



1 **A full-coverage satellite-based global atmospheric CO₂ dataset at 0.05°**
2 **resolution from 2015 to 2021 for exploring global carbon dynamics**

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26



27 **Abstract**

28 The irreversible trend for global warming underscores the necessity for accurate
29 monitoring and analysis of atmospheric carbon dynamics on a global scale. Carbon
30 satellites hold significant potential for atmospheric CO₂ monitoring. However, existing
31 studies on global CO₂ are constrained by coarse resolution (ranging from 0.25° to 2°)
32 and limited spatial coverage. In this study, we developed a new global dataset of
33 column-averaged dry-air mole fraction of CO₂ (XCO₂) at 0.05° resolution with full
34 coverage using carbon satellite observations, multi-source satellite products, and an
35 improved deep learning model. We then investigated changes in global atmospheric
36 CO₂ and anomalies from 2015 to 2021. The reconstructed XCO₂ products show a better
37 agreement with Total Carbon Column Observing Network (TCCON) measurements,
38 with R² of 0.92 and RSME of 1.54 ppm. The products also provide more accurate
39 information on the global and regional spatial patterns of XCO₂ compared to origin
40 carbon satellite monitoring and previous XCO₂ products. The global pattern of XCO₂
41 exhibited a distinct increasing trend with a growth rate of 2.32 ppm/year, reaching
42 414.00 ppm in 2021. Globally, XCO₂ showed obvious spatial variability across
43 different latitudes and continents. Higher XCO₂ concentrations were primarily
44 observed in the Northern Hemisphere, particularly in regions with intensive
45 anthropogenic activity, such as East Asia and North America. We also validated the
46 effectiveness of our XCO₂ products in detecting intensive CO₂ emission sources. The
47 XCO₂ dataset is publicly accessible on the Zenodo platform at
48 <https://doi.org/10.5281/zenodo.12706142> (Wang et al., 2024). Our findings represent a
49 promising advancement in monitoring carbon emission across various countries and
50 enhancing the understanding of global carbon dynamics.

51

52 **Keywords:** Atmospheric carbon dioxide; Satellite carbon monitoring; Deep learning;
53 OCO-2/3

54

55 **1. Introduction**

56 Carbon dioxide (CO₂) is a primary greenhouse gas (GHG). Anthropogenic
57 activities and land use change since the industrial revolution have led to a marked
58 increase in atmospheric CO₂, which is widely considered to be a major contributor to



59 climate change, reaching a record-high of 414.71 parts per million (ppm) in 2021
60 (Friedlingstein et al., 2022). The damaging global climate change caused by
61 atmospheric increases in CO₂ is severe and irreversible (IPCC, 2023; Kemp et al., 2022;
62 Solomon et al., 2009). Consequently, the Paris Agreement announced to hold “the
63 increase in the global average temperature to well below 2°C above pre-industrial levels”
64 and pursue efforts “to limit the temperature increase to 1.5°C above pre-industrial
65 levels.” It was also determined that the joined parties should submit their nationally
66 determined contributions (NDCs) to reduce CO₂ emissions.

67 Accurate monitoring of atmospheric CO₂ concentrations is crucial for measuring
68 global CO₂ emissions mitigation as well as characterizing terrestrial carbon change.
69 Currently, ground-based and airborne platform-based atmospheric CO₂ observation
70 networks, such as the Total Carbon Column Observing Network (TCCON,
71 <https://tccodata.org/>), are capable of providing CO₂ measurements with high accuracy
72 (Petzold et al., 2016; Wunch et al., 2011, 2010). However, these observation networks
73 are insufficient to fully explore the spatiotemporal patterns of atmospheric CO₂ at a
74 global scale. The launch of a series of carbon observation satellites in recent years has
75 provided favorable opportunities for continuous and large-scale atmospheric CO₂
76 observation (Buchwitz et al., 2015; Hammerling et al., 2012). The Scanning Imaging
77 Absorption Spectrometer for Atmospheric Chartography (SCIAMACHY) onboard
78 EnviSat was one of the first instruments to monitor the atmospheric column-averaged
79 dry-air mole fraction of CO₂ (XCO₂) (Bovensmann et al., 1999). The Greenhouse Gases
80 Observing Satellite (GOSAT) launched by Japan utilized the Thermal And Near-
81 Infrared Sensor for carbon Observation (TANSO) instrument to monitor XCO₂ globally,
82 providing products with a spatial resolution of 10 km every three days (Butz et al.,
83 2011). The Orbiting Carbon Observatory-2 (OCO-2) and OCO-3 launched by NASA
84 provide XCO₂ measurements at a finer spatial resolution (Crisp et al., 2017; Eldering
85 et al., 2017). These sensors are considered among the best for XCO₂ observation,
86 featuring larger overlapping swaths that cover areas of ~20×80 km² and exhibiting the
87 least retrieval absolute bias, measuring less than 0.4 ppm (Eldering et al., 2019; Taylor
88 et al., 2020). However, the narrow swath of the sensor can only cover limited spatial
89 areas, and caused by the cloud and aerosol contaminations, the data from OCO-2/3
90 always contain large amount of missing values (Taylor et al., 2016; Crisp et al., 2017).
91 These limitations obstacle the better understanding of the atmosphere-land carbon cycle
92 over large spatial scale based on satellite observation.



