \mu\text{mol m}^{-2} \text{ s}^{-1}$ , respectively), during the dry- to wet-season transition  
549 period, with much smaller fluxes observed during October 2022 and February 2023 ( $-4 \pm 2 \text{ } \mu\text{mol}$   
550  $\text{ m}^{-2} \text{ s}^{-1}$  and  $-3 \pm 2 \text{ } \mu\text{mol m}^{-2} \text{ s}^{-1}$ , respectively) when the sun was lower and temperatures were  
551 cooler. Similar seasonality can be seen for swamp scrub. Swamp forests and swamp scrub  
552 vegetative regions are dominated by deciduous bald cypress trees (Ruiz et al., 2021). The

553 majority of these conifers were observed to either lack leaves or had brown needles during  
554 October and February flights.

555 CH<sub>4</sub> emissions were largest for freshwater marshes and swamp shrublands, with the  
556 largest emissions in October 2022 during the tail end of the wet season when soils were  
557 inundated. Average October 2022 fluxes for freshwater marshes, swamp forests, swamp scrubs,  
558 and swamp shrublands were  $59 \pm 26 \text{ nmol m}^{-2} \text{ s}^{-1}$ ,  $52 \pm 24 \text{ nmol m}^{-2} \text{ s}^{-1}$ ,  $33 \pm 18 \text{ nmol m}^{-2} \text{ s}^{-1}$ , and  
559  $69 \pm 29 \text{ nmol m}^{-2} \text{ s}^{-1}$ , respectively. Much larger CH<sub>4</sub> fluxes of 42–1200 nmol m<sup>-2</sup> s<sup>-1</sup>, (44–1244  
560 mg C-CH<sub>4</sub> m<sup>-2</sup> d<sup>-1</sup>) have been recorded in both marsh and forested wetland areas of Veracruz,  
561 Mexico (Marín-Muñiz et al., 2015) and Costa Rica (Nahlik and Mitsch 2011). More comparable  
562 CH<sub>4</sub> emission rates of 48–290 nmol m<sup>-2</sup> s<sup>-1</sup> (0.05–0.3 g C-CH<sub>4</sub> m<sup>-2</sup> d<sup>-1</sup>) during periods of  
563 inundation, and  $2 \pm 1 \text{ nmol m}^{-2} \text{ s}^{-1}$  ( $0.002 \pm 0.001 \text{ g C-CH}_4 \text{ m}^{-2} \text{ d}^{-1}$ ) during dry conditions were  
564 recorded from an eddy flux tower in the Pantanal wetland of Brazil (Dalmagro et al., 2019).  
565 Recently, Murguia-Flores et al. (2023) identified a median (IQR) CH<sub>4</sub> emission rate for tropical  
566 shallow-water inland wetlands of 39.2 (7.1–180.7) nmol m<sup>-2</sup> s<sup>-1</sup>, or 40.6 (7.4–187.3) g C-CH<sub>4</sub> m<sup>-2</sup>  
567 d<sup>-1</sup>. Our measurements from freshwater wetlands in southern Florida fall within this range and  
568 close to the reported median.

### 569 *Upland Woodlands*

570 Upland woodlands exhibited a moderately high NEE ranging from -4 to -8 μmol m<sup>-2</sup> s<sup>-1</sup>.  
571 A similar range (-6 to 12 μmol m<sup>-2</sup> s<sup>-1</sup>) of NEE for slash pine plantations in subtropical Australia  
572 was also observed across wet and dry seasons (McGowan et al., 2020). This vegetation region  
573 also demonstrates a similar but less pronounced seasonal cycle of CO<sub>2</sub> fluxes as swamp forests.  
574 Although most of the upland woodlands in this region are dominated by evergreen slash pine,  
575 this seasonality may be caused by variations in PAR and by the presence of some semi-  
576 deciduous species, such as laurel oaks. It is also possible that the productivity of flood-intolerant  
577 upland woodland species was somewhat suppressed during October 2022 at the tail of the wet  
578 season when water levels were relatively high (Fig S17). Reduction of NEE for October 2022  
579 could have also been driven by increased soil respiration during the wet season when soil  
580 moisture was higher (Fig S17), (Orchard and Cook 1983, Hawkes et al., 2016).

581 Methane fluxes from upland woodlands were undetectable in April 2022, and were  
582 slightly negative at  $-9 \pm 4 \text{ nmol m}^{-2} \text{ s}^{-1}$  and  $-9 \pm 5 \text{ nmol m}^{-2} \text{ s}^{-1}$  for February 2023 and April 2023,  
583 respectively. Similar negative fluxes have also been observed during the dry season in tropical  
584 upland forests in Costa Rica (Nahlik and Mitsch, 2011) and in the Amazon (Gauci et al., 2024).  
585 In contrast, a large positive flux of methane ( $40 \pm 19 \text{ nmol m}^{-2} \text{ s}^{-1}$ ) was observed during October  
586 2022. Wet season methane emissions have been observed in upland regions of tropical and  
587 subtropical regions elsewhere, when upland forests and woodlands can switch from being a  
588 methane sink to a methane source (Meronigol and Guenther, 2008).

