



1 **Constraining anthropogenic CO₂ fluxes in mixed urban** 2 **source areas using eddy covariance and footprint-** 3 **informed ecological parameter migration**

4 Jinwen Zhang^{1,2}, Yongjian Liang^{2,3}, Yufei Yang^{1,2}, Chenglei Pei^{2,3}, Hebiao Huang⁴,
5 Bo Huang⁵, Xiufeng Lian^{1,2}, Yingyan Huang^{2,3}, Yuwen Peng^{2,3}, Chunlei Cheng^{1,2},
6 Cheng Wu^{1,2}, Zhen Zhou^{1,2}, and Mei Li^{1,2}

7 ¹College of Environment and Climate, Institute of Mass Spectrometry and Atmospheric
8 Environment, Guangdong Provincial Engineering Research Center for Online Source
9 Apportionment System of Air Pollution, Guangdong–Hongkong–Macau Joint Laboratory of
10 Collaborative Innovation for Environmental Quality, Jinan University, Guangzhou 511443, China

11 ²Key Laboratory of Regional Air Quality Monitoring, Ministry of Ecology and Environment,
12 Guangdong, Guangzhou 510308, China

13 ³Guangzhou Ecological and Environmental Monitoring Center Station, Guangdong, Guangzhou
14 510006, China

15 ⁴Guangzhou Academy of Agricultural and Rural Sciences, Guangdong, Guangzhou 510450, China

16 ⁵Guangzhou Hexin Instrument Co., Ltd., Guangdong, Guangzhou 510530, China

17 Correspondence: Mei Li (limei@jnu.edu.cn)

18 **Abstract.** Urban net CO₂ fluxes measured by eddy covariance (EC) cannot be treated
19 as anthropogenic emissions without biogenic constraints, because they integrate fossil-
20 fuel sources, biogenic processes, human respiration, and changing footprints. Existing
21 partitioning approaches often require isotopes, tracers, multi-species fluxes, or clean
22 vegetation/emission sectors unavailable at many urban sites. We developed a footprint-
23 informed ecological parameter-migration framework to constrain anthropogenic CO₂
24 fluxes (F_H) in mixed urban source areas of Guangzhou, a humid subtropical megacity.
25 A highly vegetated, weakly disturbed suburban donor site provided ecological
26 parameters, which were transferred to a mixed urban target site using dynamic
27 footprints and footprint-weighted enhanced vegetation index (EVI_{fp}). The migrated
28 parameters produced plausible diurnal and seasonal gross primary productivity (GPP)
29 and ecosystem respiration (R_{eco}), supporting the donor–target constraint. At the target
30 site, daytime biogenic uptake masked anthropogenic emissions and occasionally drove
31 net CO₂ flux toward neutral or weakly negative values. After subtracting R_{eco} , GPP, and
32 human respiration, F_H remained consistently positive throughout the day (1.80–5.14
33 $\mu\text{mol m}^{-2} \text{s}^{-1}$). F_H showed morning and evening peaks consistent with NO_x, CO, wind-
34 sector source areas, and traffic indicators, while footprint-aligned inventories provided



35 magnitude context and suggested spatial-proxy mismatch in a high-population urban
36 functional zone with relatively low on-site combustion. Uncertainty was dominated by
37 GPP light-response structure, with smaller effects from other perturbations. These
38 results demonstrate the feasibility of extracting footprint-scale F_{ff} from mixed urban
39 source areas using conventional EC, remote-sensing, and footprint data where isotopic
40 or multi-species flux observations are unavailable.

41 **1 Introduction**

42 Urban CO₂ monitoring increasingly needs to determine not only how much CO₂ cities
43 emit, but also how observed CO₂ signals are jointly shaped by fossil-fuel combustion,
44 biogenic processes, human respiration, and atmospheric transport (Sargent et al., 2018;
45 Zhang et al., 2026). This distinction is particularly important in urban landscapes where
46 green space, roads, buildings, residential activity, and water bodies are highly
47 interwoven. Although cities generally act as net CO₂ sources, urban vegetation can
48 substantially modulate local CO₂ exchange and can partially offset, mask, or reshape
49 the diurnal and seasonal signatures of anthropogenic emissions (Velasco and Roth, 2010;
50 Nordbo et al., 2012; Hardiman et al., 2017; Miller et al., 2020). Therefore, net CO₂
51 exchange observed in urban environments cannot be directly interpreted as fossil-fuel
52 or anthropogenic CO₂ emissions unless biogenic processes are explicitly constrained
53 (Wu et al., 2022; Soininen et al., 2025).

54 Eddy covariance (EC) is one of the few approaches capable of directly measuring
55 surface-atmosphere CO₂ exchange at high temporal resolution (Baldocchi, 2003;
56 Velasco and Roth, 2010). It has been widely used to quantify ecosystem carbon
57 exchange and has increasingly been applied to urban CO₂ flux observations (Järvi et al.,
58 2012; Liu et al., 2012; Matthews and Schume, 2022; Ueyama and Takano, 2022; Horne
59 et al., 2026). Urban EC studies have shown that CO₂ fluxes often exhibit pronounced
60 diurnal, seasonal, and source-area dependence, reflecting the combined influences of
61 traffic activity, building energy use, vegetation phenology, land-cover composition, and
62 meteorological conditions (Velasco et al., 2013; Ward et al., 2015; Pérez-Ruiz et al.,
63 2020). However, EC directly measures the net exchange of all CO₂ sources and sinks
64 within a dynamic flux footprint (Menzer and McFadden, 2017; Lee et al., 2021). In



65 mixed urban source areas, a daytime reduction or even reversal of net CO₂ flux does
66 not necessarily indicate weakened anthropogenic emissions, but may at least partly
67 reflect enhanced photosynthetic uptake (Sargent et al., 2018; Wu et al., 2022). Thus, the
68 key challenge for urban EC studies is not whether CO₂ fluxes can be observed, but how
69 net fluxes can be partitioned into physically interpretable anthropogenic and biogenic
70 components.

71 Previous studies have attempted to partition urban EC net CO₂ fluxes using
72 inventories, statistical methods, source-area sectors, and tracers. Early work combined
73 EC observations with bottom-up inventories to evaluate urban CO₂ emissions or
74 attribute them to traffic and domestic heating (Matese et al., 2009). Such comparisons
75 provide useful magnitude context, but inventory-based interpretation alone cannot
76 separate the biogenic and anthropogenic components embedded in EC net fluxes at the
77 dynamic footprint scale. Menzer and McFadden (2017) statistically partitioned multi-
78 year urban net CO₂ fluxes into biogenic and anthropogenic components, and Lee et al.
79 (2021) incorporated footprint-weighted land cover and temporal subsets to separate
80 gross primary productivity (GPP), ecosystem respiration (R_{eco}), vehicle emissions, and
81 building emissions. More recently, O₂:CO₂, CO, radiocarbon (¹⁴C), carbonyl sulfide
82 (COS), and multi-species EC measurements have provided stronger constraints on
83 urban CO₂ flux partitioning (Ishidoya et al., 2020; Wu et al., 2022; Soininen et al., 2025;
84 Hilland et al., 2025; Kunz et al., 2026). In particular, Soininen et al. (2025) fitted
85 ecological light-response parameters in a vegetation-dominated sector and applied them
86 to a street-sector footprint, obtaining residual anthropogenic CO₂ fluxes comparable to
87 bottom-up traffic emissions. This provides an important example in which parameters
88 derived from a relatively clean ecological source area can be used to estimate biogenic
89 contributions in a mixed or emission-dominated source area. Beyond source
90 partitioning, such footprint-scale anthropogenic flux estimates can also help evaluate
91 whether proxy-based gridded inventories allocate emissions realistically within
92 heterogeneous urban functional zones, where population density, night-time lights, or
93 road density may not scale with local combustion intensity (Gately and Hutyra, 2017;
94 Squires et al., 2020; Wu et al., 2022).

95 Nevertheless, these approaches remain difficult to generalize. Methods based on



96 ^{14}C , COS, $\text{O}_2:\text{CO}_2$, or multi-species EC usually require additional instrumentation,
97 sampling, or calibration infrastructure (Ishidoya et al., 2020; Soininen et al., 2025; Kunz
98 et al., 2026), whereas statistical and sector-transfer approaches often require clearly
99 separated vegetation-dominated and emission-dominated source areas around the tower
100 (Menzer and McFadden, 2017; Lee et al., 2021; Soininen et al., 2025). For cities
101 equipped only with conventional EC, meteorological, remote-sensing, and land-cover
102 data, this limitation is particularly important where vegetation, roads, buildings, and
103 water bodies are tightly mixed and no clean vegetation sector is available. In such
104 settings, the partitioning problem becomes: how can the biogenic component be
105 robustly estimated at the dynamic EC-footprint scale and removed from the net flux?

106 Conventional ecosystem flux partitioning usually estimates R_{eco} from nighttime
107 temperature responses and GPP from daytime light-response curves (Reichstein et al.,
108 2005; Lasslop et al., 2010). Directly fitting these ecological parameters at an urban EC
109 site is problematic because fluxes may be contaminated by traffic, building-related
110 emissions, human activity, and storage changes. Fixed-buffer vegetation indices or
111 land-cover fractions also ignore the continuous variation of EC source areas with wind
112 direction, turbulence, and atmospheric stability. Flux footprint models can link each
113 flux observation to its actual contributing source area (Kljun et al., 2015), while the
114 enhanced vegetation index (EVI) can represent canopy greenness, vegetation activity,
115 and photosynthetic potential (Huete et al., 2002; Mahadevan et al., 2008). However,
116 dynamic footprints and vegetation indices alone do not identify the baseline ecological
117 response at the target site, because the target-site net flux remains a superposition of
118 biogenic exchange and anthropogenic emissions. This motivates the hypothesis tested
119 here: ecological parameters obtained from a nearby donor site with weaker
120 anthropogenic disturbance may provide the missing ecological response reference and
121 can be transferred to a mixed urban target site when constrained by vegetation amount
122 and activity within the dynamic target footprint.

123 Such cross-site parameter migration should avoid mechanically imposing donor-
124 site parameters on the target footprint. Ecosystem-scale light-response parameters are
125 apparent canopy-scale quantities controlled by canopy structure, vegetation amount,
126 phenology, and environmental conditions, rather than fixed leaf-level constants (Sellers,



127 1985; Myneni and Williams, 1994; Huete et al., 2002). This provides the basis for
128 treating donor-site parameters as a same-region ecological-response reference when
129 donor and target footprints share broadly similar climatic and vegetation backgrounds.
130 Similar logic is used in light-use-efficiency and VPRM-style models, where vegetation
131 indices are combined with environmental scalars to estimate photosynthetic uptake and
132 urban biogenic CO₂ fluxes (Monteith, 1972; Running et al., 2004; Xiao et al., 2004;
133 Mahadevan et al., 2008; Gourdji et al., 2022). In this framework, footprint-weighted
134 EVI links remotely sensed vegetation activity to the actual EC source area (Huete et al.,
135 2002; Kljun et al., 2015; Järvi et al., 2019). For GPP, this constraint is relatively direct
136 because EVI tracks photosynthetically active vegetation amount and canopy greenness,
137 and is therefore closely linked to canopy light-absorbing capacity. For R_{eco}, EVI is used
138 more cautiously as a footprint-scale vegetated/pervious-surface mask and capacity-
139 normalization factor, rather than as a direct physiological driver. This treatment is
140 consistent with urban biogenic CO₂ models that use vegetation fractions or remote-
141 sensing vegetation proxies together with meteorological and urban-surface modifiers to
142 represent vegetation and soil respiration (Järvi et al., 2019; Hardiman et al., 2017; Wu
143 et al., 2021), and with urban soil-respiration measurements showing substantial
144 respiration from vegetated and managed pervious cover types (Decina et al., 2016).
145 Therefore, both respiration and light-response parameter migration are treated as
146 constrained approximations and evaluated through sensitivity analyses.

147 Based on this gap and rationale, we develop a footprint-informed ecological
148 parameter migration framework to separate biogenic and anthropogenic CO₂ fluxes
149 from urban EC observations in Guangzhou, China. Guangzhou is a humid subtropical
150 megacity characterized by high population density, strong traffic activity, and
151 substantial urban vegetation, making it a representative testbed for partitioning CO₂
152 fluxes in mixed urban source areas. Our previous concentration-based study in
153 Guangzhou showed that urban CO₂ variability is jointly influenced by anthropogenic
154 emissions, biogenic exchange, and sea-land breeze transport, and that vegetation
155 uptake during the growing season can weaken the apparent fossil-fuel CO₂ signal
156 (Zhang et al., 2026). Compared with concentration observations, EC can provide a more
157 direct constraint on local surface-atmosphere exchange, but only if the biogenic



158 contribution within the flux footprint can be estimated and removed.

159 To address this challenge, we use paired EC observations from two sites in
160 Guangzhou in a donor–target design. The Conghua (CH) site is located in a highly
161 vegetated area with weak anthropogenic activity and is used as a relatively clean
162 ecological donor site. The Panyu (PY) site is located in a mixed urban source area
163 composed of vegetation, roads, buildings, and water bodies and is used as the urban
164 target site. We first constrain R_{eco} and GPP light-response parameters at CH, and then
165 migrate these ecological parameters to PY using dynamic flux footprints and footprint-
166 weighted EVI. After subtracting R_{eco} , GPP, and human respiration, we define the
167 remaining flux as F_{ff} , an operational estimate of footprint-scale anthropogenic CO_2 flux.
168 We then evaluate this interpretation using independent pollutant, source-area, traffic-
169 activity, and inventory-magnitude evidence, while also assessing the dominant sources
170 of uncertainty.

171 This study addresses three questions: (i) can footprint-weighted vegetation activity
172 provide an ecologically consistent constraint for migrating ecological parameters from
173 a relatively clean donor site to a heterogeneous urban target footprint; (ii) how do
174 partitioned biogenic and anthropogenic components jointly shape net CO_2 exchange
175 within a mixed urban footprint in a humid subtropical megacity; and (iii) do the
176 operationally defined F_{ff} estimates show temporal and source-area patterns consistent
177 with footprint-scale anthropogenic activity, provide useful context for inventory spatial
178 allocation, and remain robust under key uncertainty sources? By answering these
179 questions, this study provides a transferable framework for partitioning urban CO_2
180 source and sink components using conventional EC, remote sensing, and flux footprints
181 in cities without isotopic or multi-species flux measurements. It also offers footprint-
182 scale observational constraints for interpreting anthropogenic emissions, biogenic
183 modulation, and inventory spatial allocation in mixed urban source areas.

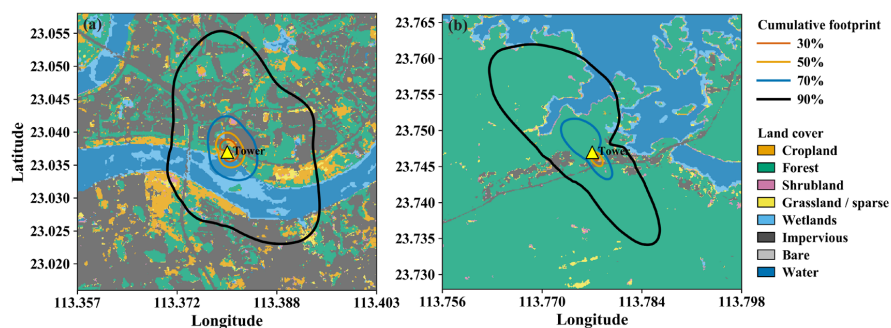
184 **2 Data and methods**

185 **2.1 Study sites**

186 This study used eddy covariance (EC) observations from two sites in Guangzhou, China:
187 Panyu (PY) and Conghua (CH). Both sites are part of an urban high-precision



188 greenhouse-gas monitoring network and are equipped with high-precision greenhouse-
189 gas concentration, CO₂ flux, and meteorological measurement systems. The site-
190 selection strategy and basic observational information have been reported by Zhang et
191 al. (2026). Guangzhou has a humid subtropical monsoon climate (according to the
192 Köppen climate classification), with a mean annual temperature of 23.7 °C and mean
193 annual precipitation of 1933.6 mm. The PY site (113.380° E, 23.037° N) is located in
194 the Guangzhou Higher Education Mega Center, where urban green space, roads,
195 buildings, water bodies, and dense human activity are closely interwoven. The CH site
196 (113.777° E, 23.747° N) is located near Liuxi River National Forest Park, where
197 population density is low and vegetation cover is high (Fig. 1). A two-dimensional flux-
198 footprint model centered on each tower and 10 m land-cover data show that the PY
199 footprint is composed of 46 % impervious surface, 44 % vegetation, and 10 %
200 water/wetland, with urban forest accounting for 60 % of the vegetated area (Kljun et
201 al., 2015; Zhang et al., 2025). The CH footprint is composed of 2 % impervious surface,
202 86 % vegetation, and 12 % water/wetland, with urban forest accounting for 96 % of the
203 vegetated area. Urban forests at both sites are dominated by evergreen broadleaf trees,
204 including Bauhinia, eucalyptus, kapok, and autumn maple. The key contrast between
205 the two sites is therefore not climate zone, but vegetation cover, urbanization intensity,
206 and anthropogenic-activity background.



207
208 **Figure 1.** Two-dimensional climatological footprint contours over land-cover classes at the
209 PY and CH sites. Panels (a) and (b) show the PY and CH sites, respectively. Background
210 land-cover classes were derived from the 2023 global 10 m land-cover product GLC_FCS10
211 (Zhang et al., 2025). The footprint model was evaluated on a 10 m spatial grid to match the
212 land-cover data. The overlaid contours represent cumulative footprint contributions of 30 %,
213 50 %, 70 %, and 90 %, and yellow triangles mark the tower locations.



214 Table S3 and Fig. S6 show that CH and PY have similar seasonal phases in
215 temperature and shortwave radiation, although PY has a warmer urban microclimatic
216 background and the month-to-month consistency in vapor pressure deficit (VPD) is
217 relatively weak. We therefore do not assume identical microclimatic conditions or
218 identical carbon-exchange magnitudes at the two sites. Rather, the site pair provides a
219 donor–target contrast under the same regional climate and broadly similar vegetation
220 background, with CH representing weaker anthropogenic disturbance and PY
221 representing a mixed urban source area.