93 Consequently, several studies have concentrated on generating spatially
94 continuous XCO₂ products based on satellite observations (He et al., 2022; Siabi et al.,
95 2019; Zhang and Liu, 2023). One potential solution is the application of diverse
96 interpolation methods (He et al., 2020; Zeng et al., 2014). Hammerling et al. (2012)
97 mapped the global distribution of CO₂ based on OCO-2 and the geostatistical method.
98 Zeng et al. (2014) developed a gap-filling model based on the space-time kriging to
99 obtain gap-filled GOAST XCO₂ data in China. However, their results encounter large
100 uncertainty in regions with sparse data coverage, due to algorithmic constraints and the
101 coarse spatial resolution of the original data. Recently, data fusion techniques have
102 gained recognition as an effective method for obtaining full-coverage XCO₂ data
103 (Sheng et al., 2023; He et al., 2022; Siabi et al., 2019; Zhang and Liu, 2023). These
104 techniques can be broadly categorized into two groups based on their underlying
105 principles. The first category leverages the spatiotemporal correlation inherent in multi-
106 source XCO₂ data, fusing them based on this spatiotemporal information (Wang et al.,
107 2023; Sheng et al., 2023). For instance, Wang et al. (2023) introduced a spatiotemporal
108 self-supervised fusion model and generate seamless global XCO₂ data at a spatial
109 resolution of 0.25°. The second category is regression-based methods, which aim to fill
110 the gap in XCO₂ data by capturing the nonlinear relationship between multi-source
111 XCO₂ measurements and related covariates (He et al., 2022; Siabi et al., 2019; Zhang
112 and Liu, 2023). This category includes a range of specific methodologies, from
113 traditional statistical and geostatistical models to advanced machine learning models.
114 Siabi et al (2019) employed the Artificial Neural Network (ANN) to establish
115 correlation between XCO₂ and eight environmental variables. Zhang and Liu (2023)
116 utilized the convolution neural networks (CNN) coupled with attention mechanisms to
117 produce full-coverage XCO₂ data across China.

118 Despite significant efforts to generate seamless XCO₂ products using satellite
119 observations, several challenges and limitations persist. Firstly, due to the sparse
120 distribution of satellite XCO₂ data, previous studies have relied on assimilation and
121 reanalysis XCO₂ data as covariates. This reliance often results in final products that
122 closely mirror the assimilation and reanalysis results, leading to an oversmoothed
123 distribution that undermines the high-resolution advantages of satellite data.
124 Furthermore, most studies that employ regression models to estimate full-coverage
125 XCO₂ are limited to regional or national scales due to the weak transferability of these