### 589 3.2.2 Other surface properties

590 Within each vegetation land classification, there still exists substantial surface  
591 heterogeneity. For example, comparing the extent of mangrove forests (Fig. 2) with the maps of  
592 observed fluxes (Fig 5), it is apparent that this ecosystem exhibits a range of midday fluxes, even  
593 within the same month. The interquartile range of CO<sub>2</sub> fluxes observed where the footprint

594 consists of 80% mangrove forest is  $-15.1$  to  $-6.5 \mu\text{mol m}^2\text{s}^{-1}$  during April 2022. Similarly, the  
 595 interquartile range of  $\text{CH}_4$  fluxes observed where the footprint consists of 80% freshwater marsh  
 596 is 13 to  $73 \text{ nmol m}^2 \text{ s}^{-1}$  during April 2022. Vegetation type alone explains 35 - 53% of variability  
 597 in observed fluxes (Fig S16). Variability in the underlying drivers of  $\text{CO}_2$  and  $\text{CH}_4$  exchange  
 598 within an ecosystem type also influences heterogeneity in fluxes. For example, the range of LAI  
 599 for footprints containing 80% or more mangrove forests in April 2022 was 1.9 to  $6.3 \text{ m}^2/\text{m}^2$  and  
 600 the range of canopy heights was 5–20 m. For footprints containing 80% or more freshwater  
 601 marsh in October 2022, the range of water depths was 6–97 cm.

**Table 1:** Spearman's correlation coefficients for  $\text{CO}_2$  and  $\text{CH}_4$  fluxes and surface properties.

Data set	$\text{CO}_2$ all	$\text{CO}_2$ mangrove <sup>a,b</sup>	$\text{CH}_4$ all	$\text{CH}_4$ freshwater marsh <sup>a</sup>
<i>LAI</i>	-0.52	-0.36	-0.16	-0.02
<i>FPAR</i>	-0.52	-0.39	-0.07	0.07
<i>EVI</i>	-0.51	-0.51	-0.23	-0.17
<i>NDVI</i>	-0.46	-0.51	-0.14	0.06
<i>Canopy Height</i>	-0.35	-0.54	-0.33	-0.11
<i>AGBM</i>	-0.35	-0.54	-0.27	-0.13
<i>Soil moisture</i>	0.17	0.05	0.27	0.16
<i>water depth</i>	0.18	-0.33	0.53	0.44
<i>PAR</i>	-0.14	0.03	-0.10	-0.11
<i>VPD</i>	-0.32	-0.12	-0.02	-0.07
<i>T</i>	-0.06	0.02	-0.05	0.13
<i>RH</i>	0.25	0.12	-0.02	0.15

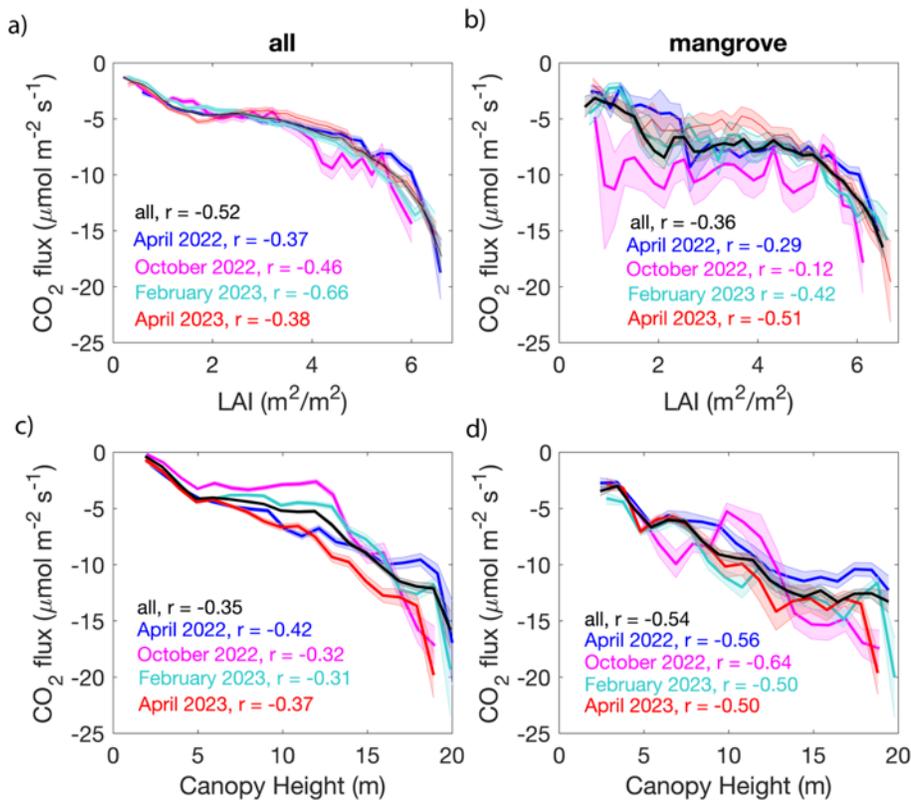
a. Considering footprints consisting of more than 80% of the given land type.

b. All mangrove ecosystems (forest, shrubland, scrub) are combined.