222 **2.2 EC flux observations and quality control**

223 EC CO₂ fluxes were measured from tall tower platforms at both sites. At PY, the EC
224 system consisted of an open-path CO₂/H₂O analyzer (LI-7500DS, LI-COR Inc., USA)
225 and a three-dimensional sonic anemometer (WindMaster Pro, Gill Inc., UK), installed
226 at 48 m. At CH, an open-path IRGASON system (Campbell Scientific Inc., USA),
227 integrating a three-dimensional sonic anemometer and an infrared CO₂/H₂O analyzer,
228 was installed at 40 m. Based on a 10 m canopy-height product (Lang et al., 2023) and
229 vector building-height data (<https://lbs.amap.com>, last access: 25 May 2026), the mean
230 effective roughness-element heights within the PY and CH footprints were 13.1 and
231 15.2 m, respectively. The measurement heights at both sites exceeded twice the mean
232 roughness-element height (Z_H , m), supporting their representativeness for turbulent
233 fluxes over the study source areas (Rotach, 1999; Grimmond and Oke, 1999; Velasco
234 and Roth, 2010). The observation period extended from 1 January 2023 to 30 April
235 2025.

236 During the observation period, the EC systems recorded CO₂/H₂O concentration,
237 three-dimensional wind components, air temperature, relative humidity, and related
238 variables at 10 Hz. Half-hourly fluxes were calculated from the covariance between
239 vertical wind speed and CO₂ mole fraction. Flux processing followed standard EC
240 procedures, including Webb-Pearman-Leuning (WPL) density correction, double
241 coordinate rotation, and 30 min detrending (Webb et al., 1980; Sabbatini et al., 2018).

242 To ensure data reliability, we applied a stepwise quality-control procedure.



Records flagged for extreme weather, instrument maintenance, or obvious malfunction were removed. Low-turbulence nighttime periods were filtered using the friction velocity (u^*) threshold method implemented in REddyProc (Papale et al., 2006; Wutzler et al., 2018). Non-stationary fluxes were excluded using the stationarity test of Foken et al. (2004), and despiking was then applied. After quality control, the valid-data fractions for net CO_2 fluxes were 51.7 % at PY and 54.5 % at CH, comparable to common ranges in urban EC studies (Matthews and Schume, 2022). The detailed quality-control procedure and retention rates are given in Sect. S1. The net fluxes were subsequently corrected for storage following the procedure described in Sect. S2 (Aubinet et al., 2001; Kooijmans et al., 2017; Kohonen et al., 2020; Soininen et al., 2025). The daily mean storage fluxes were 0.03 and 0.04 $\mu\text{mol m}^{-2} \text{s}^{-1}$ at PY and CH, respectively, much smaller than the measured fluxes. All subsequent analyses therefore used storage-corrected net CO_2 fluxes. Unless otherwise stated, storage-corrected fluxes, footprint calculations, ecological parameter fitting, parameter migration, and F_{ff} propagation were performed at the native 30 min time step. These results were aggregated to hourly means for diurnal and cross-variable analyses, and further summarized at daily, monthly, or seasonal scales where needed.

2.3 CO_2 flux partitioning framework

The net CO_2 flux observed by an urban EC system integrates multiple source and sink processes within the footprint, including fossil-fuel emissions, human respiration, ecosystem respiration, biogenic uptake, and changes in air-column storage (Wu et al., 2022; Soininen et al., 2025; Hilland et al., 2025). The storage-corrected flux is defined as:

$$F_{cr}(t) = F_{mz}(t) + F_{st}(t) \quad (1)$$

Here F_{cr} is the storage-corrected CO_2 flux ($\mu\text{mol m}^{-2} \text{s}^{-1}$), F_{mz} is the measured net CO_2 flux after quality control ($\mu\text{mol m}^{-2} \text{s}^{-1}$), and F_{st} is the storage flux ($\mu\text{mol m}^{-2} \text{s}^{-1}$). F_{cr} was then partitioned into the following components:

$$F_{cr}(t) = F_{ff}(t) + F_{hu}(t) + R_{eco}(t) - GPP(t) \quad (2)$$

Here F_{ff} is operationally defined as the anthropogenic CO_2 flux remaining after



272 subtracting ecosystem respiration (R_{eco}), gross primary productivity (GPP), and human
273 respiration (F_{hu}) from the storage-corrected net flux. F_{hu} is the human-respiration flux,
274 R_{eco} is the sum of plant and soil respiration, and GPP is the amount of CO_2 fixed by
275 photosynthesis. Within the PY footprint, direct fossil-fuel emissions are expected to be
276 dominated by road traffic and related urban activities. Because buildings within the
277 source area mainly rely on externally supplied electricity, on-site building-energy
278 combustion was not explicitly considered (Yang et al., 2025). F_{hu} was estimated from
279 population density and per-capita respiration rate (Sect. S3).

280 **2.4 Ecological flux parameterization at the donor site**

281 Ecological parameters for R_{eco} and GPP were fitted at CH and then migrated to PY as
282 described in Sect. 2.5. The donor-site ecological parameterization followed a two-step
283 framework in which nighttime respiration was fitted first and daytime GPP was then
284 inferred from the light response. This approach has been widely used for partitioning
285 EC-based NEE into ecosystem respiration and photosynthetic uptake (Reichstein et al.,
286 2005; Lasslop et al., 2010). Nighttime ecosystem respiration ($PAR < 10 \mu mol m^{-2} s^{-1}$)
287 was represented using a simplified Q10 temperature-response equation:

$$288 \quad R_{eco}(t) = R_c(t) Q_{10}^{T_a(t)/10} \quad (3)$$

289 Here T_a is the driving air temperature ($^{\circ}C$), Q_{10} is the temperature-sensitivity
290 coefficient and represents the relative increase in respiration rate for a $10^{\circ}C$
291 temperature increase, and R_c is the base level of respiration ($\mu mol m^{-2} s^{-1}$). Because
292 daytime ecosystem respiration may be suppressed by light (Wehr et al., 2016; Keenan
293 et al., 2019), daytime respiration was further represented as:

$$294 \quad R_{eco,day}(t) = R_{eco}(t) \left[1 - \beta \frac{PAR(t)}{K_L + PAR(t)} \right] \quad (4)$$

295 Here $R_{eco,day}$ is the light-modified daytime ecosystem respiration ($\mu mol m^{-2} s^{-1}$), β
296 is the maximum inhibition fraction, K_L is the light-inhibition half-saturation constant
297 ($\mu mol photon m^{-2} s^{-1}$), and PAR is photosynthetically active radiation ($\mu mol photon$
298 $m^{-2} s^{-1}$). Because PAR was not measured directly, it was converted from shortwave
299 radiation (Sects. S4 and S7). The baseline setting used $\beta = 0.15$ and $K_L = 600 \mu mol$



300 photon $\text{m}^{-2} \text{s}^{-1}$. A weak daytime-respiration-inhibition scenario, with a lower β (0.10)
301 and a higher K_L ($800 \mu\text{mol photon m}^{-2} \text{s}^{-1}$), was used to evaluate the influence of this
302 assumption on ecological-parameter fitting and F_{ff} estimation.

303 Daytime GPP was represented using a non-rectangular hyperbolic saturating light-
304 response equation:

$$305 \quad GPP(t) = \frac{\alpha PAR(t) + P_{max} - \sqrt{(\alpha PAR(t) + P_{max})^2 - 4\theta P_{max} \alpha PAR(t)}}{2\theta} \quad (5)$$

306 Here α is the initial light-use efficiency, defined as the initial slope of the GPP
307 response to PAR under low-light conditions; P_{max} is the maximum photosynthetic
308 capacity ($\mu\text{mol m}^{-2} \text{s}^{-1}$); and θ is the curvature parameter controlling the transition from
309 the low-light linear segment to the high-light saturation segment. We then used three
310 GPP model structures to evaluate how ecological structural assumptions affect CH-side
311 GPP fitting and PY-side F_{ff} estimation:

312 (1) Pure saturating structure: the non-rectangular hyperbolic form above was used
313 directly, without temperature stratification or VPD inhibition. All samples satisfying the
314 GPP-fitting criteria were fitted within a single parameter group for α , P_{max} , and θ .

315 (2) Saturating structure with VPD inhibition: VPD inhibition of maximum
316 photosynthetic capacity was introduced into the non-rectangular hyperbolic form to test
317 the influence of atmospheric dryness stress (Lasslop et al., 2010). Specifically, P_{max} was
318 replaced by an effective maximum photosynthetic capacity regulated by VPD:

$$319 \quad P_{max,VPD} = P_{max} \times f_{VPD} \quad (6)$$

$$320 \quad f_{VPD} = \exp[-k_{VPD} \max(0, VPD - VPD_0)] \quad (7)$$

321 Here $P_{max,VPD}$ is the VPD-regulated effective maximum photosynthetic capacity
322 ($\mu\text{mol m}^{-2} \text{s}^{-1}$), f_{VPD} is the VPD inhibition factor, VPD is vapor pressure deficit (kPa),
323 k_{VPD} is the VPD inhibition coefficient (kPa^{-1}), and VPD_0 is the reference threshold
324 above which inhibition is applied; VPD_0 was set to 1.0 kPa. No additional inhibition
325 was applied when VPD was below 1.0 kPa. Above this threshold, maximum
326 photosynthetic capacity declined exponentially with increasing VPD. VPD calculation
327 is described in Sect. S4.

328 (3) Saturating structure with temperature stratification: instead of adding a



continuous temperature-response function to the GPP equation, GPP-fitting samples were grouped by air temperature (T_a), and α , $P_{\max}$, and θ were fitted separately within each temperature group. The baseline analysis used Scheme A, in which the CH-side GPP-fitting samples were divided into low-, mid-, and high-temperature groups according to the tertiles of T_a :

$$q_1 = \text{quantile}(T_a, 1/3) \quad (8)$$

$$q_2 = \text{quantile}(T_a, 2/3) \quad (9)$$

Here T_a is air temperature ($^{\circ}\text{C}$), and q_1 and q_2 are the one-third and two-third quantiles of air temperature in the fitting samples. The low-temperature group is $T_a < q_1$, the mid-temperature group is $q_1 \leq T_a < q_2$, and the high-temperature group is $T_a \geq q_2$. This treatment represents differences in light-response parameters under different thermal states while avoiding an additional continuous temperature-response function under limited data and complex urban source-area conditions. To test the sensitivity to temperature-boundary selection, we used Scheme B as a structural sensitivity scenario. Scheme B kept the equations, fitting windows, and screening criteria unchanged, but expanded the mid-temperature group by 0.5°C toward both the low- and high-temperature sides.

To reduce short-term sampling variability and avoid non-physical jumps between adjacent parameter windows, ecological parameters fitted from 11 d subsets were smoothed with an 11 d moving average, following the urban green-space flux-partitioning approach of Soininen et al. (2025). The smoothed R_e , α , and $P_{\max}$ were used to estimate R_{eco} and GPP at continuous half-hourly resolution and for subsequent parameter migration, whereas Q_{10} and θ were kept fixed over the full period.

We used the saturating structure with temperature stratification as the baseline GPP scheme, and evaluated its CH-side fitting performance in Sect. 3.2.1. The pure saturating structure, the saturating structure with VPD inhibition, and Scheme B were retained as structural sensitivity experiments.

2.5 Footprint-informed ecological parameter migration

CH-to-PY migration was designed as footprint- and vegetation-constrained parameter



358 migration, not as a mechanical transfer of CH observed fluxes to PY. The CH-fitted
359 parameters are interpreted as a regional ecological-response reference under weak
360 anthropogenic disturbance, while PY biogenic fluxes are driven by PY-specific
361 temperature, radiation, VPD, and footprint-weighted EVI. This treatment follows the
362 view that EC light-response parameters are apparent ecosystem- or canopy-scale
363 quantities whose magnitudes depend on canopy structure, vegetation amount,
364 phenology, and environmental conditions (Sellers, 1985; Myneni and Williams, 1994;
365 Huete et al., 2002). It is also consistent with light-use-efficiency and satellite-driven
366 GPP models that estimate photosynthetic uptake using vegetation indices and
367 environmental scalars (Monteith, 1972; Running et al., 2004; Xiao et al., 2004;
368 Mahadevan et al., 2008).

369 For GPP-related parameters, EVI_{fp} provides a direct constraint on
370 photosynthetically active vegetation amount and canopy activity. For R_{eco} , EVI_{fp} has a
371 different role: it acts as a footprint-scale vegetation/pervious-surface mask and a
372 capacity-normalization factor for vegetation-associated respiration. Impervious
373 surfaces within the footprint are therefore not assumed to contribute the same
374 respiratory capacity as vegetated surfaces. Under the shared regional climate and
375 broadly similar evergreen broadleaf vegetation background of CH and PY, the CH-
376 derived respiration capacity per unit EVI_{fp} is used as a first-order reference for patchy
377 urban green spaces at PY. This does not imply that soil moisture, substrate supply,
378 irrigation, management intensity, or urban heat-island effects are fully captured by EVI;
379 these factors are treated as part of the structural boundary of the donor–target migration
380 framework and are considered when interpreting the sensitivity results.

381 Because the EC source area changes at half-hourly scales with wind direction,
382 stability, and turbulence conditions, land-cover fractions or vegetation indices within a
383 fixed buffer cannot represent the actual contributing source area of each flux
384 observation. We therefore calculated two-dimensional flux-footprint weights for each
385 time step and used these weights to construct footprint-weighted land-cover fractions
386 (LD_{fp}) and footprint-weighted EVI (EVI_{fp}) following the procedure described in Sect.
387 S5. Land-cover data were taken from the global 10 m GLC_FCS10 product (Zhang et



al., 2025). EVI was obtained from the Harmonized Landsat Sentinel-2 (HLS) vegetation-index products, with 30 m spatial resolution and a revisit cycle of approximately 2–3 d (Neigh, 2021; Neigh, 2025). Compared with the Normalized Difference Vegetation Index (NDVI), EVI is less prone to saturation in high-biomass areas and is more stably related to canopy greenness, the fraction of absorbed photosynthetically active radiation (fAPAR), and seasonal GPP dynamics (Huete et al., 2002; Xiao et al., 2004).

Operationally, we treated the CH temperature-stratified ecological response as the reference function and allowed amplitude-type parameters to vary with vegetation amount and activity in the target footprint. The scaling was applied to ecological capacity parameters rather than to observed fluxes; therefore, migrated PY GPP and R_{eco} were calculated using PY-specific radiation, temperature, VPD, and dynamic EVI_{fp} . Capacity-type parameters fitted at CH within temperature group g , including α , P_{max} , and R_e , were therefore normalized by CH EVI_{fp} :

$$p_{norm,CH}(g) = \frac{P_{CH}(g)}{EVI_{fp,CH}(g)} \quad (10)$$

Here g denotes the temperature group (low, mid, or high), $P_{CH}(g)$ is the ecological parameter fitted at CH within temperature group g , and $P_{norm,CH}(g)$ is the parameter normalized by CH EVI_{fp} . The median $P_{norm,CH}(g)$ within each temperature group was then used as the CH-to-PY parameter-migration coefficient to reduce the influence of anomalous fitting windows or extreme parameter values:

$$C_{norm,CH}(g) = median[p_{norm,CH}(g)] \quad (11)$$

$C_{norm,CH}(g)$ is the CH-to-PY parameter-migration coefficient for temperature group g . The corresponding PY EVI_{fp} was then used to reconstruct ecological parameters at PY:

$$P_{PY}(t) = C_{norm,CH}(g(t)) \times EVI_{fp,PY}(t) \quad (12)$$

Here $P_{PY}(t)$ is the migrated ecological parameter at PY for time t . For non-amplitude parameters that are not suitable for EVI proportional scaling, the representative CH statistic within the same temperature group was used. This treatment



preserves the response structure of CH parameters to thermal state while allowing parameter magnitudes to adjust with vegetation activity within the actual PY footprint. Time-step-resolved LD_{fp} , EVI_{fp} , and EVI_{fp} -based migrated parameters together provide the source-area constraint for CH-to-PY parameter migration.

2.6 Sensitivity-based uncertainty assessment

To test the robustness of PY F_{ff} to parameter propagation, processing choices, and ecological model structure, we constructed a targeted sensitivity-based uncertainty framework. This framework was not intended to provide a complete probability distribution for all error sources, nor were the scenarios treated as equally probable samples. Instead, controlled perturbations were applied to processing steps with clear physical meaning to identify the dominant sources of F_{ff} uncertainty.

Uncertainty propagation included two levels. The first level was CH-side ecological-parameter bootstrapping: for each CH scenario, respiration and light-response parameters were resampled 2000 times and migrated to PY using EVI_{fp} , yielding $q05$, $q50$, and $q95$ estimates of R_{eco} , GPP, and F_{ff} . The second level was scenario sensitivity: single-factor or structural-alternative scenarios were constructed for the u^* threshold, storage flux, EVI time window, GPP model structure, temperature grouping, daytime-respiration inhibition, CH-side shortwave-radiation input, and human-respiration emissions, and were compared with the baseline scheme.

The final propagation ensemble contained 14 CH-side scenarios and 3 human-respiration scenarios, forming 42 propagation combinations (Sect. S6). CH0 is the baseline scheme. The CH-side scenarios evaluate input and processing sensitivity, ecological model-structure choices, parameter-migration settings, and CH-side shortwave-radiation representativeness. Human-respiration emissions were represented by base, low50, and high50 scenarios, with low50 and high50 equal to 0.5 and 1.5 times the baseline human-respiration emission, respectively.

For any CH scenario s , human-respiration scenario r , bootstrap member b , and time t , F_{ff} was propagated as:

$$F_{ff}(b, s, r, t) = F_{cr}(s, t) - R_{eco}(b, s, t) + GPP(b, s, t) - F_{hu}(r, t) \quad (13)$$



Here $F_{ff}(b, s, r, t)$ is the propagated F_{ff} estimate for the corresponding scenario and bootstrap member; $F_{cr}(s, t)$ is the storage-corrected net CO_2 flux at PY under scenario s ; $R_{eco}(b, s, t)$ is ecosystem respiration driven by migrated parameters; $GPP(b, s, t)$ is gross primary productivity driven by migrated parameters; and $F_{hu}(r, t)$ is the human-respiration CO_2 flux.