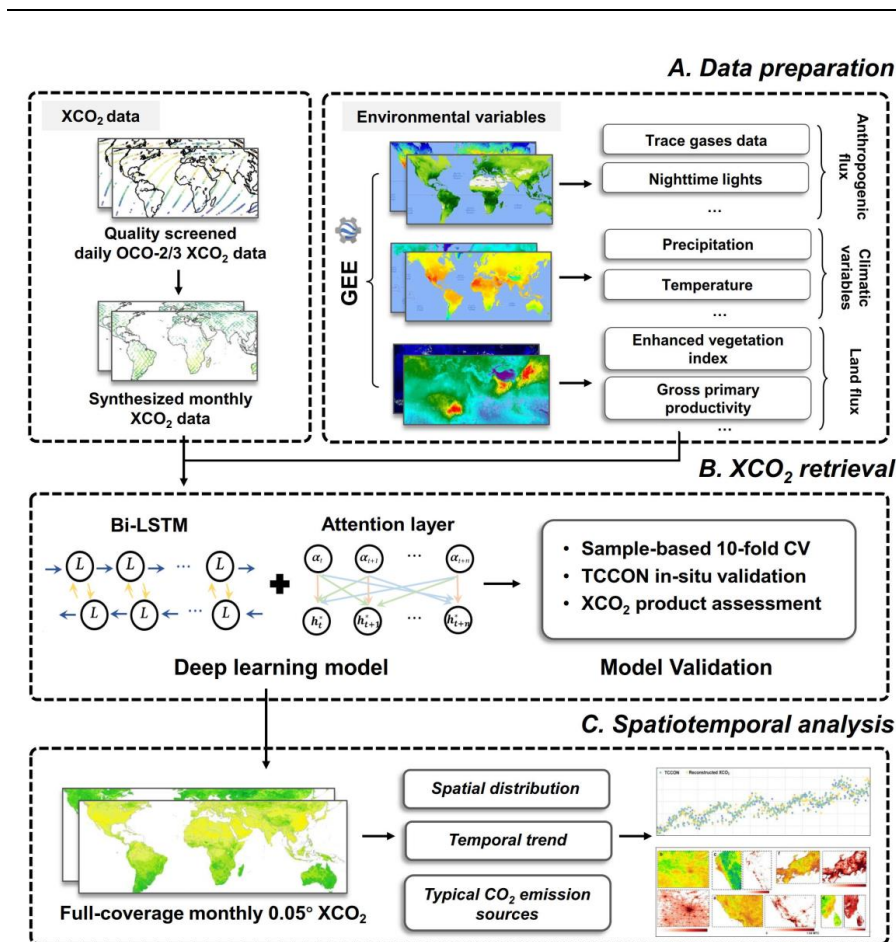


126 models. Consequently, there is a scarcity of research investigating global patterns of
127 XCO₂ based on statistical models. Additionally, the current generation of XCO₂
128 products typically features a coarse spatial resolution, ranging from approximately 0.25°
129 to 2°. In conclusion, there is an urgent need to develop global full-coverage XCO₂
130 products with a fine spatial resolution. This development should leverage satellite
131 carbon monitoring and advanced methods that exhibit spatiotemporal transferability to
132 overcome the aforementioned limitations.

133 In this study, we leveraged time-series OCO-2/3 XCO₂ data and various related
134 environmental variables retrieved from multi-source satellites to generate global full-
135 coverage XCO₂ products. The advanced deep learning method was adopted to model
136 time-series XCO₂ and incorporate terrestrial flux, anthropogenic flux and climatic
137 impacts into the parameterization process. Our XCO₂ products achieved full global
138 coverage with a spatial resolution of 0.05° and a monthly temporal resolution from 2015
139 to 2021. We also validated our XCO₂ products against in-situ XCO₂ data and other
140 XCO₂ products. Based on our high-resolution products, we explored the spatial and
141 temporal pattern of atmospheric CO₂ globally and identified regions with intense CO₂
142 emission. Our findings aim to enhance the understanding of carbon dynamics on a
143 global scale through data reconstruction and analysis.

144 **2. Materials and methods**

145 In this study, we utilized Google Earth Engine (GEE) to integrate OCO-2/3 XCO₂
146 data and multiple environmental variables as data inputs. In addition, the attention-
147 based Bidirectional Long Short-Term Memory (At-BiLSTM) model was trained for
148 building the relationship between OCO-2/3 XCO₂ and the related environmental
149 variables. Then, we reconstructed the global monthly XCO₂ and validated the accuracy
150 of the products against TCCON XCO₂ data and the original OCO-2/3 XCO₂ data. We
151 also analyzed the spatial and temporal variation of XCO₂ over the globe and detect the
152 intense CO₂ emission regions. The methodology framework is shown in Fig.1.