602 Spearman's correlation coefficients between 1 km averaged GHG fluxes and a variety of  
 603 surface and atmospheric variables are shown in Table 1. For surface data sets (LAI, FPAR, EVI,  
 604 NDVI, Canopy Height, AGBM, soil moisture, and water depth) correlation coefficients are  
 605 calculated between GHG fluxes and footprint weighted variables. Vapor pressure deficit (VPD),  
 606 temperature (T), and relative humidity (RH) are based on airborne temperature and water vapor  
 607 measurements. Photosynthetically active radiation (PAR) was estimated from the NOAA High-  
 608 Resolution Rapid Refresh (HRRR) Model product at 3 km resolution interpolated to the 1 km  
 609 averaged flight tracks. Spearman's correlation coefficients were used over Pearson's coefficients  
 610 because many of the relationships between the environmental and surface variables are non-  
 611 linear (Fig. 7, Fig 8).

612 The strongest predictors of  $\text{CO}_2$  fluxes for all flux data were LAI and FPAR (Table 1, Fig  
 613 7). However, within mangrove ecosystems (forest, shrublands, and scrub), canopy height and  
 614 above ground biomass had the most robust relationship with  $\text{CO}_2$  fluxes. In contrast,  $\text{CH}_4$  fluxes  
 615 over all flight tracks and over freshwater marshes both correlate best with EDEN water depth  
 616 (Table 1, Fig 8). The largest  $\text{CH}_4$  fluxes and EDEN water depths were over water management

617 regions outside of ENP or BCNP boundaries (Fig 3, Fig 5, Fig S17). The influence of water on  
 618 the methane emissions is not surprising, as numerous studies have observed larger methane  
 619 emissions in coastal wetlands during the wet season when soils are inundated and conditions in  
 620 the soil become more anaerobic and ideal for methanogenesis (e.g. Nahlik and Mitsch 2011;  
 621 Beringer et al., 2013; Marín-Muñiz et al., 2015; Dalmagro et al., 2019; Hondula et al., 2021).  
 622 This relationship confirms that higher CH<sub>4</sub> fluxes during October 2022 were likely due to  
 623 inundation. This flight period occurred following Hurricane Ian. Analysis of EDEN water data  
 624 sets since 2002 suggests that water levels were slightly elevated relative to the October average,  
 625 but comparable to many other years on record (Fig S19).



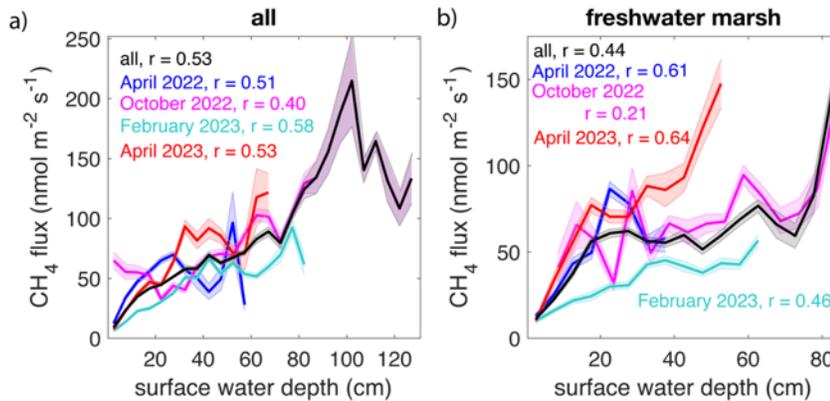
**Figure 7:** Relationships between CO<sub>2</sub> flux and leaf area index (LAI) (a, b) and between CO<sub>2</sub> flux and Canopy Height (c, d) for all 10 second averaged data (a, c), and for data where footprints were comprised of 80% mangroves (mangrove forests, mangrove shrublands, and mangrove scrubs) (b, d). CO<sub>2</sub> data are averaged over 0.25  $\text{m}^2 \text{m}^{-2}$  LAI bins or 2 m canopy height bins. Solid black, blue, magenta, turquoise, and red lines are the average for all flights, April 2022, October 2022, February 2023, and April 2023, respectively. Shaded areas are the 95% confidence interval of the mean.

626 Interestingly, we observe a slightly negative relationship between VPD and CO<sub>2</sub> flux.  
 627 This is likely both due to the generally high humidity and water availability in the region (50—  
 628 90%) and because VPD was higher during the April months when there was greater sunlight

629 availability and the deciduous bald cypress trees were green. Similar relationships between VPD  
 630 and mangrove NEE have also been reported for mangroves in the Yucatan (Alvarado-Barrientos  
 631 et al., 2020).

632 Some relationships between fluxes and environmental variables likely come about  
 633 because of redundant and non-causal correlations. For example, the negative relationship  
 634 between canopy height and CH<sub>4</sub> fluxes, are likely due to correlations between surface variables.  
 635 In this case, greater canopy height does not cause lower CH<sub>4</sub> emissions, but areas with greater  
 636 canopy heights are often mangrove forests and areas with less surface water extent where there  
 637 are low CH<sub>4</sub> emissions. Many of the variables tested, such as EVI, NDVI, Canopy Height, LAI,  
 638 and AGBM also co-vary with each other.