The central estimate at each time was the bootstrap median, and the parameter-propagation range was represented by q05–q95. Scenario uncertainty was evaluated as the difference from the baseline CH0 + human base configuration:

$$\Delta F_{ff}(s, t) = F_{ff}(s, base, t) - F_{ff}(CH0, base, t) \quad (14)$$

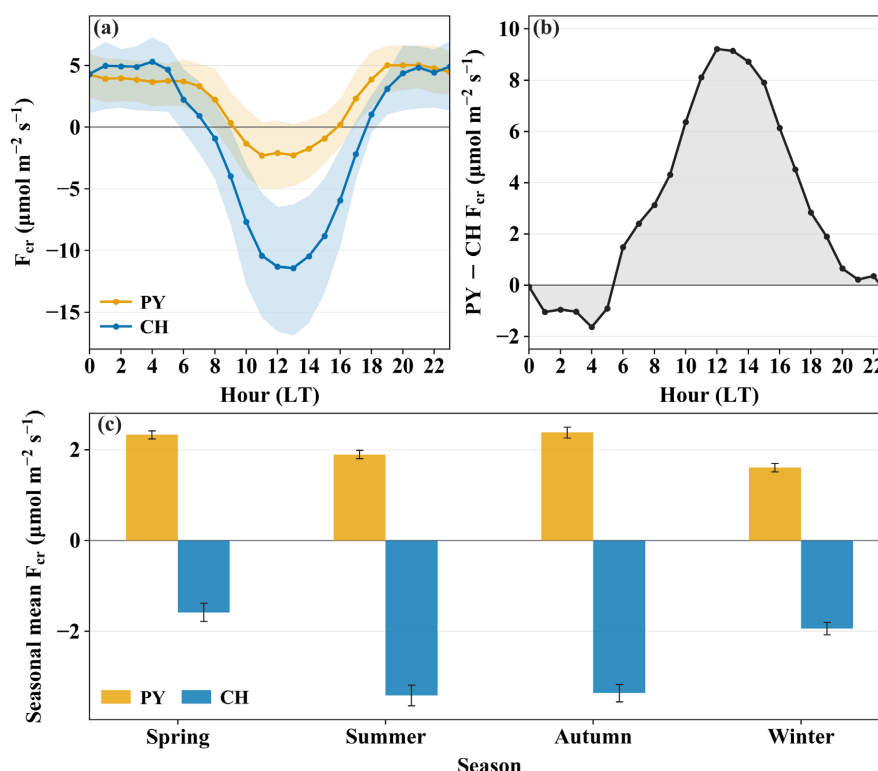
Here $F_{ff}(s, base, t)$ denotes the bootstrap-median F_{ff} under CH scenario s and the base human-respiration setting. $\Delta F_{ff}(s, t)$ denotes the F_{ff} offset for scenario s relative to baseline CH0 at time t . Positive values indicate that the scenario increases F_{ff} , whereas negative values indicate that it decreases F_{ff} . ΔF_{ff} was then summarized at overall, hourly, and monthly scales to diagnose how different uncertainty sources affect the mean, diurnal variation, and seasonal variation of F_{ff} .

3 Results and discussion

3.1 Net CO_2 flux variability and source-area controls

3.1.1 Diurnal and seasonal patterns of net CO_2 flux

Hourly storage-corrected CO_2 fluxes (F_{cr}) during 2023–2024 showed a clear urban–suburban contrast (Fig. 2a–b). Overall, the mean F_{cr} at PY was $2.04 \mu mol m^{-2} s^{-1}$, indicating a year-round net CO_2 source, whereas the mean F_{cr} at CH was $-2.57 \mu mol m^{-2} s^{-1}$, indicating a year-round net CO_2 sink. Urban EC studies generally show that urban surfaces are net CO_2 sources, with anthropogenic emissions, especially traffic, building-energy use, and residential activities, providing important contributions to urban CO_2 fluxes. By contrast, suburban or highly vegetated areas are more likely to show pronounced biogenic uptake signals (Crawford et al., 2011; Järvi et al., 2012). The annual mean difference between PY and CH therefore reflects the control exerted by source-area urbanization and vegetation cover on net CO_2 exchange.



473

474 **Figure 2.** Diurnal and seasonal characteristics of corrected CO₂ flux (F_{cr}) at the PY and CH
475 sites. **(a)** Mean diurnal F_{cr} at PY and CH; shading indicates the interquartile range (Q25–Q75).
476 **(b)** Mean diurnal PY–CH contrast in F_{cr} . **(c)** Seasonal mean F_{cr} at PY and CH for spring (March–
477 May), summer (June–August), autumn (September–November), and winter (December–
478 February), with error bars indicating the standard error. Positive values indicate net CO₂ release
479 to the atmosphere, whereas negative values indicate net uptake.

480 At the diurnal scale, both sites showed higher nighttime and lower daytime fluxes,
481 but with very different amplitudes. At CH, F_{cr} was mostly positive at night, became
482 rapidly negative as radiation increased during the day, and reached the strongest uptake
483 around 12:00–13:00, with a mean of about $-11 \mu\text{mol m}^{-2} \text{s}^{-1}$. At PY, daytime F_{cr} also
484 decreased, but only approached weakly negative or near-neutral values around midday.
485 This suggests that vegetation uptake was still detectable at PY, but its expression in the
486 net flux was partly masked by persistent urban anthropogenic emissions. Similar
487 behavior has been reported for a tropical residential neighborhood in Singapore and
488 other urban EC studies: urban vegetation can absorb part of the daytime CO₂, but it



usually does not fully offset anthropogenic emissions, so urban sites generally remain net sources (Velasco et al., 2013; Velasco and Roth, 2010).

At the seasonal scale, mean F_{cr} values at PY were 2.33, 1.89, 2.38, and 1.61 $\mu\text{mol m}^{-2} \text{s}^{-1}$ in spring, summer, autumn, and winter, respectively (Fig. 2c). PY therefore remained a net CO_2 source throughout the year, although net emissions were lower in summer. At CH, F_{cr} was negative in all seasons, with spring, summer, autumn, and winter means of -1.59 , -3.41 , -3.36 , and $-1.94 \mu\text{mol m}^{-2} \text{s}^{-1}$, respectively. Stronger uptake during the warm growing-season months is consistent with the long summer carbon-sink duration observed in subtropical urban vegetation and the seasonally active photosynthesis of South China subtropical evergreen forests (Zhang et al., 2019; Wang et al., 2023). Multi-year urban and suburban EC observations also show that sites with higher vegetation cover usually exhibit stronger growing-season CO_2 uptake, whereas seasonal variation at urban sites is jointly controlled by anthropogenic emissions and biogenic processes (Järvi et al., 2012; Menzer and McFadden, 2017).

Seasonal diurnal patterns further show that biogenic regulation of net flux is stronger during the growing season (Fig. S1). CH exhibited the largest diurnal amplitude and strongest midday uptake in summer, consistent with high-vegetation sites having both stronger ecosystem respiration and stronger photosynthetic uptake during the growing season. Although PY remained a net CO_2 source throughout the year, its diurnal amplitude in summer and autumn was also higher than in spring and winter, indicating that clear biogenic modulation remains embedded in the net flux of this mixed urban source area. In other words, low daytime F_{cr} and enhanced growing-season amplitude at PY should not be interpreted as weaker anthropogenic emissions, but as the net result of anthropogenic emissions and vegetation uptake acting together.

3.1.2 Footprint land cover and sectoral controls

Because urban EC source areas vary with wind direction, turbulence state, and boundary-layer conditions, footprint modelling combined with high-resolution land-cover data is needed to interpret spatial heterogeneity in urban CO_2 fluxes (Crawford and Christen, 2015; Kljun et al., 2015).



Wind-sector analysis showed that F_{cr} at PY had clear source-area dependence (Fig. 3). High F_{cr} mainly occurred in the NE, N, and NW sectors, with mean values of about 3.59, 2.83, and 2.52 $\mu\text{mol m}^{-2} \text{s}^{-1}$, respectively, clearly higher than those in the W and SW sectors. Correspondingly, impervious fractions were relatively high in the N and NW sectors, about 33 % and 37 %, respectively. Although the NE impervious fraction was slightly lower than those in the N and NW sectors, the NE sector still contained a strong mixture of urban hard surfaces and forest patches. Correlations across the eight wind-sector means showed that sector-mean F_{cr} at PY was strongly positively correlated with impervious fraction ($r = 0.77$). F_{cr} was also positively correlated with forest fraction ($r = 0.52$), while impervious and forest fractions were themselves positively correlated ($r = 0.61$). This result indicates that the positive forest- F_{cr} correlation at PY should not be interpreted as forest cover increasing CO_2 emissions. It more likely reflects collinearity caused by the spatial coexistence of campus forests, roads, buildings, and hard surfaces in and around the PY university-town area. In other words, forest patches at PY are not independent biogenic source areas; they are often embedded within urbanized source areas and therefore carry part of the urban-activity source-area signal at the sector scale.

By contrast, the directional pattern at CH was more consistent with biogenic control. F_{cr} was mostly negative across CH sectors, with the strongest net uptake in the W, SW, and S sectors, where mean F_{cr} values were about -7.44 , -6.74 , and -6.76 $\mu\text{mol m}^{-2} \text{s}^{-1}$, respectively; these sectors generally had higher forest fractions. Sector-scale correlations showed a negative relationship between F_{cr} and forest fraction at CH ($r = -0.76$), indicating that source areas with higher forest cover generally corresponded to stronger net CO_2 uptake. The comparison between CH and PY therefore demonstrates that the influence of land cover on F_{cr} is site dependent. At CH, forest cover is closer to a true biogenic control. At PY, forest, impervious surface, and human-activity source areas are tightly coupled, so correlation and causality must be separated carefully. Similarly, Pérez-Ruiz et al. (2020) showed that different urban land-cover types influence the response of CO_2 fluxes to environmental drivers, highlighting land-cover heterogeneity as an important factor in explaining flux differences within cities.

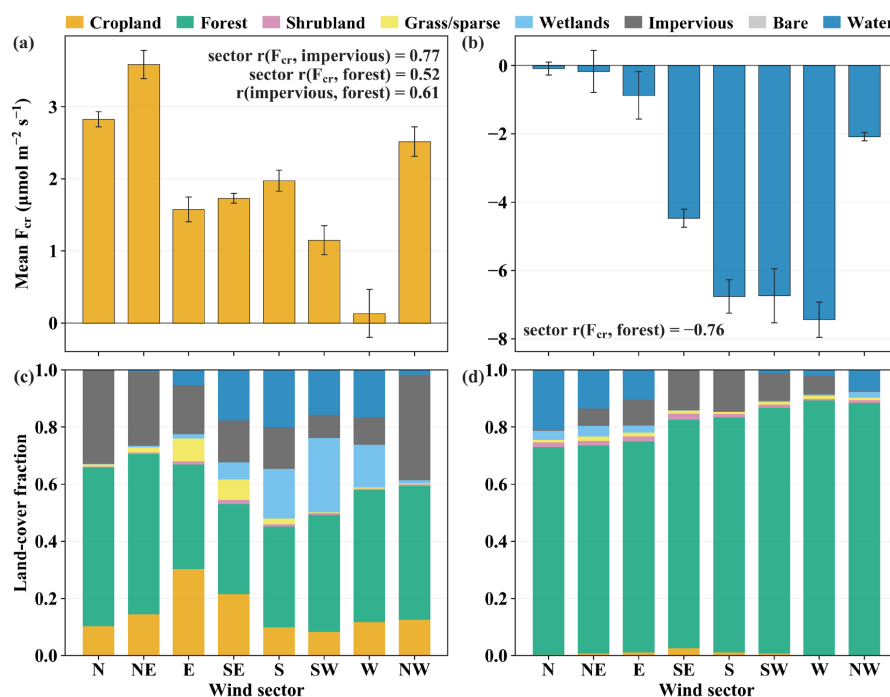


Figure 3. Directional variation in corrected CO₂ flux (F_{cr}) and footprint-weighted land-cover composition at the PY and CH sites. Panels (a) and (b) show sectoral mean F_{cr} at PY and CH, respectively; error bars indicate the standard error across hourly values. Panels (c) and (d) show the corresponding footprint-weighted land-cover fractions for PY and CH, respectively. The eight sectors were defined as 45° wind-direction bins centered on the cardinal and intercardinal directions, with N spanning 337.5°–360° and 0°–22.5°; land-cover fractions were calculated by averaging the footprint-weighted land-cover composition of all valid records within each sector.

Sector-hour heat maps and hourly correlation analyses further support this interpretation (Figs. S2 and S3). High F_{cr} at PY was concentrated not only in more urbanized directions such as N, NE, and NW, but also during nighttime, evening, and morning-rush periods, indicating that its diurnal variation was jointly regulated by wind-sector source area, local activity, and biogenic uptake. Hourly correlations also showed a positive relationship between F_{cr} and impervious fraction at PY, suggesting that hard surfaces and built-up source areas are important indicators of higher net CO₂ emissions. F_{cr} at PY was also positively correlated with forest fraction, but this likely reflects the co-occurrence of urban green space with roads and buildings, given their sectoral collinearity with impervious surfaces (Fig. 3). At CH, most sectors became



566 clearly negative during the day, and F_{cr} was negatively correlated with forest fraction,
567 more closely reflecting biogenic uptake control in a highly vegetated source area.
568 Beyond land cover, F_{cr} at both sites was significantly negatively correlated with
569 shortwave radiation, showing that enhanced daytime radiation and vegetation
570 photosynthesis were important drivers of lower net CO_2 flux. Correlations of F_{cr} with
571 temperature, VPD, relative humidity, and wind speed more likely reflect the combined
572 effects of diurnal phase, turbulent mixing, and biogenic processes, and should not be
573 interpreted as independent causal effects of single variables.

574 Overall, source-area analysis shows that F_{cr} at PY is jointly shaped by mixed urban
575 land cover, especially the co-occurrence of impervious surfaces and campus green
576 space, together with wind-sector source areas and environmental conditions, whereas
577 F_{cr} at CH mainly reflects biogenic uptake control by high forest cover. F_{cr} can therefore
578 reveal the spatiotemporal heterogeneity of urban net CO_2 exchange, but it cannot
579 directly represent anthropogenic CO_2 flux. For mixed urban source areas such as PY,
580 where green space and built-up land are highly interwoven, R_{eco} , GPP, and human
581 respiration must be removed before anthropogenic CO_2 flux can be more robustly
582 extracted.

583 **3.2 Footprint-informed ecological parameterization**

584 **3.2.1 Donor-site respiration and GPP light-response fitting**

585 Having established CH as a highly vegetated and weakly disturbed donor site, we next
586 evaluated whether its respiration and GPP light-response parameters could provide a
587 stable ecological reference for parameter migration. This step tests whether the donor-
588 site fitting produced physiologically interpretable parameters and whether the selected
589 GPP structure adequately captured CH-side photosynthetic variability.

590 In the baseline scheme, nighttime ecosystem respiration was represented using a
591 fixed Q_{10} value of 1.64. Seasonal variation was therefore expressed mainly through the
592 fitted base respiration coefficient R_e , rather than Q_{10} itself. Nighttime respiration fitting
593 at CH showed acceptable agreement between observations and simulations, with $R =$
594 0.608 , $RMSE = 1.419 \mu mol m^{-2} s^{-1}$, and $bias = -0.004 \mu mol m^{-2} s^{-1}$ (Fig. S4a). Thus,



595 under the present data-screening and parameterization framework, the nighttime
596 respiration model showed no obvious systematic overestimation or underestimation and
597 provided a basis for subsequent daytime respiration extrapolation and parameter
598 migration.

599 CH-side GPP light-response fitting further showed that the temperature-stratified
600 scheme captured photosynthetic variation reasonably well. The baseline GPP fit had R
601 $= 0.659$, $RMSE = 4.904 \mu\text{mol m}^{-2} \text{s}^{-1}$, and $MAE = 3.679 \mu\text{mol m}^{-2} \text{s}^{-1}$, outperforming
602 the pure saturating structure and the saturating structure with VPD inhibition (Fig. S4b–
603 d). The ecological parameters in the CH baseline scheme were not constant; they varied
604 over time and among temperature groups (Fig. S5). R_c fluctuated during the year, with
605 monthly medians of about 0.96–1.36. Among the GPP light-response parameters, α and
606 P_{max} also varied with time and thermal state, with monthly medians of about 0.026–
607 0.052 and 16.8–43.8 $\mu\text{mol m}^{-2} \text{s}^{-1}$, respectively. These results indicate interpretable
608 ecological variability in CH respiration and light-response parameters, rather than
609 dominance by random fitting error. θ was treated as a fixed shape parameter in the
610 baseline scheme, with a value of 0.2, and is therefore not interpreted as an independent
611 seasonal parameter.

612 Comparison among GPP structural schemes further supports the temperature-
613 stratified structure as the baseline. Compared with the pure saturating structure and the
614 saturating structure with VPD inhibition, the temperature-stratified scheme had a higher
615 R and lower RMSE and MAE, indicating that it better captured hourly variation in the
616 CH GPP target. In the saturating + VPD inhibition structure, the fitted k_{VPD} was close
617 to zero. This suggests that, under the humid subtropical background and screened
618 samples used here, the explicit VPD inhibition term did not provide additional
619 explanatory power beyond the temperature-stratified structure. This result does not
620 deny the physiological relevance of VPD; rather, it indicates that differences in thermal
621 state better explained light-response variation in this dataset. We therefore selected the
622 temperature-stratified structure as the baseline and used the pure saturating and
623 saturating + VPD inhibition structures as structural sensitivity scenarios in the
624 uncertainty analysis.



625 3.2.2 EVI_{fp}-constrained parameter migration to the urban target

626 We evaluated whether the donor–target design met the ecological comparability
627 conditions required for EVI_{fp}-constrained parameter migration. This evaluation focused
628 on three aspects: whether CH could provide a relatively clean ecological-response
629 reference, whether CH and PY shared a comparable regional environmental and
630 phenological background, and whether vegetation amount and activity in the PY
631 footprint could be represented by dynamic EVI_{fp}. The footprint land-cover contrast
632 shown in Fig. 3c–d and Table S3 supports CH as a vegetation-dominated donor and PY
633 as a heterogeneous urban target, indicating that PY cannot directly use CH observed
634 fluxes or unscaled CH parameters.