639 The relationships between remotely sensed vegetation and soil properties and carbon  
 640 fluxes demonstrate the potential predictive power of remote sensing for greenhouse gas fluxes.  
 641 Incorporation of remotely sensed data sets into a predictive machine learning model of southern  
 642 Florida CO<sub>2</sub> and CH<sub>4</sub> fluxes is a part of ongoing work.



**Figure 8:** Relationships between CH<sub>4</sub> flux and EDEN surface water depth all 10 second averaged data (a), and for data where footprints were comprised of 80% freshwater marsh (b). CH<sub>4</sub> fluxes are averaged over water depth in 5 cm bins. Solid black, blue, magenta, turquoise, and red lines are the average for all flights, April 2022, October 2022, February 2023, and April 2023, respectively. Shaded areas are 95% confidence intervals. All 10-second averaged data include areas outside of the national park boundaries where vegetation data area available.

643

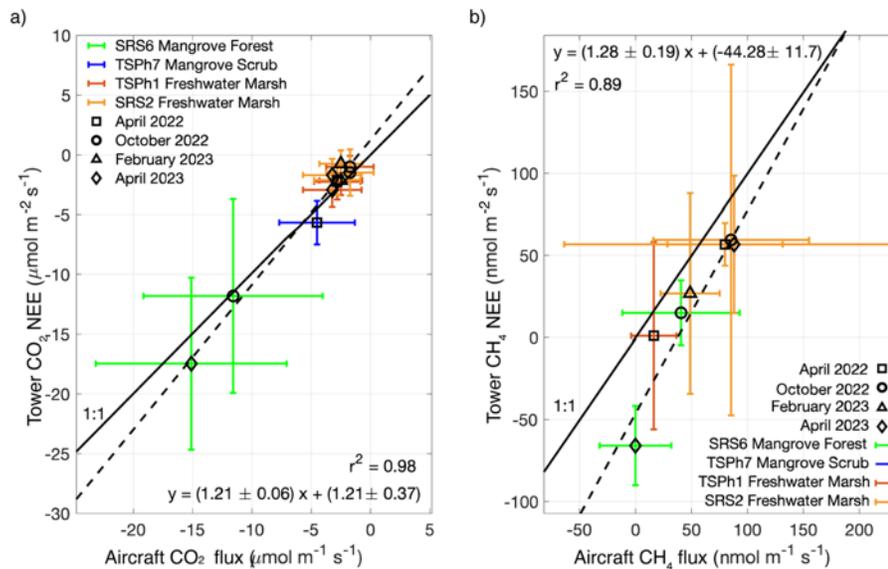
### 644 3.3 Flux tower comparisons

645 Comparison of airborne fluxes to EC flux tower measurements requires careful  
 646 consideration of flux footprints (Hannun et al., 2020). Only one flight had a flux footprint that  
 647 directly overlapped with the mangrove forest tower site (SRS-6) while that tower site was active,  
 648 and measurements only overlapped for several seconds, making a direct comparison with the  
 649 flux towers impossible. Indirect comparison of airborne and EC tower fluxes is complicated by  
 650 surface variability and resulting heterogeneity of fluxes. Each airborne flux footprint typically

651 consisted of several vegetation types and a further range of surface properties (LAI, canopy  
 652 heights, soil moisture, surface water extent, etc.). In contrast, the EC tower footprints typically  
 653 covered only single ecosystem type with greater homogeneity.

654 We indirectly compare the average monthly fluxes of CH<sub>4</sub> and CO<sub>2</sub> from eddy covariance  
 655 tower ground sites (when available) to airborne flux measurements with footprints consisting  
 656 primarily of similar surface properties as the ground sites. Airborne CO<sub>2</sub> and CH<sub>4</sub> fluxes were  
 657 averaged over all flight days during a given month after selecting for data that met certain criteria  
 658 for comparison with the EC tower. For CH<sub>4</sub> flux comparisons with SRS-2 and TS/Ph-1, airborne  
 659 data were filtered to only include points where freshwater marsh constituted more than 80% of  
 660 the footprint and the footprint weighted average water depth was within 10 cm of the EC tower  
 661 for the given month (Table S6). For SRS-6 comparison, airborne fluxes were included in the  
 662 average if footprints contained more than 80% mangrove forest. Airborne CO<sub>2</sub> fluxes were  
 663 averaged during each flight day where footprints constituted over 80% of the EC tower land  
 664 classification and had a footprint weighted LAI within 1 m<sup>2</sup>/m<sup>2</sup> of the EC tower footprint LAI for  
 665 the given month (Table S7). EC tower fluxes were averaged for all available data during a given  
 666 month from 10:00—17:00 local time (LT).