635 Environmental and vegetation diagnostics further support this constrained-
636 migration design. Figure S6 shows that CH and PY have highly consistent seasonal
637 phases in air temperature and daytime shortwave radiation. VPD has a similar seasonal
638 magnitude at the two sites, but its month-to-month consistency is relatively weak,
639 indicating shared regional seasonal forcing but non-identical microclimatic conditions.
640 This microclimatic difference does not invalidate the migration framework because CH
641 meteorological time series were not imposed on PY. Instead, the migrated PY biogenic
642 fluxes were driven by PY-specific temperature, radiation, and VPD, while CH provided
643 the vegetation-normalized ecological-response reference. Table S3 also shows broadly
644 similar urban-forest types at the two sites, whereas vegetation cover and anthropogenic
645 influence differ substantially. Consistently, Fig. 4 shows that EVI_{fp} is overall higher at
646 CH than at PY, as expected from the higher vegetation cover at CH. Although EVI_{fp} at
647 PY is lower, it retains a clear seasonal pattern, with higher values in summer and lower
648 values in winter and spring. This recognizable phenological signal supports the use of
649 EVI_{fp} for scaling ecological capacity in the dynamic PY footprint. For R_{eco}, EVI_{fp}
650 should be interpreted as scaling vegetation-associated respiratory capacity within the
651 dynamic footprint, rather than assuming that all urban surfaces or soil processes respond
652 linearly to canopy greenness. Thus, the similar seasonal phases in temperature and
653 radiation support CH as a same-region donor, whereas PY's warmer background and



weaker VPD consistency define the extrapolation boundary.

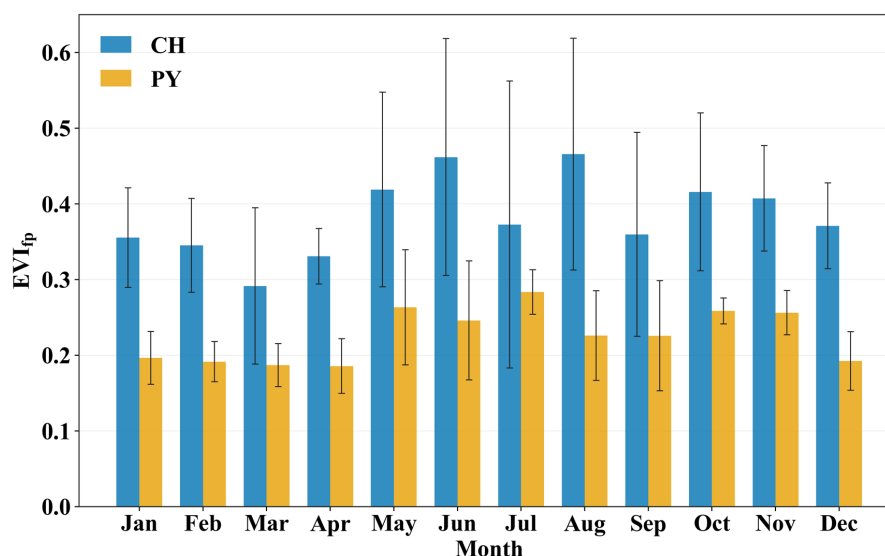


Figure 4. Monthly variation of footprint-weighted vegetation activity at the CH and PY sites. Bars show the monthly mean footprint-weighted enhanced vegetation index (EVI_{fp}) for CH and PY, and error bars indicate ± 1 standard deviation.

The migrated PY ecological-parameter ranges are consistent with these diagnostics. Under the baseline scheme, the 5 %–95 % range of α at PY was 0.007–0.029, the 5 %–95 % range of $P_{\max}$ was 5.30–13.89 $\mu\text{mol m}^{-2} \text{s}^{-1}$, and the 5 %–95 % range of R_c was 0.351–0.762. Compared with the CH donor site, the capacity-type parameters at PY were substantially reduced but did not approach zero, consistent with the presence of urban green space in the PY footprint and with lower vegetation cover and activity than at CH. All migrated parameters were non-negative. Migrated GPP showed a typical daytime increase and approached zero at night, whereas R_{eco} remained positive throughout the day and varied more smoothly (Fig. 5). Recent urban flux-partitioning work provides the closest benchmark for the migrated PY biogenic fluxes. Soininen et al. (2025) reported median GPP peaks of about 8.8–11.9 $\mu\text{mol m}^{-2} \text{s}^{-1}$ for a street-sector footprint with substantial vegetation cover, together with a mean ecosystem respiration of about 3.9 $\mu\text{mol m}^{-2} \text{s}^{-1}$. The migrated PY maximum GPP of 10.83 $\mu\text{mol m}^{-2} \text{s}^{-1}$ is therefore comparable to this urban mixed-source GPP benchmark. For R_{eco} , additional magnitude support comes from urban and subtropical vegetated



674 surfaces. Soil respiration ranged from 2.6 to 6.7 $\mu\text{mol m}^{-2} \text{s}^{-1}$ across vegetated and
675 managed cover types in greater Boston (Decina et al., 2016), and annual mean soil
676 respiration in Hefei park and campus green spaces was 2.51–2.64 $\mu\text{mol m}^{-2} \text{s}^{-1}$ (Tao et
677 al., 2016). Eddy-covariance flux partitioning in a subtropical Chinese urban forest
678 reported mean R_{eco} of about 3.0 $\mu\text{mol m}^{-2} \text{s}^{-1}$, with condition-specific means ranging
679 from 1.9 to 4.0 $\mu\text{mol m}^{-2} \text{s}^{-1}$ under heatwave and drought conditions (Yang et al., 2025).
680 Humid subtropical forests in South China also show comparable total soil-respiration
681 magnitudes of about 2.7–3.0 $\mu\text{mol m}^{-2} \text{s}^{-1}$ after unit conversion (Yi et al., 2007). The
682 migrated PY maximum R_{eco} of 4.54 $\mu\text{mol m}^{-2} \text{s}^{-1}$ therefore falls within a plausible range
683 for vegetated and managed pervious surfaces. This comparison is a magnitude-
684 consistency check rather than direct validation, but it supports the view that the EVI_{fp}
685 constraint did not generate unrealistically large biogenic fluxes. Nevertheless, R_{eco}
686 remains more indirectly constrained by EVI_{fp} than GPP.

687 Taken together, Fig. 3c–d, Fig. 4, Table S3, and Fig. S6 support the ecological
688 consistency of the CH-to-PY migration design: CH provides a cleaner ecological-
689 response reference under a shared regional background, while PY requires EVI_{fp} -based
690 scaling because of its lower vegetation activity and stronger source-area heterogeneity.
691 The migration hypothesis is supported by three lines of evidence: comparable regional
692 climatic and phenological forcing, EVI_{fp} -based separation of vegetation-associated
693 source areas from impervious surfaces, and migrated GPP and R_{eco} magnitudes that fall
694 within reported urban green-space and mixed-source ranges. Direct independent
695 validation of donor-site parameter migration remains limited, so this approach is treated
696 as an ecologically constrained parameterization rather than a universally validated
697 empirical law. Its influence on F_{ff} is further evaluated through the sensitivity analyses
698 in Sect. 3.5.

699 **3.3 Partitioned flux dynamics at the urban target site**

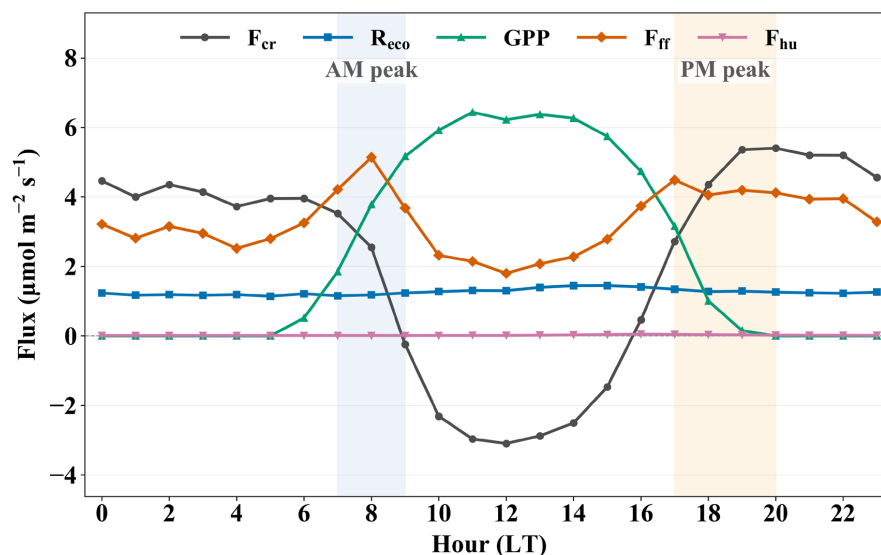
700 **3.3.1 Diurnal partitioning of CO_2 flux components**

701 Using the biogenic components constrained above, we partitioned the corrected CO_2
702 flux (F_{cr}) at PY and examined the resulting anthropogenic CO_2 flux (F_{ff}). This step links



703 the ecological correction to the question of how biogenic uptake and anthropogenic
704 emissions jointly shape net exchange in a mixed urban footprint.

705 Figure 5 shows that F_{cr} at PY has typical characteristics of a mixed urban green-
706 space source area. It is positive at night and in the evening, decreases substantially
707 during the day as GPP increases, and becomes negative around midday.
708 Correspondingly, GPP reaches high values during 11:00–14:00, with an hourly mean
709 peak of about $6.44 \mu\text{mol m}^{-2} \text{s}^{-1}$. R_{eco} shows a smaller diurnal amplitude and remains
710 around $1.14\text{--}1.45 \mu\text{mol m}^{-2} \text{s}^{-1}$. These results show that, although PY is an urban site,
711 its net flux still contains a pronounced biogenic uptake signal. In other words, negative
712 daytime F_{cr} does not mean that anthropogenic CO_2 emissions disappear; it means that
713 biogenic uptake offsets part of the anthropogenic emissions at the net-exchange level.



714
715 **Figure 5.** Main-case diurnal decomposition of corrected CO_2 flux at the PY site. Lines show
716 hourly means of corrected flux (F_{cr}), ecosystem respiration (R_{eco}), gross primary productivity
717 (GPP), anthropogenic CO_2 flux (F_{ff}), and human respiration flux (F_{hu}). Shaded intervals mark
718 the morning (07:00–09:00 LT) and evening (17:00–20:00 LT) peak windows in local time.

719 After partitioning, F_{ff} remained positive throughout the day, with hourly means of
720 about $1.80\text{--}5.14 \mu\text{mol m}^{-2} \text{s}^{-1}$ and a 24 h mean of about $3.29 \mu\text{mol m}^{-2} \text{s}^{-1}$. Its diurnal
721 variation showed a clear anthropogenic-activity rhythm: the 07:00–09:00 morning-rush
722 mean was about $4.34 \mu\text{mol m}^{-2} \text{s}^{-1}$, with the daily maximum at 08:00; the 17:00–20:00



723 evening-rush mean was about $4.22 \mu\text{mol m}^{-2} \text{s}^{-1}$; and the 11:00–14:00 midday mean
724 decreased to about $2.08 \mu\text{mol m}^{-2} \text{s}^{-1}$. This double-peak structure is consistent with the
725 rhythms of commuting, campus travel, campus services, and surrounding road traffic
726 in the Guangzhou Higher Education Mega Center, and with urban EC source-
727 partitioning studies showing that fossil-fuel or anthropogenic CO_2 fluxes often exhibit
728 traffic- or activity-related peaks (Lee et al., 2021; Wu et al., 2022).

729 In magnitude, F_{ff} at PY lies between two relevant urban green-space references.
730 Yang et al. (2025) reported a low-intensity traffic-emission background of $0\text{--}1.93 \mu\text{mol}$
731 $\text{m}^{-2} \text{s}^{-1}$ for a campus urban-forest environment in Nanjing, which is closer to a green-
732 space setting than to a dense traffic corridor. In contrast, Soininen et al. (2025) studied
733 a Helsinki footprint influenced by both urban green space and a busy street sector,
734 where light-response-curve partitioning inferred anthropogenic CO_2 fluxes of $3.1\text{--}10.9$
735 $\mu\text{mol m}^{-2} \text{s}^{-1}$. The PY F_{ff} range of $1.80\text{--}5.14 \mu\text{mol m}^{-2} \text{s}^{-1}$ falls between these two
736 references, consistent with the source-area character of the Guangzhou Higher
737 Education Mega Center, where green space, roads, buildings, and water bodies are
738 interwoven.

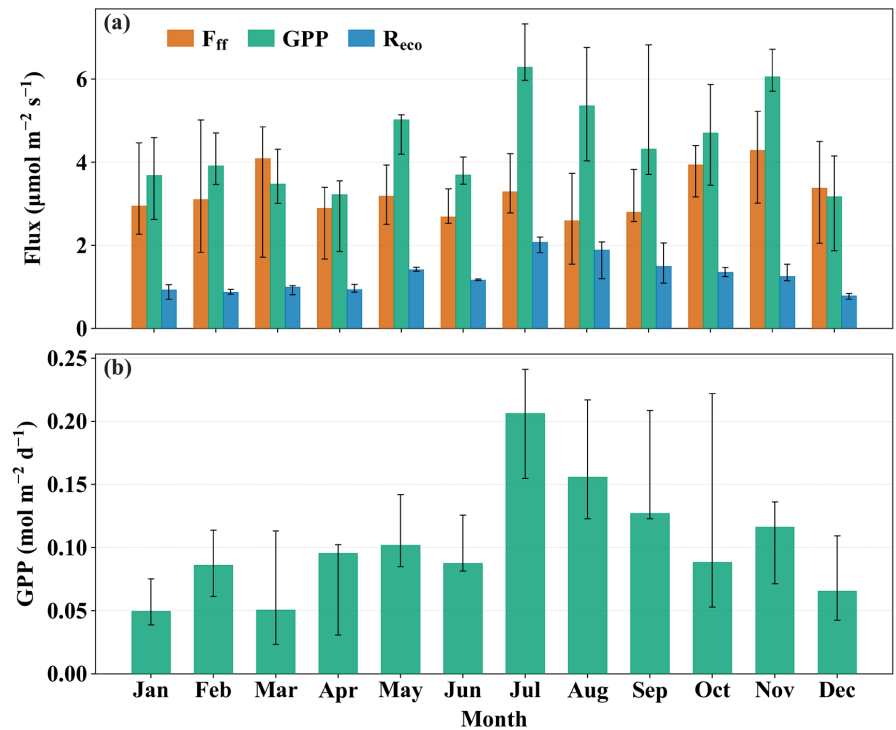
739 These results show how biogenic and anthropogenic components jointly shape the
740 observed net exchange at PY. During midday, strong GPP offsets a substantial part of
741 the anthropogenic source and can drive F_{cr} close to zero or negative values. During the
742 morning and evening periods, weaker biogenic uptake and enhanced anthropogenic
743 activity allow F_{ff} to dominate the net-flux signal. Thus, the urban net CO_2 flux is not
744 controlled by a single source process, but by the changing balance between persistent
745 anthropogenic emissions and temporally variable biogenic uptake.

746 3.3.2 Monthly variation in F_{ff} and its potential drivers

747 At the monthly scale, F_{ff} at PY remained positive in all months, indicating a persistent
748 anthropogenic CO_2 source signal within the PY footprint. Figure 6a shows that monthly
749 median F_{ff} was about $2.59\text{--}4.29 \mu\text{mol m}^{-2} \text{s}^{-1}$, with lower values in August and higher
750 values in October–November. The lowest monthly median occurred in August, about
751 $2.59 \mu\text{mol m}^{-2} \text{s}^{-1}$, while October and November increased to about 3.94 and $4.29 \mu\text{mol}$



752 $\text{m}^{-2} \text{s}^{-1}$, respectively. This range indicates moderate month-to-month variability
753 superimposed on a persistent anthropogenic source signal.



754
755 **Figure 6.** Monthly variations in daily aggregated partitioned CO_2 fluxes and daytime
756 cumulative GPP at the PY site. **(a)** Monthly medians of daily aggregated anthropogenic CO_2
757 flux (F_{ff}), daytime mean GPP, and ecosystem respiration (R_{eco}), with error bars showing the
758 interquartile range (Q25–Q75) derived from daily values within each month. **(b)** Monthly
759 medians of daily daytime cumulative GPP ($\text{mol m}^{-2} \text{d}^{-1}$), also shown with interquartile ranges.

760 This monthly pattern should not be attributed to a single factor. PY is located in a
761 university-town area, where summer and winter vacations, the start of semester, campus
762 services, and surrounding road traffic can influence local anthropogenic emissions.
763 August falls within the summer vacation, when teaching, commuting, and internal
764 campus service activities are reduced; this may help explain the low F_{ff} values. In
765 October–November, regular teaching activities and surrounding road activity resume,
766 providing a reasonable anthropogenic-activity background for higher F_{ff} . However, the
767 PY EC footprint does not cover only the campus interior; it also includes roads,
768 residential areas, water bodies, and urban service sources. Vacation effects may



769 therefore contribute to the monthly variation, but they should be interpreted together
770 with other footprint-scale sources and environmental controls rather than as a single
771 dominant driver of F_{ff} .

772 Meteorological and ecological conditions also modulate month-to-month F_{ff}
773 variation. Precipitation was notably high in August, which may have further reduced F_{ff}
774 by reducing travel, changing road-operating conditions, or altering activity intensity
775 within the effective footprint. At the same time, vegetation activity and GPP remained
776 strong in August, and ecological correction and changes in source-area vegetation
777 weight may also have affected the monthly median F_{ff} (Figs. 4 and S7). In contrast,
778 precipitation decreased substantially in October–November, and the combination of
779 resumed campus activity and less rainfall is consistent with stronger anthropogenic
780 sources. Monthly variation in PY F_{ff} should therefore be interpreted as the combined
781 result of anthropogenic activity, meteorological conditions, ecological correction, and
782 dynamic footprints, rather than as a direct response to a single driver.

783 **3.3.3 Eco-meteorological controls on GPP and R_{eco}**

784 Monthly variation in GPP and R_{eco} provides ecological context for understanding why
785 the apparent net CO_2 source strength and the inferred F_{ff} vary differently through the
786 year. Figures 6a–b show that GPP and R_{eco} at PY both strengthened in the warm season
787 and weakened in winter. This is consistent with the basic understanding that GPP in EC
788 flux partitioning is jointly controlled by radiation, vegetation activity, and water status,
789 whereas R_{eco} is mainly regulated by temperature and substrate availability (Reichstein
790 et al., 2005; Lasslop et al., 2010).

791 For GPP, the daytime cumulative GPP shown in Fig. 6b is more appropriate for
792 explaining seasonality because GPP is intrinsically a daytime process, whereas all-day
793 averages are affected by nighttime zeros, daylength, and the distribution of valid
794 samples. Daytime cumulative GPP at PY peaked in July, with a monthly median of
795 about $0.206 \text{ mol m}^{-2} \text{ d}^{-1}$. It remained relatively high in August–September but did not
796 continue to increase, and it decreased clearly in winter. High PAR, temperature, and
797 EVI_{fp} in July jointly supported strong photosynthesis. In August–September, heavy



798 rainfall and cloudy conditions may have reduced effective radiation, while elevated
799 VPD during some periods may also have induced stomatal regulation and limited
800 further GPP enhancement (Fig. S7). The Nanjing campus urban-forest study of Yang et
801 al. (2025) also showed that urban-forest GPP and R_{eco} are sensitive to radiation,
802 temperature, and water status. Thus, warm-season enhancement of PY GPP and its
803 fluctuations in rainy months are reasonable and should not be interpreted as a
804 monotonic response to temperature or precipitation alone.

805 Monthly variation in R_{eco} more directly reflects temperature control. Monthly
806 median R_{eco} at PY was about $0.79\text{--}2.07 \mu\text{mol m}^{-2} \text{s}^{-1}$, higher in summer and early
807 autumn and lower in winter, consistent with a Q10-type respiration model and seasonal
808 temperature variation. It is important to note that R_{eco} in this study was not fitted directly
809 at PY. It was obtained by migrating CH-side respiration parameters with EVI_{fp}
810 constraints and driving them with PY temperature. Its warm-season enhancement and
811 winter reduction therefore indicate that the migrated ecological respiration component
812 has reasonable physiological meaning.

813 Overall, the monthly biogenic components at PY show clear eco-meteorological
814 consistency: GPP increases with stronger radiation and greater vegetation activity under
815 warm-season conditions, but is modulated by rainfall, cloudiness, and VPD, whereas
816 R_{eco} mainly follows seasonal temperature variation. This supports the physiological
817 plausibility of the migrated biogenic components. More importantly for flux
818 interpretation, warm-season vegetation activity can reduce the apparent net CO_2 source
819 strength even when F_{ff} remains positive, whereas weaker winter biogenic uptake allows
820 the anthropogenic component to be more directly expressed in F_{cr} .

821 **3.4 Independent evidence for the anthropogenic interpretation of F_{ff}**

822 Sections 3.1–3.3 show that PY F_{ff} remained positive throughout the day and year and
823 exhibited a clear morning peak and weaker evening peak. Because F_{ff} is inferred after
824 ecological correction, its anthropogenic attribution requires independent evaluation. We
825 therefore evaluate F_{ff} using three complementary lines of evidence: pollutant diurnal
826 patterns, wind-sector land cover and traffic activity, and footprint-weighted inventories



827 that provide magnitude context and insight into possible spatial-allocation mismatch.
828 These comparisons are not used to validate F_{ff} as a direct traffic-emission estimate or
829 to attribute it to a single source category; rather, they provide independent consistency
830 checks for its anthropogenic-emission interpretation (Velasco and Roth, 2010;
831 Crawford and Christen, 2015; Wu et al., 2022).

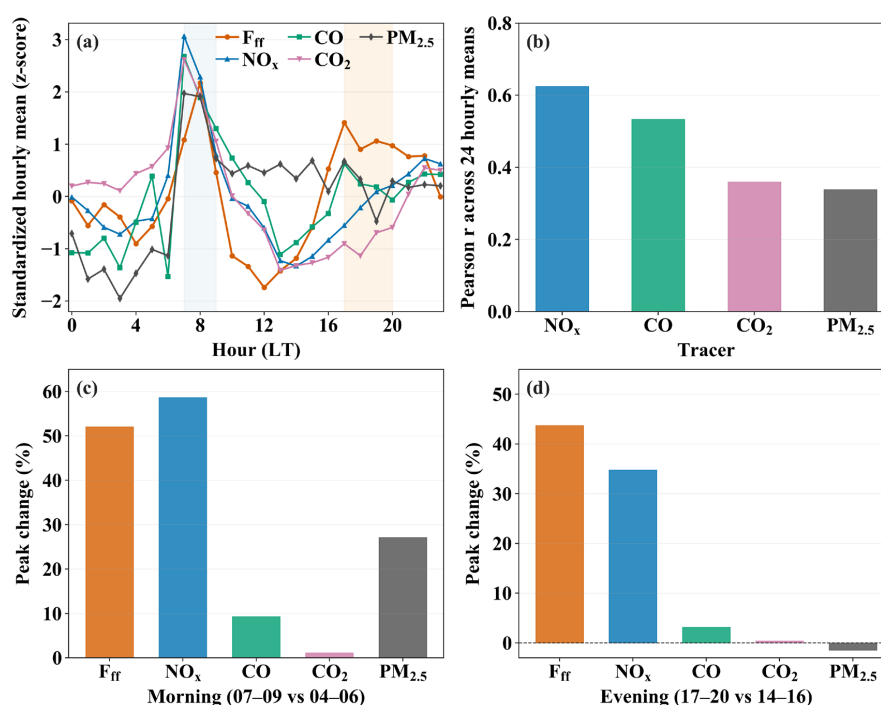
832 **3.4.1 Consistency with pollutant diurnal patterns**

833 Figure 7 compares the 24 h standardized diurnal patterns of F_{ff} with NO_x , CO, CO_2 , and
834 $PM_{2.5}$. Z-score standardization removes differences in units and magnitudes among
835 variables and focuses on phase, peak timing, and relative shape, rather than absolute
836 concentrations or fluxes. Figure 7a shows that F_{ff} has a clear morning peak during
837 07:00–09:00 and increases again during 17:00–20:00. NO_x and CO also increase
838 synchronously during the morning peak. Shape-correlation results further show that F_{ff}
839 has the highest 24 h standardized-pattern correlation with NO_x , with Pearson r about
840 0.62; its correlation with CO is the second highest, about 0.53; and its correlations with
841 CO_2 and $PM_{2.5}$ are weaker, about 0.36 and 0.34, respectively (Fig. 7b). This ranking is
842 physically consistent with urban anthropogenic emissions. NO_x is sensitive to motor
843 vehicles and local combustion processes (Parrish, 2006; Juchem et al., 2025), CO is
844 commonly used as an auxiliary tracer for combustion sources (Turnbull et al., 2011),
845 whereas CO_2 concentration and $PM_{2.5}$ are more strongly affected by boundary-layer
846 height, regional transport, secondary formation, and non-traffic sources (Xueref-Remy
847 et al., 2018; Guo et al., 2014). Compared with CO_2 concentration alone, the diurnal
848 consistency of F_{ff} with NO_x and CO therefore provides stronger support for a traffic- or
849 combustion-related anthropogenic CO_2 flux.

850 Peak-window analysis further strengthens this interpretation. During the morning
851 peak (07:00–09:00) relative to the 04:00–06:00 background, F_{ff} increased by about
852 52 %, and NO_x increased by about 59 %, with similar amplitudes. CO and $PM_{2.5}$ also
853 showed positive increases, but with smaller amplitudes (Fig. 7c). During the evening
854 peak from 17:00 to 20:00, F_{ff} and NO_x increased by about 44 % and 35 % relative to
855 14:00–16:00, respectively, whereas CO_2 and $PM_{2.5}$ showed weaker changes (Fig. 7d).



856 These results indicate that the morning and evening F_{ff} peaks are unlikely to be artifacts
857 of the partitioning procedure; instead, they show an activity rhythm consistent with
858 traffic- and combustion-related pollutants. Similarly, Wu et al. (2022) emphasized that
859 the fossil-fuel component in urban EC CO_2 fluxes should be evaluated using CO , ^{14}C ,
860 or high-resolution emission information, rather than inferred from the net CO_2 flux
861 pattern alone.



862 **Figure 7.** Diurnal-shape consistency and peak-window enhancement characteristics of the
863 anthropogenic CO_2 flux (F_{ff}) and selected atmospheric tracers at the PY site. **(a)** Standardized
864 hourly mean (z-score) curves for F_{ff} and the selected tracers. Each series was first aggregated
865 independently to hourly means and then standardized to highlight shape similarity rather than
866 magnitude. The shaded bands indicate the morning (07:00–09:00 LT) and evening (17:00–
867 20:00 LT) peak period. **(b)** Pearson correlation coefficients between the standardized 24 h
868 shape of F_{ff} and each tracer, computed across the 24 h mean values. **(c)** Morning peak-window
869 enhancement, expressed as the percentage change from the pre-window (04:00–06:00 LT) to
870 the peak window (07:00–09:00 LT). **(d)** Evening peak-window enhancement, expressed as
871 the percentage change from the pre-window (14:00–16:00 LT) to the peak window (17:00–
872 20:00 LT).

874 3.4.2 Wind-sector, land-cover, and traffic evidence

875 Morning-peak F_{ff} differed substantially among the eight wind sectors (Fig. 8a). The NE



sector was highest, with a mean of about $6.16 \mu\text{mol m}^{-2} \text{s}^{-1}$; the N sector was second, about $4.81 \mu\text{mol m}^{-2} \text{s}^{-1}$; and the SW sectors also remained relatively high, about $4.46 \mu\text{mol m}^{-2} \text{s}^{-1}$. The W sector was lowest, about $2.48 \mu\text{mol m}^{-2} \text{s}^{-1}$. Thus, morning-peak F_{ff} was not enhanced uniformly across all wind directions, but was concentrated in specific source-area directions.

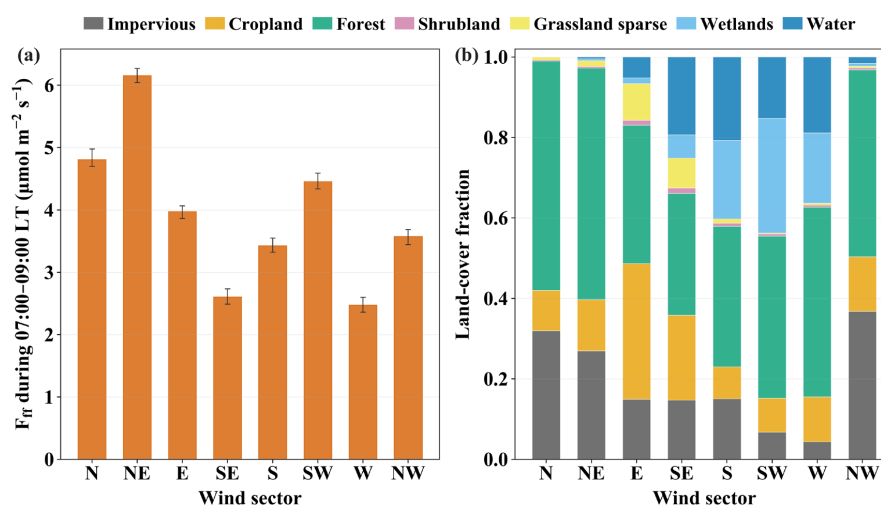


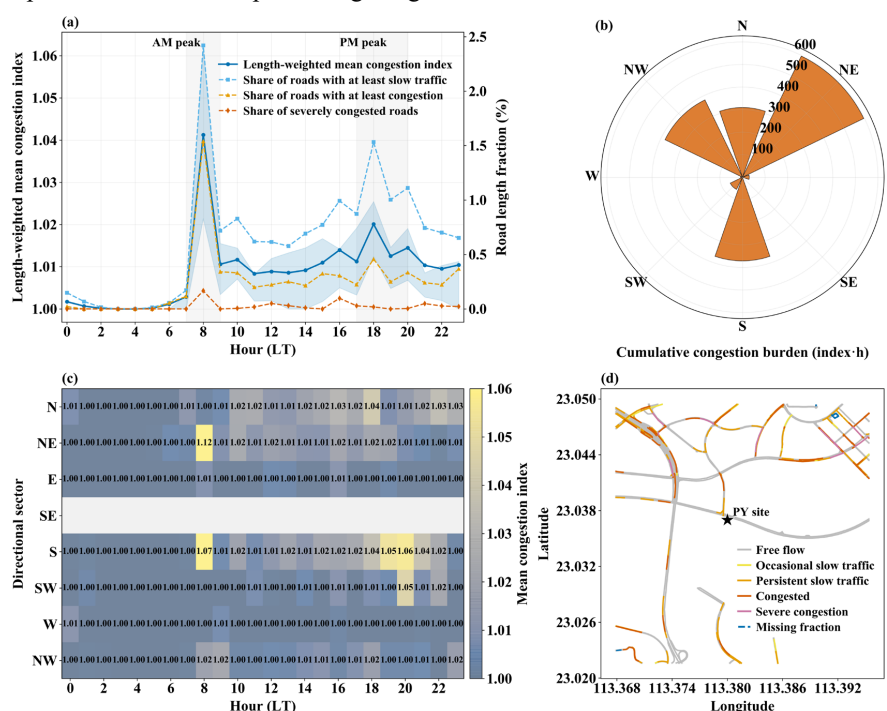
Figure 8. Morning-peak anthropogenic CO₂ flux and footprint-weighted land-cover composition by wind sector at the PY site. **(a)** Mean anthropogenic CO₂ flux (F_{ff}) during 07:00–09:00 LT for eight wind-direction sectors, using the same 45° sector classification as in Fig. 3. Error bars indicate the interquartile range (Q25–Q75). **(b)** Corresponding footprint-weighted land-cover composition for each sector during the same morning-peak period.

Footprint-weighted land cover shows that high F_{ff} is broadly consistent with urbanized and mixed activity source areas, but is not controlled by impervious fraction alone (Fig. 8b). The N, NE, and NW sectors had impervious fractions of about 33 %, 27 %, and 37 %, respectively, and also high forest fractions of about 46 %–58 %. This indicates that urban green space, roads, and hard surfaces are closely interwoven in the PY source area. The simultaneously high F_{ff} and high impervious fraction in the NE and N sectors support their urban-activity source character. In contrast, the SW sector had a lower impervious fraction but higher F_{ff} , suggesting that local road nodes, parking entrances, campus gates, or short-term traffic aggregation areas can generate strong anthropogenic CO₂ signals even under a lower impervious-background fraction in



897 mixed urban source areas.

898 Road-congestion indicators provide additional supporting evidence for this
899 interpretation (Fig. 9). The extended congestion dataset covers 331 hourly road
900 snapshots from 23:00 LT on 6 May to 13:00 LT on 24 May 2026, including 713 road
901 segments within approximately 1.5 km of the PY site. These data are not synchronous
902 with the main EC analysis period (2023–2024) and are therefore not used for hourly
903 validation of F_{ff} or for conversion to traffic CO₂ emission fluxes. Instead, they provide
904 an auxiliary description of typical road-traffic timing, directional contrast, and line-
905 source distribution around the PY university-town area. Details on data source,
906 representativeness, and processing are given in Sect. S8.



907
908 **Figure 9.** Traffic-activity indicators used to interpret the temporal, directional, and spatial
909 behavior of the PY anthropogenic CO₂ flux (F_{ff}). **(a)** Diurnal variation of the length-weighted
910 mean congestion index and the fractions of road length with at least slow traffic, at least
911 congestion, and severe congestion. **(b)** Eight-sector rose plot of the cumulative congestion
912 burden (index·h). **(c)** Hour × directional-sector heat map of the mean congestion index. **(d)**
913 Spatial distribution of road-segment mean congestion class surrounding the PY site, used to
914 identify the road segments responsible for the directional congestion patterns shown in panel
915 (b).



916 Figure 9a shows a clear traffic-activity rhythm. The length-weighted mean
917 congestion index and the fractions of road length with slow or congested traffic
918 increased during the morning and evening traffic-relevant periods, with the strongest
919 and sharpest enhancement around 08:00 LT and a broader but weaker increase after
920 approximately 16:00 LT. The overall congestion amplitude remained modest, indicating
921 that most roads were free-flowing or only mildly slow during the monitoring period.
922 Nevertheless, the timing and relative strength of the congestion enhancements are
923 consistent with the high-value windows of F_{ff} and NO_x , supporting the interpretation
924 that the double-peak structure of F_{ff} is broadly associated with road-traffic activity.

925 The directional traffic evidence further supports the sectoral interpretation of F_{ff} .
926 The cumulative congestion burden is unevenly distributed among directions and is
927 strongest in the NE sector, with additional contributions from the N, NW, and S sectors
928 (Fig. 9b). The hour-sector heat map shows a pronounced NE morning enhancement
929 around 08:00 LT, consistent with the highest morning-peak F_{ff} in the NE sector (Fig.
930 8a). The S and SW sectors show stronger congestion from evening to nighttime, helping
931 explain why traffic-related activity can also contribute to relatively high F_{ff} in these
932 directions despite their lower impervious-background fraction. The road-segment map
933 further links these directional patterns to specific road corridors near PY, especially to
934 the NE and along several southern to southwestern road sections (Figs. 9d and S8).

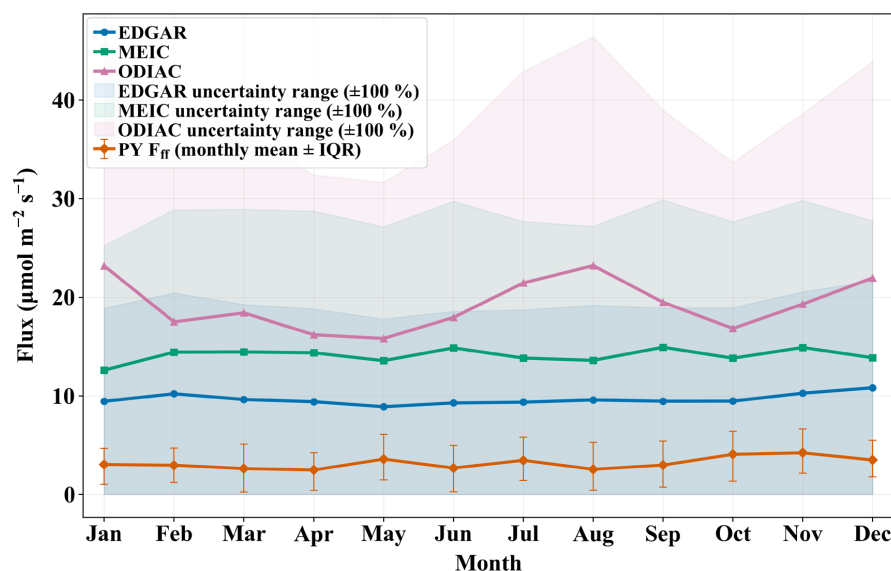
935 Taken together, Figs. 8, 9, and S8 indicate that spatial differences in morning-peak
936 F_{ff} at PY reflect the combined effects of urbanized land cover, local road or gate-related
937 line sources, and dynamic EC footprints. The agreement among F_{ff} , pollutant peaks,
938 footprint-weighted land cover, sectoral congestion burden, and road-segment
939 congestion patterns strengthens the anthropogenic interpretation of F_{ff} , while non-
940 monotonic relationships among these indicators reflect the complexity of mixed
941 university-town source areas.