667 Comparisons indicate relatively good agreement between EC flux tower and airborne  
 668 flux measurements (Fig. 9). These comparisons provide a validation of our airborne CWT fluxes,  
 669 as well as confirm that surface water extent, LAI, and vegetation class indeed capture much of  
 670 the observed variability in carbon exchange. Tower comparisons with latent heat (LE) and  
 671 sensible heat (H) are discussed in Supplementary Information section S4 and Figures S6-S8.



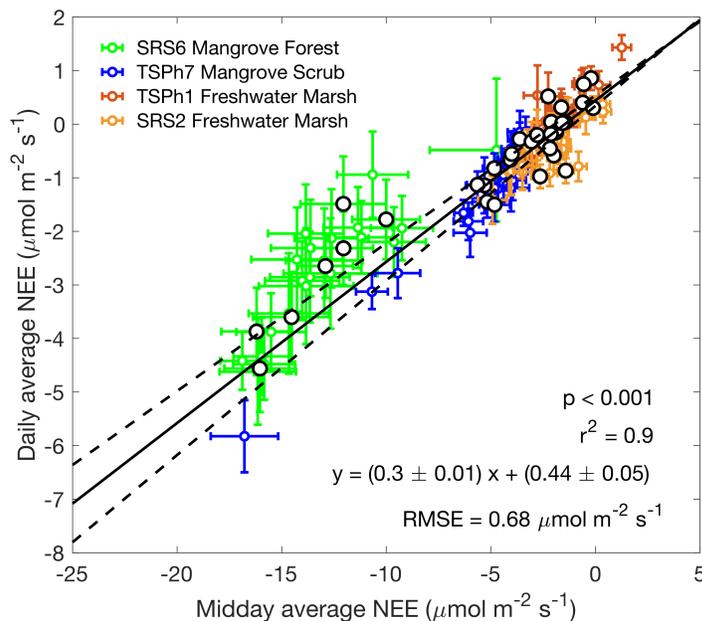
**Figure 9:** Comparison of EC fluxes from flux tower sites with airborne fluxes measured from the King Air for a) CO<sub>2</sub> and b) CH<sub>4</sub>. EC tower CH<sub>4</sub> and CO<sub>2</sub> NEE (flux – storage) for a given month were averaged over all available data between 10:00 and 16:00 LT. Airborne CH<sub>4</sub> fluxes were averaged over all flight days for data with the same vegetation class and surface water as the EC tower. Airborne CO<sub>2</sub> fluxes were averaged during each flight day after filtering data for the same vegetation class and LAI as the EC tower. EC towers SRS-6, SRS=2, TS/Ph-1, and TS/Ph-7 sample mangrove forest, freshwater marsh, freshwater marsh, and mangrove scrub, respectively. Error bars are one standard deviation. The dashed line is the 1:1 line.