942 3.4.3 Inventory magnitude context and spatial-allocation implications

943 Figure 10 compares PY F_{ff} with footprint-weighted monthly estimates from three
944 widely used bottom-up inventories: the Open-Data Inventory for Anthropogenic



945 Carbon Dioxide (ODIAC; Oda et al., 2018), the Emissions Database for Global
946 Atmospheric Research (EDGAR; Crippa et al., 2024), and the Multi-resolution
947 Emission Inventory model for Climate and air pollution research (MEIC; Xu et al.,
948 2024). Detailed information on the inventories and calculation procedure is provided in
949 Sect. S5. Inventory central estimates are generally higher than the observation-inferred
950 F_{ff} : EDGAR is about $9\text{--}11\text{ }\mu\text{mol m}^{-2}\text{ s}^{-1}$, MEIC about $13\text{--}15\text{ }\mu\text{mol m}^{-2}\text{ s}^{-1}$, and ODIAC
951 about $16\text{--}23\text{ }\mu\text{mol m}^{-2}\text{ s}^{-1}$, whereas monthly PY F_{ff} is about $2.59\text{--}4.29\text{ }\mu\text{mol m}^{-2}\text{ s}^{-1}$.
952 This several-fold difference should be interpreted in the context of both spatial-support
953 mismatch and spatial-proxy mismatch. Global and regional gridded inventories do not
954 directly observe emissions within the EC footprint. Instead, they allocate national,
955 regional, or sectoral activity data to grid cells using proxy variables such as population
956 density, night-time lights, road networks, point-source locations, and sectoral activity
957 distributions. ODIAC distributes non-point fossil-fuel CO_2 emissions mainly using
958 satellite-observed night-time lights after separately treating large point sources (Oda et
959 al., 2018). EDGAR uses sector-specific spatial proxies, including population, road
960 networks, facility locations, and land-use information, to construct gridded emissions
961 (Crippa et al., 2024). MEIC improves sectoral and subnational representation, but its
962 gridded fields are still spatially allocated inventory estimates rather than direct
963 footprint-scale observations (Xu et al., 2024). This proxy-allocation issue is particularly
964 relevant for the PY footprint. The Guangzhou Higher Education Mega Center is a high-
965 population-density and brightly lit urban functional zone with dense road infrastructure,
966 but its on-site fossil-fuel combustion intensity per capita is likely lower than that of
967 industrial, commercial, or heavily trafficked urban cores (Sect. 3.3.1). Buildings within
968 the source area mainly rely on externally supplied electricity, and many roads are low-
969 speed campus-service roads or have restricted access (Sect. S8). Population density,
970 night-time lights, and road density may therefore over-represent local combustion
971 emissions when used as spatial allocation proxies. In such a setting, footprint-weighted
972 inventory values can be several times larger than the EC-inferred F_{ff} because broader
973 urban emissions are partly assigned to a high-proxy-intensity functional zone with
974 relatively low on-site combustion.



975

976 **Figure 10.** Monthly comparison between the PY anthropogenic CO₂ flux (F_{ff}) and footprint-
 977 aligned bottom-up inventory estimates. The PY observation represents the CH0 main
 978 scenario with the base human-respiration setting and is summarized as the monthly mean
 979 with the interquartile range (Q25–Q75). EDGAR, MEIC, and ODIAC are shown as monthly
 980 central estimates. The shaded bands show an illustrative $\pm 100\%$ envelope to visualize broad
 981 urban-grid inventory uncertainty, rather than inventory-specific confidence intervals.

982 To illustrate the possible magnitude of city- or grid-scale inventory differences,
 983 Fig. 10 uses a $\pm 100\%$ range as an illustrative uncertainty envelope, not as an inventory-
 984 specific confidence interval. Although the inventory central estimates are several times
 985 higher than PY F_{ff} , the observation-inferred monthly F_{ff} remains within the illustrative
 986 $\pm 100\%$ inventory envelopes shown in Fig. 10. This indicates that the discrepancy is
 987 large, but still falls within the broad local/grid-scale uncertainty context reported for
 988 urban fossil-fuel CO₂ inventories. Previous studies show that fossil-fuel CO₂ inventory
 989 uncertainty is strongly scale dependent: it is relatively small at national or regional
 990 aggregate scales but can increase substantially at city, neighborhood, or 0.1° grid scales.
 991 Gately and Hutyra (2017) reported that different inventories can differ by 50 %–250 %
 992 at city scale and that many 0.1° grids differ by more than 100 %. More importantly for
 993 the present study, spatial proxy variables themselves can introduce local positive bias
 994 when they do not represent the true distribution of combustion activity. Squires et al.



995 (2020) compared EC measurements of traffic-dominated NO_x and CO fluxes in central
996 Beijing with MEIC and found that MEIC substantially overestimated emissions within
997 the EC footprint, providing direct evidence that proxy-based inventories can have
998 positive biases for traffic-related emissions in Chinese urban centers. Wu et al. (2022)
999 similarly emphasized that EC-based source decomposition is needed to evaluate high-
1000 resolution urban CO_2 inventories at the footprint scale rather than relying on net CO_2
1001 flux or gridded inventories alone.

1002 From this perspective, the lower PY F_{ff} relative to EDGAR, MEIC, and ODIAC
1003 should not be interpreted simply as evidence that the EC-based estimate is biased low.
1004 It more likely reflects a combination of footprint-scale ecological correction, local
1005 functional-zone representativeness, and proxy-based spatial allocation errors in gridded
1006 inventories. A model-side contribution to the difference is also possible. Under the
1007 present sign convention, underestimated GPP or overestimated R_{eco} would shift F_{ff}
1008 downward, whereas overestimated GPP would shift F_{ff} upward. The potential influence
1009 of ecological-correction uncertainty on F_{ff} magnitude is examined in Sect. 3.5. The
1010 inventory comparison therefore provides a magnitude reference and a diagnostic of
1011 spatial-allocation plausibility, rather than numerical validation of either the inventories
1012 or F_{ff} . The large inventory– F_{ff} difference at PY further indicates that spatial-proxy
1013 mismatch can be substantial in special-function urban zones, where population density,
1014 night-time lights, and road density do not necessarily scale with local fossil-fuel
1015 combustion intensity.

1016 Overall, the anthropogenic-emission attribution of PY F_{ff} is constrained by
1017 multiple independent lines of evidence. Temporally, F_{ff} is consistent with the diurnal
1018 patterns of NO_x and CO, and the morning-peak enhancement is close to that of NO_x .
1019 Spatially, high F_{ff} in the eight-sector analysis occurs mainly in NE, N, and SW
1020 directions, where urban activity or road-traffic contributions are stronger, and is broadly
1021 consistent with footprint-weighted land cover, sectoral congestion load, and road-
1022 segment congestion distribution. In magnitude, F_{ff} is lower than bottom-up inventory
1023 central estimates, but this comparison provides an urban-emission background, a
1024 plausibility boundary, and a diagnostic of possible proxy-allocation mismatch in a



1025 special-function urban zone. Thus, PY F_{ff} is not an isolated residual produced only by
1026 model calculation. Rather, it is an ecologically corrected anthropogenic CO_2 flux
1027 estimate supported by multiple consistency checks across pollutant timing, source-area
1028 structure, traffic activity, and inventory magnitude context.

1029 **3.5 Uncertainty sources and robustness of F_{ff}**

1030 Donor–target parameter migration is the core structural assumption of the
1031 framework. Its uncertainty arises from differences in canopy structure, vegetation
1032 composition, soil moisture, substrate availability, and management intensity between
1033 CH and PY, which cannot be fully captured by EVI. Because F_{ff} is obtained after
1034 subtracting migrated biogenic fluxes and human respiration from the storage-corrected
1035 net flux, we treat the migrated biogenic components as constrained approximations and
1036 evaluate how key assumptions affect F_{ff} . This uncertainty is especially relevant for R_{eco} ,
1037 for which EVI_{fp} mainly constrains the vegetation-associated respiratory capacity within
1038 the footprint but cannot fully represent soil moisture, substrate supply, irrigation, or
1039 management-driven differences between CH and PY. These unresolved R_{eco} controls
1040 are treated as part of the structural boundary of the donor–target migration framework,
1041 rather than as separately identifiable uncertainty sources in the present sensitivity
1042 design. The analysis focuses on whether sensitivity scenarios mainly change the
1043 absolute magnitude of F_{ff} or also alter its positive sign across seasons and its morning–
1044 evening peak structure.

1045 **3.5.1 Relative contributions of uncertainty sources**

1046 Figure 11 summarizes the mean F_{ff} offsets of each sensitivity scenario relative to the
1047 baseline. Overall, GPP model structure is the dominant source of variation in F_{ff}
1048 magnitude. The pure saturating light-response structure and the saturating + VPD
1049 inhibition structure reduce mean F_{ff} by about 0.546 and 0.547 $\mu mol\ m^{-2}\ s^{-1}$, respectively,
1050 clearly larger than the effects of other processing steps. By comparison, CH-side
1051 shortwave-radiation perturbations of -10% and $+10\%$ change mean F_{ff} by about
1052 $+0.092$ and $-0.085\ \mu mol\ m^{-2}\ s^{-1}$, respectively; the EVI 5 d window reduces F_{ff} by about



0.077 $\mu\text{mol m}^{-2} \text{s}^{-1}$; the u^* threshold within $\pm 20\%$ changes F_{ff} by about -0.068 to $+0.021$ $\mu\text{mol m}^{-2} \text{s}^{-1}$; Scheme B and weak daytime-respiration inhibition have effects of about ± 0.041 $\mu\text{mol m}^{-2} \text{s}^{-1}$; and the $\pm 10\%$ storage-flux perturbation is close to zero, within about 0.005 $\mu\text{mol m}^{-2} \text{s}^{-1}$.

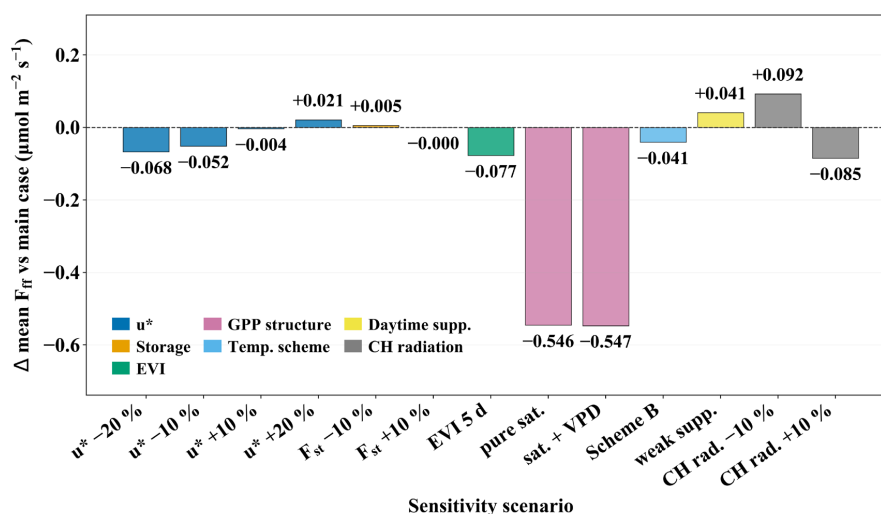


Figure 11. Module sensitivity of the PY anthropogenic CO_2 flux (F_{ff}) relative to the main case. Bars show the difference in the overall mean F_{ff} between each sensitivity scenario and the main case, using the base human-respiration setting. Colors denote uncertainty modules, including u^* threshold, storage correction, EVI temporal window, GPP model structure, temperature-group scheme, daytime respiration suppression, and CH-only shortwave-radiation perturbation.

This result shows that uncertainty in PY F_{ff} mainly arises from the biogenic-uptake term, especially the structural estimation of daytime GPP, rather than from small additive terms such as storage flux or u^* filtering. This directly addresses the inventory-comparison concern raised in Sect. 3.4.3: ecological-correction uncertainty can shift the absolute magnitude of F_{ff} , but the tested biogenic-correction scenarios do not remove the several-fold gap between F_{ff} and the footprint-weighted inventory central estimates. The CH-only radiation-input perturbation has a larger effect than storage flux and daytime-respiration inhibition, but remains substantially smaller than the GPP-structure difference. Thus, CH radiation representativeness is an input uncertainty that should be reported, but it is not the dominant factor controlling the F_{ff} conclusions. This dominance is expected because GPP is the key daytime subtraction term in the F_{ff}



1075 calculation; different light-response structures directly alter the daytime biogenic-
1076 uptake estimate and therefore the amplitude of diurnal F_{ff} variation. Previous EC flux-
1077 partitioning studies also show that nighttime respiration extrapolation and daytime GPP
1078 model structure can strongly affect NEE partitioning, especially during periods of
1079 strong daytime biogenic uptake, when structural differences are often more important
1080 than random parameter error (Reichstein et al., 2005; Lasslop et al., 2010; Desai et al.,
1081 2008). It is also notable that the pure saturating structure and the saturating + VPD
1082 inhibition structure have almost identical effects on F_{ff} . This is consistent with the CH-
1083 side model comparison in Sect. 3.2, which showed that the explicit VPD inhibition term
1084 did not add explanatory power beyond the pure saturating structure in the present
1085 dataset. Figure 11 therefore not only identifies GPP structure as the main uncertainty
1086 source, but also further supports the temperature-stratified light-response structure as
1087 the baseline scheme.

1088 **3.5.2 Robustness of diurnal and seasonal patterns**

1089 Figure 12a compares the diurnal spread of F_{ff} among the baseline and 13 sensitivity or
1090 structural-alternative scenarios with parameter-bootstrap uncertainty. The scenario
1091 spread is larger than the bootstrap uncertainty during most hours. At the hourly scale,
1092 the mean scenario q90–q10 spread is about $0.47 \mu\text{mol m}^{-2} \text{s}^{-1}$, whereas the mean
1093 bootstrap q90–q10 spread is about $0.21 \mu\text{mol m}^{-2} \text{s}^{-1}$; the former is about 2.3 times the
1094 latter. Thus, within the present framework, model structure and processing-scenario
1095 choices explain the overall uncertainty in F_{ff} more strongly than bootstrap resampling
1096 of parameters alone. This result is consistent with previous flux-partitioning studies
1097 showing that parameter-estimation error is important, but light-response structure and
1098 data-processing choices can produce systematic shifts in partitioning results (Papale et
1099 al., 2006; Lasslop et al., 2010; Richardson and Hollinger, 2007).

1100 In terms of diurnal variation, scenario spread is more pronounced during the
1101 morning and evening peaks and the daytime transition periods. The largest scenario
1102 q90–q10 spread occurs at 16:00, 17:00, and 08:00, corresponding to the afternoon
1103 transition when biogenic uptake weakens, the evening peak, and the morning peak. This



gives the F_{ff} uncertainty a clear process meaning: at night, GPP is close to zero and F_{ff} is mainly controlled by net flux and respiration terms; during daytime and evening, it is affected by both ecological-correction choices and anthropogenic-activity rhythms, so scenario differences are larger. Despite these amplitude differences, the morning and evening peaks persist in all scenarios, and the relatively low midday F_{ff} pattern is not changed. Thus, scenario uncertainty mainly affects F_{ff} magnitude after daytime biogenic-uptake correction, rather than the presence of a diurnal structure associated with urban anthropogenic activity.

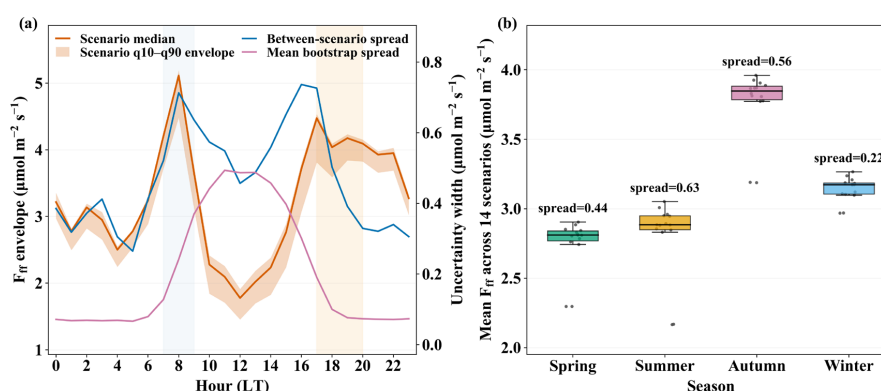


Figure 12. Integrated uncertainty-spread diagnostics for the PY anthropogenic CO_2 flux (F_{ff}). (a) Diurnal comparison between the scenario envelope and bootstrap uncertainty under the base human-respiration setting. The colored uncertainty ribbon, bounded by the q10 and q90 lines, represents the envelope of hourly mean F_{ff} across the 14 sensitivity scenarios; the solid line shows the scenario median. The between-scenario spread (q90–q10) and the mean within-scenario bootstrap spread (q90–q10) are also shown. Pale vertical background bands highlight the morning (07:00–09:00 LT) and evening (17:00–20:00 LT) traffic-relevant periods. (b) Seasonal distributions of scenario-mean F_{ff} across the 14 CH scenarios under the base human-respiration setting. Boxplots summarize the spread among scenarios for each season, and points denote individual scenario means.

Figure 12b further shows that the seasonal uncertainty in F_{ff} is also linked to biogenic processes. Across the 14 scenarios, the scenario range is largest in summer, followed by autumn, and smallest in winter. This is consistent with the seasonality of the biogenic components: warm-season GPP and R_{eco} are stronger, so ecological model structure and radiation input have larger effects on F_{ff} . In winter, biogenic processes are weaker, and differences caused by model structure and input perturbations are reduced. In other words, F_{ff} uncertainty is not randomly distributed. It is concentrated in warm-



1130 season and daytime periods, when ecological correction is strongest and component
1131 separation is most difficult.

1132 This pattern also provides a boundary test for the CH-to-PY migration assumption:
1133 scenario spread is largest when ecological correction is strongest, but the positive sign
1134 across seasons and the morning–evening peak structure remain stable.

1135 **3.5.3 Interpretive boundaries of F_{ff}**

1136 Core structural scenarios further clarify the interpretive boundary of F_{ff} (Fig. S9a). The
1137 baseline scheme, Scheme B, and the weak daytime-respiration-inhibition scenario
1138 produce similar diurnal shapes, whereas the pure saturating and saturating + VPD
1139 inhibition structures yield lower F_{ff} values, especially during daytime and evening. Thus,
1140 GPP structure mainly controls the inferred magnitude of F_{ff} , while the morning peak,
1141 midday low, and evening peak remain robust. This supports using F_{ff} as a footprint-
1142 scale anthropogenic-activity signal, but its absolute value remains conditional on the
1143 ecological model structure and donor–target migration assumptions.