672

## 673 3.4. Estimates of net carbon fluxes

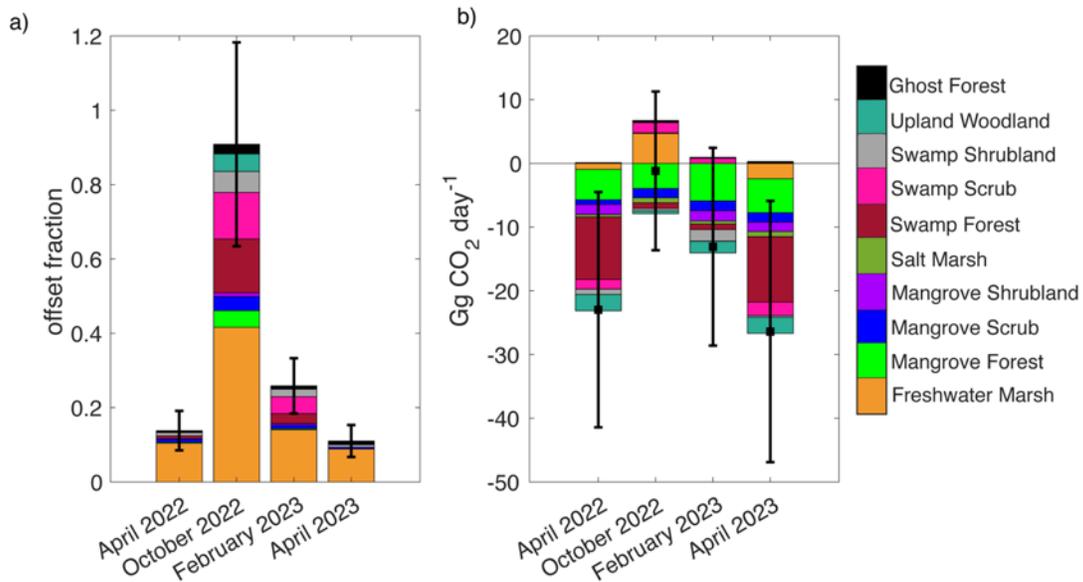
674

675 The balance between CH<sub>4</sub> emissions and CO<sub>2</sub> uptake partially determines the impact of  
 676 wetland ecosystems on carbon sequestration and climate change mitigation. To estimate the net  
 677 impact of the entire Everglades region on carbon exchange during our sampling periods, we first  
 678 scaled CH<sub>4</sub> fluxes by their CO<sub>2</sub> equivalent global warming potential (GWP) using a factor of  
 679 27.9 g CH<sub>4</sub>/g CO<sub>2</sub> in accordance with the latest IPCC report (Forster et al., 2021) and then  
 680 calculated the net CO<sub>2</sub> equivalent exchange rate and the fraction of total CO<sub>2</sub> uptake that is  
 681 offset by CH<sub>4</sub> emissions for each vegetation region based on disaggregated fluxes. These values  
 682 were then scaled by the total area of each vegetation region adequately sampled in ENP and  
 683 BCNP (Fig. S20, Fig. S21). During daytime sampling periods, the 6,237 km<sup>2</sup> area of south  
 684 Florida represented by the sampled land classes has a total CO<sub>2</sub> equivalent exchange rate of  $-5.3$   
 685  $\pm 2.6$  to  $-2.7 \pm 1.5$  Gg CO<sub>2</sub> hr<sup>-1</sup>, with CH<sub>4</sub> emissions offsetting CO<sub>2</sub> by  $3 \pm 1\%$  to  $14 \pm 4\%$ ,  
 686 depending on the month. The largest CH<sub>4</sub> emissions relative to CO<sub>2</sub> uptake occurred during the  
 687 October 2022 deployment (Fig. S21). However, midday airborne flux measurements do not  
 688 include nighttime CO<sub>2</sub> respiration which is required for an estimate of daily carbon exchange.



**Figure 10:** Linear fit of daily 24 hr average net ecosystem exchange (NEE) and the average daytime NEE between LT 10:00 and 17:00 for all tower sites in the flight domain. Each marker represents a monthly average from data available between 2020 and 2023. Larger black markers are monthly averages during April, February, and October. The line of best fit and the 95% confidence interval are solid and dotted lines, respectively. Error bars are  $1\sigma$ .

689



**Figure 11:** (a) GWP-scaled CH<sub>4</sub> emissions as a fraction of the CO<sub>2</sub> uptake for each month across the BlueFlux experimental domain. Colors indicate the contribution of each ecosystem classification. CH<sub>4</sub> offset fractions are weighted by the area of each ecosystem type. (b) Average daily net CO<sub>2</sub> equivalent uptake for each deployment period, calculated as the sum of CO<sub>2</sub> and GWP-scaled CH<sub>4</sub> fluxes. Total daily CO<sub>2</sub> fluxes were calculated by from the daytime airborne measurements using the linear fit of daily integrated NEE and daytime NEE from Figure 10. CH<sub>4</sub> fluxes were assumed to be constant throughout the day so average disaggregated CH<sub>4</sub> fluxes for each deployment period were scaled by 24 hours. Ecosystem contributions are scaled by area. Error bars are 1σ.

690 Tower flux observations constrain the diurnal cycle of carbon exchange in several  
 691 locations. We use this diurnal temporal information from the Everglades tower network to  
 692 extrapolate daytime fluxes (LT 10:00–17:00) to total daily carbon exchange in ENP and BCNP  
 693 as in Hannun et al. (2020) (Figure S22). The tower sites in the BlueFlux/FCE LTER domain are  
 694 located in three different vegetation regions that represent the three dominant vegetation types,  
 695 including tall riverine mangrove forests (SRS-6), scrub mangroves (TS/Ph-7), and freshwater  
 696 marshes (SRS-2 and TS/Ph-1). These tower sites demonstrate a linear relationship between the  
 697 total CO<sub>2</sub>-NEE between LT 10:00 and 17:00 and the total daily integrated NEE (Fig. 10,  $r =$   
 698 0.94). A similar relationship was also derived for EC tower sites in the mid-Atlantic region over  
 699 several different ecosystem types (Hannun et al., 2020). We scale the average ecosystem CO<sub>2</sub>  
 700 fluxes measured by CARAFE during LT 10:00–17:00 (Fig. 6) by the relationship shown in

701 Figure 10 to estimate the 24 hr average NEE for each vegetation class during the months of our  
702 flight measurements.

703 The CH<sub>4</sub> average midday flux measured by the tower sites was not significantly different  
704 from the daily average (Figure S23). Thus, CH<sub>4</sub> fluxes are treated as constant throughout the day  
705 and average daytime CH<sub>4</sub> fluxes were assumed to be representative of daily averaged fluxes.