1144 Human-respiration sensitivity provides a separate boundary on the non-fossil
1145 anthropogenic term (Fig. S9b–c). Even when human respiration is set to 0.5 or 1.5 times
1146 the baseline, the mean change in F_{ff} is only about $\pm 0.01 \mu\text{mol m}^{-2} \text{s}^{-1}$, and the maximum
1147 hourly effect is about $\pm 0.026 \mu\text{mol m}^{-2} \text{s}^{-1}$. Human respiration should therefore be
1148 explicitly subtracted, but it is not a dominant uncertainty under the current PY source-
1149 area background. Overall, PY F_{ff} is best interpreted as an ecologically constrained,
1150 footprint-scale anthropogenic CO_2 flux estimate: robust as a relative temporal and
1151 source-area signal, but not a model-independent observation or direct inversion of local
1152 traffic or inventory emissions.

1153 **4 Conclusions**

1154 Using paired EC observations from the CH and PY sites in Guangzhou, this study
1155 developed a footprint-informed ecological parameter-migration framework for
1156 partitioning CO_2 fluxes in mixed urban source areas. The framework uses CH as a
1157 relatively clean ecological donor, transfers vegetation-normalized ecological



parameters to PY using dynamic footprints and footprint-weighted enhanced vegetation index (EVI_{fp}), and estimates footprint-scale anthropogenic CO_2 flux (F_{ff}) after ecological and human-respiration corrections.

The CH-to-PY migration was supported by a consistent ecological evidence chain. The two sites share a similar regional climatic and phenological background but differ strongly in vegetation cover and anthropogenic influence. After EVI_{fp} -based scaling, the migrated PY gross primary productivity (GPP) and ecosystem respiration (R_{eco}) showed plausible diurnal and seasonal patterns, with magnitudes consistent with reported urban green-space and mixed-source fluxes. These results support the donor-target design as an ecologically consistent constraint on biogenic components, while the migrated fluxes should still be interpreted as constrained approximations rather than directly validated ecological observations.

The partitioned components reveal how biogenic and anthropogenic processes jointly shape net CO_2 exchange at the urban target site. PY remained a net CO_2 source overall, but daytime storage-corrected net CO_2 flux (F_{cr}) could approach neutral values or become weakly negative when GPP was strong. After ecological correction, F_{ff} remained positive throughout the day, with hourly means of $1.80\text{--}5.14 \mu\text{mol m}^{-2} \text{s}^{-1}$ and a clear morning peak and weaker evening peak. Thus, urban EC net flux cannot be interpreted directly as anthropogenic emissions because vegetation uptake can mask part of the anthropogenic signal.

The inferred F_{ff} showed temporal and source-area patterns consistent with footprint-scale anthropogenic activity, including diurnal agreement with combustion-related tracers and high morning-peak values in sectors with stronger urban activity or traffic-related source contributions. Footprint-weighted inventory comparisons provided an order-of-magnitude context, but should not be interpreted as strict inventory validation. The large inventory- F_{ff} difference at PY further suggests that proxy-based spatial allocation can be problematic in special-function urban zones, where population density, night-time lights, and road density do not necessarily scale with local fossil-fuel combustion intensity.

The uncertainty analysis showed that F_{ff} was most sensitive to GPP light-response



1188 structure, whereas other input, screening, parameterization, storage, and human-
1189 respiration choices had smaller effects. Scenario differences mainly affected the
1190 absolute magnitude of F_{ff} , especially during daytime and the warm season, but did not
1191 change its positive sign across seasons or its morning–evening peak structure. F_{ff} should
1192 therefore be interpreted as an ecologically constrained, footprint-scale anthropogenic
1193 CO_2 flux estimate, not as a model-independent direct emission observation.

1194 Several limitations remain. The uncertainty framework is based on controlled
1195 sensitivity scenarios rather than a complete probability distribution of all error sources.
1196 The CH-to-PY migration depends on ecological similarity, EVI_{fp} -based scaling,
1197 footprint and remote-sensing representativeness, CH-side radiation input, and site-level
1198 microclimatic contrasts. These factors define the main boundary for extending the
1199 framework to other urban sites. F_{ff} should be used as a tower-footprint-scale
1200 observational constraint and relative-change indicator, rather than as a direct
1201 replacement for city-total inventories. Future work combining multi-site EC networks
1202 with traffic, energy-use, mobile, isotopic, or auxiliary tracer observations would help
1203 further test and extend this framework.

1204 This study demonstrates the feasibility of an EC-based CO_2 flux-partitioning
1205 approach for urban areas where green spaces and anthropogenic sources are highly
1206 mixed. By combining standard EC CO_2 fluxes with dynamic footprints, high-resolution
1207 land cover, and open satellite vegetation-index products, the framework lowers the
1208 observational barrier for source–sink partitioning in data-scarce and rapidly urbanizing
1209 regions, including many cities in the Global South. Beyond flux partitioning, it helps
1210 reveal anthropogenic CO_2 emission signals masked by biogenic processes, while
1211 providing footprint-scale constraints for diagnosing potential spatial-proxy mismatch
1212 in gridded inventories and informing urban carbon-monitoring network design.

1213 **Code and data availability.** The source code for the two-dimensional flux-footprint
1214 model used in this study is available at <https://footprint.kljun.net> (Kljun et al., 2015).
1215 The global 10 m land-cover product `GLC_FCS10` used in this study has been published
1216 on Zenodo and can be accessed at <https://doi.org/10.5281/zenodo.14729665> (last access:
1217 25 May 2026) (Zhang et al., 2025). The global 10 m canopy-height product used in this



study is publicly available and can be downloaded from https://langnico.github.io/globalcanopyheight/assets/tile_index.html (Lang et al., 2023). The Harmonized Landsat Sentinel-2 (HLS) 30 m vegetation-index products used in this study are publicly available at https://doi.org/10.5067/HLS/HLSL30_V1.002 and <https://doi.org/10.5067/HLS/HLSS30.002> (last access: 25 May 2026) (Neigh, 2021; Neigh, 2025). The road-traffic congestion and vector building-height data around the PY site used in this study are available from <https://lbs.amap.com> (last access: 25 May 2026). The national hourly population-density spatial-distribution dataset used in this study can be accessed at <https://doi.org/10.12078/2022091601> (Xu, 2022). Additional data and information used in this study are available from the corresponding author upon request.

Author contributions. JWZ and ML designed the study. JWZ, YJL, YFY, CLP, HBH, BH, XFL, YYH, YWP, CLC, CW, ZZ, and ML contributed to data collection and data analysis. JWZ designed and performed the model simulations. JWZ and ML wrote the paper with contributions from all coauthors. All the authors revised the paper and edited the text.

Competing interests. The contact author has declared that none of the authors has any competing interests.

Disclaimer. Publisher's note: Copernicus Publications remains neutral with regard to jurisdictional claims made in the text, published maps, institutional affiliations, or any other geographical representation in this paper. The authors bear the ultimate responsibility for providing appropriate place names. Views expressed in the text are those of the authors and do not necessarily reflect the views of the publisher.

Acknowledgements. The authors would like to thank the personnel who participated in data collection, instrument maintenance, and logistic support during the field campaign.

Financial support. This research has been supported by the National Natural Science Foundation of China (grant no. 42477273) and the National Key R&D Program of China (grant no. 2022YFE0209500).

References

- Aubinet, M., Chermanne, B., Vandenhaute, M., Longdoz, B., Yernaux, M., and Laitat, E.: Long term carbon dioxide exchange above a mixed forest in the Belgian Ardennes, Agr. Forest Meteorol., 108, 293-315, [https://doi.org/10.1016/S0168-1923\(01\)00244-1](https://doi.org/10.1016/S0168-1923(01)00244-1), 2001.
- Baldocchi, D. D.: Assessing the eddy covariance technique for evaluating carbon dioxide



- exchange rates of ecosystems: past, present and future, *Global Change Biol.*, 9, 479-492,
<https://doi.org/10.1046/j.1365-2486.2003.00629.x>, 2003.
- Crawford, B. and Christen, A.: Spatial source attribution of measured urban eddy covariance
CO₂ fluxes, *Theor. Appl. Climatol.*, 119, 733-755, [https://doi.org/10.1007/s00704-014-1124-](https://doi.org/10.1007/s00704-014-1124-0)
0, 2015.
- Crawford, B., Grimmond, C. S. B., and Christen, A.: Five years of carbon dioxide fluxes
measurements in a highly vegetated suburban area, *Atmos. Environ.*, 45, 896-905,
<https://doi.org/10.1016/j.atmosenv.2010.11.017>, 2011.
- Crippa, M., Guizzardi, D., Pagani, F., Banja, M., Muntean, M., Schaaf, E., Monforti-Ferrario,
F., Becker, W. E., Quadrelli, R., Risquez Martin, A., Taghavi-Moharamli, P., Köykkä, J.,
Grassi, G., Rossi, S., Melo, J., Oom, D., Branco, A., San-Miguel, J., Manca, G., Pisoni, E.,
Vignati, E., and Pekar, F.: GHG emissions of all world countries, Publications Office of the
European Union, <https://doi.org/10.2760/4002897>, 2024.
- Decina, S. M., Hutyra, L. R., Gately, C. K., Getson, J. M., Reinmann, A. B., Short Gianotti, A.
G., and Templer, P. H.: Soil respiration contributes substantially to urban carbon fluxes in the
greater Boston area, *Environmental Pollution*, 212, 433-439,
<https://doi.org/10.1016/j.envpol.2016.01.012>, 2016.
- Desai, A. R., Richardson, A. D., Moffat, A. M., Kattge, J., Hollinger, D. Y., Barr, A., Falge, E.,
Noormets, A., Papale, D., Reichstein, M., and Stauch, V. J.: Cross-site evaluation of eddy
covariance GPP and RE decomposition techniques, *Agr. Forest Meteorol.*, 148, 821-838,
<https://doi.org/10.1016/j.agrformet.2007.11.012>, 2008.
- Foken, T., Göckede, M., Mauder, M., Mahrt, L., Amiro, B., and Munger, W.: Post-Field Data
Quality Control, in: *Handbook of Micrometeorology: A Guide for Surface Flux Measurement
and Analysis*, edited by: Lee, X., Massman, W., and Law, B., Springer Netherlands,
Dordrecht, 181-208, https://doi.org/10.1007/1-4020-2265-4_9, 2004.
- Gately, C. K. and Hutyra, L. R.: Large Uncertainties in Urban-Scale Carbon Emissions, *J.
Geophys. Res.-Atmos.*, 122, 11242–11260, <https://doi.org/10.1002/2017jd027359>, 2017.
- Gourdji, S. M., Karion, A., Lopez-Coto, I., Ghosh, S., Mueller, K. L., Zhou, Y., Williams, C. A.,
Baker, I. T., Haynes, K. D., and Whetstone, J. R.: A Modified Vegetation Photosynthesis and
Respiration Model (VPRM) for the Eastern USA and Canada, Evaluated With Comparison
to Atmospheric Observations and Other Biospheric Models, *J. Geophys. Res.-Biogeosci.*,
127, e2021JG006290, <https://doi.org/10.1029/2021JG006290>, 2022.
- Grimmond, C. S. B. and Oke, T. R.: Aerodynamic Properties of Urban Areas Derived from
Analysis of Surface Form, *J. Appl. Meteorol.*, 38, 1262-1292, [https://doi.org/10.1175/1520-0450\(1999\)038<1262:APOUAD>2.0.CO;2](https://doi.org/10.1175/1520-0450(1999)038<1262:APOUAD>2.0.CO;2), 1999.
- Guo, S., Hu, M., Zamora, M. L., Peng, J., Shang, D., Zheng, J., Du, Z., Wu, Z., Shao, M., and
Zeng, L.: Elucidating severe urban haze formation in China, *P. Natl. Acad. Sci. USA*, 111,
17373-17378, <https://doi.org/10.1073/pnas.1419604111>, 2014.
- Hardiman, B. S., Wang, J. A., Hutyra, L. R., Gately, C. K., Getson, J. M., and Friedl, M. A.:
Accounting for urban biogenic fluxes in regional carbon budgets, *Sci. Total Environ.*, 592,
366-372, <https://doi.org/10.1016/j.scitotenv.2017.03.028>, 2017.
- Hilland, R., Hashemi, J., Stagakis, S., Brunner, D., Constantin, L., Kljun, N., Kunz, A. K.,
Molinier, B., Hammer, S., Emmenegger, L., and Christen, A.: Sectoral attribution of
greenhouse gas and pollutant emissions using multi-species eddy covariance on a tall tower



- 1296 in Zurich, Switzerland, *Atmos. Chem. Phys.*, 25, 14279–14299, [https://doi.org/10.5194/acp-](https://doi.org/10.5194/acp-25-14279-2025)
1297 [25-14279-2025](https://doi.org/10.5194/acp-25-14279-2025), 2025.
- 1298 Horne, J. P., Richardson, S. J., Murphy, S. L., Kenion, H. C., Haupt, B. J., Ahlswede, B. J.,
1299 Miles, N. L., and Davis, K. J.: The INFLUX network – eddy covariance in and around an
1300 urban environment, *Earth Syst. Sci. Data*, 18, 823–843, [https://doi.org/10.5194/essd-18-823-](https://doi.org/10.5194/essd-18-823-2026)
1301 [2026](https://doi.org/10.5194/essd-18-823-2026), 2026.
- 1302 Huete, A., Didan, K., Miura, T., Rodriguez, E. P., Gao, X., and Ferreira, L. G.: Overview of the
1303 radiometric and biophysical performance of the MODIS vegetation indices, *Remote Sens.*
1304 *Environ.*, 83, 195–213, [https://doi.org/10.1016/S0034-4257\(02\)00096-2](https://doi.org/10.1016/S0034-4257(02)00096-2), 2002.
- 1305 Ishidoya, S., Sugawara, H., Terao, Y., Kaneyasu, N., Aoki, N., Tsuboi, K., and Kondo, H.:
1306 O₂:CO₂ exchange ratio for net turbulent flux observed in an urban area of Tokyo, Japan, and
1307 its application to an evaluation of anthropogenic CO₂ emissions, *Atmos. Chem. Phys.*, 20,
1308 5293–5308, <https://doi.org/10.5194/acp-20-5293-2020>, 2020.
- 1309 Järvi, L., Nordbo, A., Junninen, H., Riikonen, A., Moilanen, J., Nikinmaa, E., and Vesala, T.:
1310 Seasonal and annual variation of carbon dioxide surface fluxes in Helsinki, Finland, in 2006–
1311 2010, *Atmos. Chem. Phys.*, 12, 8475–8489, <https://doi.org/10.5194/acp-12-8475-2012>, 2012.
- 1312 Järvi, L., Havu, M., Ward, H. C., Bellucco, V., McFadden, J. P., Toivonen, T., Heikinheimo, V.,
1313 Kolari, P., Riikonen, A., and Grimmond, C. S. B.: Spatial Modeling of Local-Scale Biogenic
1314 and Anthropogenic Carbon Dioxide Emissions in Helsinki, *J. Geophys. Res.-Atmos.*, 124,
1315 8363–8384, <https://doi.org/10.1029/2018JD029576>, 2019.
- 1316 Juchem, H., Maier, F., Levin, I., Jordan, A., Pöhler, D., Rosendahl, C., Della Coletta, J.,
1317 Preunkert, S., and Hammer, S.: Challenges and benefits of using NO_x as a quantitative proxy
1318 for fossil fuel CO₂ in an urban area based on radiocarbon measurements, *Atmos. Chem. Phys.*,
1319 25, 18373–18388, <https://doi.org/10.5194/acp-25-18373-2025>, 2025.
- 1320 Keenan, T. F., Migliavacca, M., Papale, D., Baldocchi, D., Reichstein, M., Torn, M., and
1321 Wutzler, T.: Widespread inhibition of daytime ecosystem respiration, *Nat. Ecol. Evol.*, 3,
1322 407–415, <https://doi.org/10.1038/s41559-019-0809-2>, 2019.
- 1323 Kljun, N., Calanca, P., Rotach, M. W., and Schmid, H. P.: A simple two-dimensional
1324 parameterisation for Flux Footprint Prediction (FFP), *Geosci. Model Dev.*, 8, 3695–3713,
1325 <https://doi.org/10.5194/gmd-8-3695-2015>, 2015.
- 1326 Kohonen, K. M., Kolari, P., Kooijmans, L. M. J., Chen, H., Seibt, U., Sun, W., and Mammarella,
1327 I.: Towards standardized processing of eddy covariance flux measurements of carbonyl
1328 sulfide, *Atmos. Meas. Tech.*, 13, 3957–3975, <https://doi.org/10.5194/amt-13-3957-2020>,
1329 2020.
- 1330 Kooijmans, L. M. J., Maseyk, K., Seibt, U., Sun, W., Vesala, T., Mammarella, I., Kolari, P.,
1331 Aalto, J., Franchin, A., Vecchi, R., Valli, G., and Chen, H.: Canopy uptake dominates
1332 nighttime carbonyl sulfide fluxes in a boreal forest, *Atmos. Chem. Phys.*, 17, 11453–11465,
1333 <https://doi.org/10.5194/acp-17-11453-2017>, 2017.
- 1334 Kunz, A. K., Hammer, S., Aigner, P., Bignotti, L., Borchardt, L., Chen, J., Della Coletta, J.,
1335 Emmenegger, L., Eritt, M., Gutiérrez, X., Hashemi, J., Hilland, R., Holst, C., Jordan, A.,
1336 Kljun, N., Kneißl, R., Lan, C., Legendre, V., Levin, I., Loubet, B., Mauder, M., Molinier, B.,
1337 Preunkert, S., Ramonet, M., Stagakis, S., and Christen, A.: ¹⁴C-based separation of fossil and
1338 non-fossil CO₂ fluxes in cities using relaxed eddy accumulation: results from tall-tower
1339 measurements in Zurich, Paris, and Munich, *Atmos. Chem. Phys.*, 26, 4967–5003,