706 Figure 11 shows the resulting daily CH<sub>4</sub> offsets to CO<sub>2</sub> uptake and net daily CO<sub>2</sub>  
707 equivalent exchange for each deployment period across ENP and BCNP. CH<sub>4</sub> emissions relative  
708 to CO<sub>2</sub> uptake are lowest during April 2023, with CH<sub>4</sub> emissions offsetting  $11 \pm 4\%$  of CO<sub>2</sub>  
709 uptake and a total net carbon exchange rate of  $-26 \pm 20$  Gg CO<sub>2</sub>-eq d<sup>-1</sup>. The largest CH<sub>4</sub> emissions  
710 relative to CO<sub>2</sub> uptake are during October 2022, with CH<sub>4</sub> emissions offsetting  $91 \pm 27\%$  of CO<sub>2</sub>  
711 uptake and a net carbon exchange rate of  $-1 \pm 12$  Gg CO<sub>2</sub>-eq d<sup>-1</sup>. This range of uncertainty in the  
712 net carbon exchange is large, and as such the region may either serve as a net source or net sink  
713 of carbon during October. The largest source of CH<sub>4</sub> emissions relative to CO<sub>2</sub> uptake are the  
714 freshwater marshes, particularly during the October 2022 wet season, when we estimate that  
715 they serve as a 5 Gg CO<sub>2</sub>-eq d<sup>-1</sup> source of carbon. Mangrove forests contribute a relatively small  
716 amount of CH<sub>4</sub> emission and provide the largest net sink of carbon cumulatively across all  
717 deployment periods with an average net carbon flux of  $-5 \pm 2$  Gg CO<sub>2</sub>-eq d<sup>-1</sup>. Swamp forests  
718 provide the largest net carbon sink during the dry/growing season (almost double than that of  
719 mangroves), as observed during April 2022 and April 2023.

720 Carbon exchange in southern Florida wetlands exhibits strong seasonality over the  
721 measurement period, with the ENP and BCNP region potentially serving as a net source of  
722 carbon to the atmosphere during the wet season and periods of high inundation. Mangrove and  
723 cypress swamp forests are large atmospheric sinks of carbon for the region, despite their  
724 relatively small extents. The importance of cypress swamp forests in CO<sub>2</sub> removal is likely even  
725 greater than this study reflects, as we do not have measurements in the summer when leaf area is  
726 at a peak.

727 The uncertainty in our estimates of daily carbon balance over ENP and BCNP is large, as  
728 it includes the propagated uncertainty in the net monthly fluxes due to variability within an  
729 ecosystem type (i.e due to LAI, water depth, soil moisture, tidal cycles, temperature, salinity,  
730 effects of recent hurricanes and recovery, etc), uncertainty in airborne fluxes, uncertainty in the  
731 disaggregation of fluxes, and uncertainty in the diurnal upscaling. However, these results serve  
732 as a first EC-based estimate of net carbon fluxes over the entire Everglades and Big Cypress area  
733 and are demonstrative of the temporal differences in carbon exchange across the region during  
734 different months, as well as the potential importance of the different vegetation regions in the  
735 carbon balance.

736 There are several additional limitations to this diurnal upscaling analysis that should be  
737 noted. Each tower only covers an extremely limited area within an ecosystem type, while the  
738 data sampled from the flights includes a much larger range of conditions and surface variability  
739 (LAI, canopy height, water depth, tidal influences, salinity, etc.). We have thusly chosen to use a  
740 uniform diurnal scaling factor for each vegetation type rather than a separate tower-derived  
741 relationship for mangrove forests, mangrove scrub, and freshwater marshes as we believe it to be  
742 more representative of the observed variability of conditions throughout our flight domain.

743 Results using a distinct relationship for mangrove forests, mangrove scrub, and freshwater  
744 marshes can be found in Figure S24.

745 The domain of swamp forests, swamp scrubs, and upland woodlands were outside the  
746 tower FCE LTER study area, and are not represented in deriving the diurnal upscaling  
747 relationship. Eddy covariance flux tower measurements conducted in BCNP during 2012—2014  
748 measured average CO<sub>2</sub> NEE of  $-108 \pm 5$  g C m<sup>-2</sup> month<sup>-1</sup>,  $-48 \pm 3$  g C m<sup>-2</sup> month<sup>-1</sup>, and  $-68 \pm 5$  g  
749 C m<sup>-2</sup> month<sup>-1</sup> during April months at a cypress swamp (swamp forest), dwarf cypress (swamp  
750 scrub), and pine upland (upland woodland) site, respectively (Shoemaker et al., 2015). We obtain  
751 lower average estimates for these vegetation areas during our 2022—2023 April deployments of  
752  $-85 \pm 40$  g C m<sup>-2</sup> month<sup>-1</sup>,  $-16 \pm 10$  g C m<sup>-2</sup> month<sup>-1</sup>, and  $-51 \pm 30$  g C m<sup>-2</sup> month<sup>-1</sup>, respectively,  
753 when scaling using the relationship derived in Figure 10. While the 2012–2014 tower  
754 measurements for swamp forests and upland woodlands are within the range of uncertainty of  
755 our estimates, the tower measurement from the swamp scrub site is not. It is not clear if this  
756 difference is due to error in our analysis, the decade long separation between tower and flight  
757 measurements, or flux variability over this ecosystem type not captured by the tower.

758 It should also be noted that in tidal wetland regions like the Everglades, the net ecosystem  
759 carbon balance is affected by lateral aqueous transport of carbon in addition to NEE (Troxler et  
760 al., 2013). Carbon initially taken up in one area may be stored in above and below ground  
761 biomass, soils, and sediments, or it may laterally flow from the area of initial uptake to later be  
762 reemitted or stored in soils and sediments downstream (Bouillon et al., 2008; Alongi and  
763 Mukhopadhyay, 2015; Rosentreter et al., 2018b). Aquatic lateral transport in the Everglades has  
764 been estimated to be relatively small (~10%) compared to the NEE of mangrove forests (Troxler  
765 et al., 2013). However, freshwater marshes in the Everglades store a substantial amount of  
766 carbon ( $400\text{--}650$  g C m<sup>-2</sup> yr<sup>-1</sup>), with almost all the carbon input through aqueous lateral transport  
767 (Troxler et al., 2013). In this study we discuss the net vertical carbon exchange largely from an  
768 atmospheric perspective, but it should not be taken as the complete story of carbon storage in the  
769 Everglades.

## 770 **5 Conclusions**

771 Airborne eddy covariance with continuous wavelet transforms can resolve heterogenous  
772 fluxes over a diverse mosaic of ecosystems across the coastal landscape of southern Florida. The  
773 largest CO<sub>2</sub> uptake fluxes were observed during April 2022 and 2023 over cypress swamp forests  
774 and over mangrove forests during all sampling periods. During the tail-end of the wet season and  
775 near maximum water levels (October 2022 campaign), we observed the largest CH<sub>4</sub> emission  
776 fluxes from all vegetation types. Across all deployments, we recorded the largest CH<sub>4</sub> fluxes  
777 from freshwater marshes and freshwater swamp shrublands. Additionally, we see some evidence  
778 for CH<sub>4</sub> uptake during the dry season in salt marshes and upland forests. Upscaling average  
779 ecosystem fluxes over the sample domain, we estimate average net CO<sub>2</sub>-eq fluxes of  $-4 \pm 3$  g  
780 CO<sub>2</sub>-eq m<sup>-2</sup> d<sup>-1</sup> in April and  $-0.2 \pm 2$  g CO<sub>2</sub>-eq m<sup>-2</sup> d<sup>-1</sup> in October (area-integrated rates of  $-26 \pm 20$   
781 Gg CO<sub>2</sub>-eq d<sup>-1</sup> and  $-1 \pm 12$  Gg CO<sub>2</sub>-eq d<sup>-1</sup>, respectively).

782 Our findings highlight the role of freshwater swamp forests and mangrove forests as  
783 extremely productive coastal ecosystems. Observed rates of CO<sub>2</sub> uptake and CH<sub>4</sub> emission for  
784 these ecosystems fall within the range of observations for mangroves and swamp forests in

785 similar subtropical and tropical regions globally. Results support the importance of these  
786 ecosystems as important carbon stores and the potential for additional GHG emissions resulting  
787 from hurricanes, coastal development, and hydrological management. However, the diversity and  
788 vulnerability of these ecosystems necessitates continued ongoing research into the carbon storage  
789 potential and the effects of restoration and degradation on the role of swamp and mangrove  
790 forests in the global coastal carbon cycle.

791 Combined with landcover information like vegetation type, leaf area, canopy height,  
792 vegetation indices, and surface water depth, airborne fluxes can help elucidate the underlying  
793 causes of the observed variability in carbon fluxes. In particular, surface water depth in the  
794 freshwater wetlands was strongly positively correlated with CH<sub>4</sub> emissions. In a heavily water  
795 managed area like southern Florida, policy decisions related to agriculture and hydrology may  
796 overlap with greenhouse gas reduction strategies. Moreover, the ongoing large-scale  
797 hydrological restoration of the Greater Everglades under the Comprehensive Everglades  
798 Restoration Plan (CERP) will likely have significant effects on vegetation dynamics, especially  
799 carbon storage and sequestration potential, thereby influencing the role of wetlands in climate  
800 change mitigation and adaptation. The relationship between surface water and methane emission  
801 in this study relied heavily on the long-term Everglades Depth Estimation Network data set.  
802 However, such high-resolution surface information is currently extremely limited globally.  
803 Improvements in high resolution remotely sensed soil moisture and surface water data will be  
804 critical for ongoing research into relationships between regional hydrology and global methane  
805 emissions. Other surface information, such as high-density coastal wetland salinity maps, would  
806 also be beneficial to this analysis.

807 A limitation of this study is the lack of observations during the peak of the wet season  
808 (May-September). Additional airborne flux measurements in southern Florida are needed to  
809 better constrain seasonality (particularly wet-season fluxes), diurnal cycles, and tidal influences.  
810 These efforts would improve the ongoing carbon budget analysis of coastal wetlands in the  
811 Everglades region and add an understanding of the carbon sink and source capacity of these  
812 ecosystems exposed to increasing impacts of sea-level rise and climate change.

813 The importance of vulnerable coastal wetland ecosystems to the CO<sub>2</sub> and CH<sub>4</sub> global  
814 budgets highlights the need for continued and sustained measurements in these regions. Airborne  
815 eddy covariance, especially paired with remote-sensing surface information, represents a  
816 powerful tool for constraining biogenic carbon cycles.

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837

### 838 **Open Research**

839 The airborne data used for all analysis in the study (concentrations, fluxes, meteorology, and aircraft  
840 navigational parameters) are openly available at ORNL DAAC via  
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842

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