- 1340 <https://doi.org/10.5194/acp-26-4967-2026>, 2026.
- 1341 Lang, N., Jetz, W., Schindler, K., and Wegner, J. D.: A high-resolution canopy height model of
1342 the Earth, *Nat Ecol Evol*, 7, 1778-1789, <https://doi.org/10.1038/s41559-023-02206-6>, 2023.
- 1343 Lasslop, G., Reichstein, M., Papale, D., Richardson, A. D., Arneth, A., Barr, A., Stoy, P., and
1344 Wohlfahrt, G.: Separation of net ecosystem exchange into assimilation and respiration using
1345 a light response curve approach: critical issues and global evaluation, *Global Change Biol.*,
1346 16, 187-208, <https://doi.org/10.1111/j.1365-2486.2009.02041.x>, 2010.
- 1347 Lee, K., Hong, J.-W., Kim, J., and Hong, J.: Partitioning of net CO₂ exchanges at the city-
1348 atmosphere interface into biotic and abiotic components, *MethodsX*, 8, 101231,
1349 <https://doi.org/10.1016/j.mex.2021.101231>, 2021.
- 1350 Liu, H. Z., Feng, J. W., Järvi, L., and Vesala, T.: Four-year (2006–2009) eddy covariance
1351 measurements of CO₂ flux over an urban area in Beijing, *Atmos. Chem. Phys.*, 12, 7881-
1352 7892, <https://doi.org/10.5194/acp-12-7881-2012>, 2012.
- 1353 Mahadevan, P., Wofsy, S. C., Matross, D. M., Xiao, X., Dunn, A. L., Lin, J. C., Gerbig, C.,
1354 Munger, J. W., Chow, V. Y., and Gottlieb, E. W.: A satellite-based biosphere parameterization
1355 for net ecosystem CO₂ exchange: Vegetation Photosynthesis and Respiration Model (VPRM),
1356 *Global Biogeochem. Cy.*, 22, <https://doi.org/10.1029/2006GB002735>, 2008.
- 1357 Matese, A., Gioli, B., Vaccari, F. P., Zaldei, A., and Miglietta, F.: Carbon Dioxide Emissions of
1358 the City Center of Firenze, Italy: Measurement, Evaluation, and Source Partitioning, *J. Appl.*
1359 *Meteorol. Clim.*, 48, 1940-1947, <https://doi.org/10.1175/2009JAMC1945.1>, 2009.
- 1360 Matthews, B. and Schume, H.: Tall tower eddy covariance measurements of CO₂ fluxes in
1361 Vienna, Austria, *Atmos. Environ.*, 274, 118941,
1362 <https://doi.org/10.1016/j.atmosenv.2022.118941>, 2022.
- 1363 Menzer, O. and McFadden, J. P.: Statistical partitioning of a three-year time series of direct
1364 urban net CO₂ flux measurements into biogenic and anthropogenic components, *Atmos.*
1365 *Environ.*, 170, 319-333, <https://doi.org/10.1016/j.atmosenv.2017.09.049>, 2017.
- 1366 Miller, J. B., Lehman, S. J., Verhulst, K. R., Miller, C. E., Duren, R. M., Yadav, V., Newman,
1367 S., and Sloop, C. D.: Large and seasonally varying biospheric CO₂ fluxes in the Los Angeles
1368 megacity revealed by atmospheric radiocarbon, *P. Natl. Acad. Sci. USA*, 117, 26681-26687,
1369 <https://doi.org/10.1073/pnas.2005253117>, 2020.
- 1370 Monteith, J. L.: Solar radiation and productivity in tropical ecosystems, *J. Appl. Ecol.*, 9, 747-
1371 766, <https://doi.org/10.2307/2401901>, 1972.
- 1372 Myneni, R. B. and Williams, D. L.: On the relationship between FAPAR and NDVI, *Remote*
1373 *Sens. Environ.*, 49, 200-211, [https://doi.org/10.1016/0034-4257\(94\)90016-7](https://doi.org/10.1016/0034-4257(94)90016-7), 1994.
- 1374 Neigh, C.: HLS Operational Land Imager Vegetation Indices Daily Global 30m v2.0, NASA
1375 Land Processes Distributed Active Archive Center [dataset],
1376 https://doi.org/10.5067/HLS/HLSSL30_V1.002, 2025.
- 1377 Neigh, C., Ju, J., Roger, J.-C., Skakun, S., Vermote, E., Claverie, M., Dungan, J., Yin, Z., Freitag,
1378 B., and Justice, C.: HLS Sentinel-2 Multi-spectral Instrument Surface Reflectance Daily
1379 Global 30m v2.0, NASA Land Processes Distributed Active Archive Center [dataset],
1380 <https://doi.org/10.5067/HLS/HLSS30.002>, 2021.
- 1381 Nordbo, A., Järvi, L., Haapanala, S., Wood, C. R., and Vesala, T.: Fraction of natural area as
1382 main predictor of net CO₂ emissions from cities, *Geophys. Res. Lett.*, 39,
1383 <https://doi.org/10.1029/2012GL053087>, 2012.



- 1384 Oda, T., Maksyutov, S., and Andres, R. J.: The Open-source Data Inventory for Anthropogenic
1385 CO₂, version 2016 (ODIAC2016): a global monthly fossil fuel CO₂ gridded emissions data
1386 product for tracer transport simulations and surface flux inversions, *Earth Syst. Sci. Data*, 10,
1387 87-107, <https://doi.org/10.5194/essd-10-87-2018>, 2018.
- 1388 Papale, D., Reichstein, M., Canfora, E., Aubinet, M., Bernhofer, C., Longdoz, B., Kutsch, W.,
1389 Rambal, S., Valentini, R., and Vesala, T.: Towards a more harmonized processing of eddy
1390 covariance CO₂ fluxes: algorithms and uncertainty estimation, *Biogeosciences*, 3, 571–583,
1391 <https://doi.org/10.5194/bg-3-571-2006>, 2006.
- 1392 Parrish, D. D.: Critical evaluation of US on-road vehicle emission inventories, *Atmos. Environ.*,
1393 40, 2288-2300, <https://doi.org/10.1016/j.atmosenv.2005.11.033>, 2006.
- 1394 Pérez-Ruiz, E. R., Vivoni, E. R., and Templeton, N. P.: Urban land cover type determines the
1395 sensitivity of carbon dioxide fluxes to precipitation in Phoenix, Arizona, *PLoS One*, 15,
1396 e0228537, <https://doi.org/10.1371/journal.pone.0228537>, 2020.
- 1397 Reichstein, M., Falge, E., Baldocchi, D., Papale, D., Aubinet, M., Berbigier, P., Bernhofer, C.,
1398 Buchmann, N., Gilmanov, T., Granier, A., Grünwald, T., Havránková, K., Ilvesniemi, H.,
1399 Janous, D., Knohl, A., Laurila, T., Lohila, A., Loustau, D., Matteucci, G., Meyers, T.,
1400 Miglietta, F., Ourcival, J.-M., Pumpanen, J., Rambal, S., Rotenberg, E., Sanz, M., Tenhunen,
1401 J., Seufert, G., Vaccari, F., Vesala, T., Yakir, D., and Valentini, R.: On the separation of net
1402 ecosystem exchange into assimilation and ecosystem respiration: review and improved
1403 algorithm, *Global Change Biol.*, 11, 1424-1439, <https://doi.org/10.1111/j.1365-2486.2005.001002.x>, 2005.
- 1405 Richardson, A. D. and Hollinger, D. Y.: A method to estimate the additional uncertainty in gap-
1406 filled NEE resulting from long gaps in the CO₂ flux record, *Agr. Forest Meteorol.*, 147, 199-
1407 208, <https://doi.org/10.1016/j.agrformet.2007.06.004>, 2007.
- 1408 Rotach, M. W.: On the influence of the urban roughness sublayer on turbulence and dispersion,
1409 *Atmos. Environ.*, 33, 4001-4008, [https://doi.org/10.1016/S1352-2310\(99\)00141-7](https://doi.org/10.1016/S1352-2310(99)00141-7), 1999.
- 1410 Running, S. W., Nemani, R. R., Heinsch, F. A., Zhao, M., Reeves, M., and Hashimoto, H.: A
1411 Continuous Satellite-Derived Measure of Global Terrestrial Primary Production, *BioScience*,
1412 54, 547-560, [https://doi.org/10.1641/0006-3568\(2004\)054\[0547:Acsmog\]2.0.Co;2](https://doi.org/10.1641/0006-3568(2004)054[0547:Acsmog]2.0.Co;2), 2004.
- 1413 Sabbatini, S., Mammarella, I., Arriga, N., Fratini, G., Graf, A., Hörtnagl, L., Ibrom, A., Longdoz,
1414 B., Mauder, M., and Merbold, L.: Eddy covariance raw data processing for CO₂ and energy
1415 fluxes calculation at ICOS ecosystem stations, *International Agrophysics*, 32, 495-515,
1416 <https://doi.org/10.1515/intag-2017-0043>, 2018.
- 1417 Sargent, M., Barrera, Y., Nehrkorn, T., Hutyrá, L. R., Gately, C. K., Jones, T., McKain, K.,
1418 Sweeney, C., Hegarty, J., Hardiman, B., Wang, J. A., and Wofsy, S. C.: Anthropogenic and
1419 biogenic CO₂ fluxes in the Boston urban region, *P. Natl. Acad. Sci. USA*, 115, 7491-7496,
1420 <https://doi.org/10.1073/pnas.1803715115>, 2018.
- 1421 Sellers, P. J.: Canopy reflectance, photosynthesis and transpiration, *International Journal of*
1422 *Remote Sensing*, 6, 1335-1372, <https://doi.org/10.1080/01431168508948283>, 1985.
- 1423 Soininen, J., Kohonen, K.-M., Rantala, P., Kulmala, L., Aaltonen, H., and Järvi, L.: Carbon
1424 uptake of an urban green space inferred from carbonyl sulfide fluxes, *npj Clim. Atmos. Sci.*,
1425 8, <https://doi.org/10.1038/s41612-025-00958-5>, 2025.
- 1426 Squires, F. A., Nemitz, E., Langford, B., Wild, O., Drysdale, W. S., Acton, W. J. F., Fu, P.,
1427 Grimmond, C. S. B., Hamilton, J. F., Hewitt, C. N., Hollaway, M., Kotthaus, S., Lee, J.,



- 1428 Metzger, S., Pingintha-Durden, N., Shaw, M., Vaughan, A. R., Wang, X., Wu, R., Zhang, Q.,
1429 and Zhang, Y.: Measurements of traffic-dominated pollutant emissions in a Chinese megacity,
1430 *Atmos. Chem. Phys.*, 20, 8737-8761, <https://doi.org/10.5194/acp-20-8737-2020>, 2020.
- 1431 Tao, X., Cui, J., Dai, Y., Wang, Z., and Xu, X.: Soil respiration responses to soil physiochemical
1432 properties in urban different green-lands: A case study in Hefei, China, *International Soil and*
1433 *Water Conservation Research*, 4, 224-229, <https://doi.org/10.1016/j.iswcr.2016.08.001>, 2016.
- 1434 Turnbull, J. C., Tans, P. P., Lehman, S. J., Baker, D., Conway, T. J., Chung, Y., Gregg, J., Miller,
1435 J. B., Southon, J. R., and Zhou, L. X.: Atmospheric observations of carbon monoxide and
1436 fossil fuel CO₂ emissions from East Asia, *J. Geophys. Res.-Atmos.*, 116,
1437 <https://doi.org/10.1029/2011JD016691>, 2011.
- 1438 Ueyama, M. and Takano, T.: A decade of CO₂ flux measured by the eddy covariance method
1439 including the COVID-19 pandemic period in an urban center in Sakai, Japan, *Environmental*
1440 *Pollution*, 304, 119210, <https://doi.org/10.1016/j.envpol.2022.119210>, 2022.
- 1441 Velasco, E. and Roth, M.: Cities as Net Sources of CO₂: Review of Atmospheric CO₂ Exchange
1442 in Urban Environments Measured by Eddy Covariance Technique, *Geography Compass*, 4,
1443 1238-1259, <https://doi.org/10.1111/j.1749-8198.2010.00384.x>, 2010.
- 1444 Velasco, E., Roth, M., Tan, S., Quak, M., Nabarro, S., and Norford, L.: The role of vegetation
1445 in the CO₂ flux from a tropical urban neighbourhood, *Atmos. Chem. Phys.*, 13, 10185-10202,
1446 <https://doi.org/10.5194/acp-13-10185-2013>, 2013.
- 1447 Wang, L., Yu, M., Ye, S., and Yan, J.: Seasonal patterns of carbon and water flux responses to
1448 precipitation and solar radiation variability in a subtropical evergreen forest, South China,
1449 *Agr. Forest Meteorol.*, 342, 109760, <https://doi.org/10.1016/j.agrformet.2023.109760>, 2023.
- 1450 Ward, H. C., Kotthaus, S., Grimmond, C. S. B., Bjorkegren, A., Wilkinson, M., Morrison, W.
1451 T. J., Evans, J. G., Morison, J. I. L., and Iamarino, M.: Effects of urban density on carbon
1452 dioxide exchanges: Observations of dense urban, suburban and woodland areas of southern
1453 England, *Environmental Pollution*, 198, 186-200,
1454 <https://doi.org/10.1016/j.envpol.2014.12.031>, 2015.
- 1455 Webb, E. K., Pearman, G. I., and Leuning, R.: Correction of flux measurements for density
1456 effects due to heat and water vapour transfer, *Quarterly Journal of the Royal Meteorological*
1457 *Society*, 106, 85-100, <https://doi.org/10.1002/qj.49710644707>, 1980.
- 1458 Wehr, R., Munger, J. W., McManus, J. B., Nelson, D. D., Zahniser, M. S., Davidson, E. A.,
1459 Wofsy, S. C., and Saleska, S. R.: Seasonality of temperate forest photosynthesis and daytime
1460 respiration, *Nature*, 534, 680-683, <https://doi.org/10.1038/nature17966>, 2016.
- 1461 Wu, D., Lin, J. C., Duarte, H. F., Yadav, V., Parazoo, N. C., Oda, T., and Kort, E. A.: A model
1462 for urban biogenic CO₂ fluxes: Solar-Induced Fluorescence for Modeling Urban biogenic
1463 Fluxes (SMUrF v1), *Geosci. Model Dev.*, 14, 3633-3661, <https://doi.org/10.5194/gmd-14-3633-2021>, 2021.
- 1465 Wu, K., Davis, K. J., Miles, N. L., Richardson, S. J., Lauvaux, T., Sarmiento, D. P., Balashov,
1466 N. V., Keller, K., Turnbull, J., and Gurney, K. R.: Source decomposition of eddy-covariance
1467 CO₂ flux measurements for evaluating a high-resolution urban CO₂ emissions inventory,
1468 *Environ. Res. Lett.*, 17, 074035, <https://doi.org/10.1088/1748-9326/ac7c29>, 2022.
- 1469 Wutzler, T., Lucas-Moffat, A., Migliavacca, M., Knauer, J., Sickel, K., Šigut, L., Menzer, O.,
1470 and Reichstein, M.: Basic and extensible post-processing of eddy covariance flux data with
1471 REddyProc, *Biogeosciences*, 15, 5015-5030, <https://doi.org/10.5194/bg-15-5015-2018>, 2018.



- 1472 Xiao, X., Hollinger, D., Aber, J., Goltz, M., Davidson, E. A., Zhang, Q., and Moore, B.:
1473 Satellite-based modeling of gross primary production in an evergreen needleleaf forest,
1474 Remote Sens. Environ., 89, 519-534, <https://doi.org/10.1016/j.rse.2003.11.008>, 2004.
- 1475 Xu, R., Tong, D., Xiao, Q., Qin, X., Chen, C., Yan, L., Cheng, J., Cui, C., Hu, H., and Liu, W.:
1476 MEIC-global-CO₂: A new global CO₂ emission inventory with highly-resolved source
1477 category and sub-country information, Sci. China Earth Sci., 67, 450-465,
1478 <https://doi.org/10.1007/s11430-023-1230-3>, 2024.
- 1479 Xu, X.: National hourly population density spatial distribution dataset, Resource and
1480 Environmental Science Data Registration and Publication System [dataset],
1481 <https://doi.org/10.12078/2022091601>, 2022.
- 1482 Xueref-Remy, I., Dieudonné, E., Vuillemin, C., Lopez, M., Lac, C., Schmidt, M., Delmotte, M.,
1483 Chevallier, F., Ravetta, F., Perrussel, O., Ciais, P., Bréon, F.-M., Broquet, G., Ramonet, M.,
1484 Spain, T. G., and Ampe, C.: Diurnal, synoptic and seasonal variability of atmospheric CO₂ in
1485 the Paris megacity area, Atmos. Chem. Phys., 18, 3335-3362, [https://doi.org/10.5194/acp-](https://doi.org/10.5194/acp-18-3335-2018)
1486 [18-3335-2018](https://doi.org/10.5194/acp-18-3335-2018), 2018.
- 1487 Yang, S., Kong, F., Yin, H., Zou, J., Järvi, L., and Sun, J.: Short-term responses of urban forest
1488 carbon dynamics to combined heatwave and drought in subtropical China, Agr. Forest
1489 Meteorol., 372, 110728, <https://doi.org/10.1016/j.agrformet.2025.110728>, 2025.
- 1490 Yi, Z., Fu, S., Yi, W., Zhou, G., Mo, J., Zhang, D., Ding, M., Wang, X., and Zhou, L.:
1491 Partitioning soil respiration of subtropical forests with different successional stages in south
1492 China, Forest Ecology and Management, 243, 178-186,
1493 <https://doi.org/10.1016/j.foreco.2007.02.022>, 2007.
- 1494 Zhang, J., Liang, Y., Pei, C., Huang, B., Huang, Y., Lian, X., Song, S., Cheng, C., Wu, C., Zhou,
1495 Z., Li, J., and Li, M.: Atmospheric CO₂ dynamics in a coastal megacity: spatiotemporal
1496 patterns, sea-land breeze impacts, and anthropogenic-biogenic emission partitioning, Atmos.
1497 Chem. Phys., 26, 3253-3276, <https://doi.org/10.5194/acp-26-3253-2026>, 2026.
- 1498 Zhang, K., Gong, Y., Fa, H., and Zhao, M.: CO₂ Flux Characteristics of Different Plant
1499 Communities in a Subtropical Urban Ecosystem, Sustainability, 11, 4879,
1500 <https://doi.org/10.3390/su11184879>, 2019.
- 1501 Zhang, X., Liu, L., Zhao, T., Zhang, W., Guan, L., Bai, M., and Chen, X.: GLC_FCS10: a global
1502 10 m land-cover dataset with a fine classification system from Sentinel-1 and Sentinel-2
1503 time-series data in Google Earth Engine, Earth Syst. Sci. Data, 17, 4039-4062,
1504 <https://doi.org/10.5194/essd-17-4039-2025>, 2025.
- 1505