153

154 **Figure 1.** The workflow for mapping and exploring global XCO₂ dynamics and drivers.

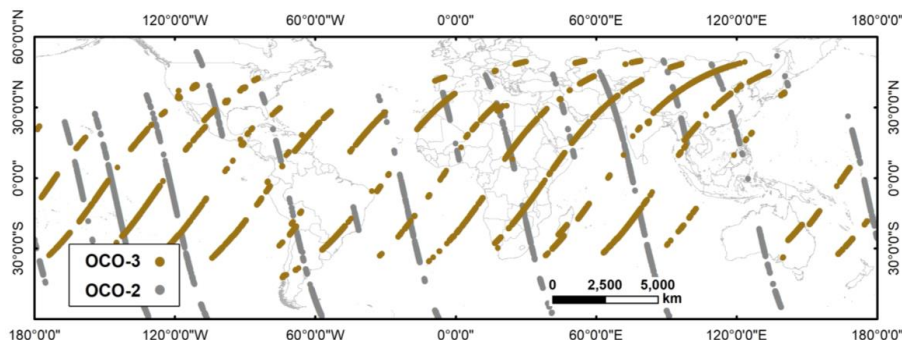
155 **2.1 Datasets**

156 **2.1.1 OCO XCO₂ data**

157 In this study, we utilized the satellite-based XCO₂ data from OCO-2 and OCO-3,
 158 covering the period from December 2014 to December 2021. The OCO-2/3 measure at
 159 three near-infrared wavelength bands, that are 0.76 μm Oxygen A-band, 1.61 μm weak
 160 CO₂, and 2.06 μm strong CO₂ bands (Crisp et al., 2004). The full physics retrieval
 161 algorithm was used to retrieve the XCO₂ based on the observation of the two satellites
 162 (Crisp et al., 2021). Previous studies (Taylor et al., 2023) suggested that the OCO-2 and
 163 OCO-3 XCO₂ measurements are in broad consistency and can therefore be used
 164 together in scientific analyses. The OCO-3 Level 2 XCO₂ Lite version 10.4r data
 165 (OCO3_L2_Lite_FP V10.4r) from 2020 to 2021 and the OCO-2 Level 2 XCO₂ Lite



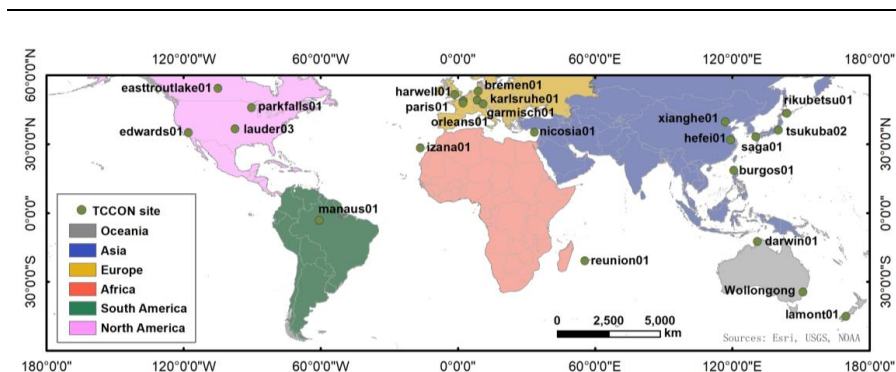
166 version 11r (OCO2_L2_Lite_FP V11r) from 2015 to 2019 were downloaded from
167 Goddard Earth Sciences Data and Information Services Center (GES DISC,
168 <https://disc.gsfc.nasa.gov/>). The products were aggregated as a daily file (Fig. 2) with a
169 spatial resolution of $2.25 \text{ km} \times 1.29 \text{ km}$ (O'Dell et al., 2018). The XCO₂ data were
170 quality filtered, and only good-quality data (i.e., xco2_quality_flag=0) were considered.
171 To generate the monthly products with a spatial resolution of $0.05^\circ \times 0.05^\circ$, we
172 converted the daily data to monthly data by averaging the sparse XCO₂ data within a
173 range of $0.05^\circ \times 0.05^\circ$ over one month.



174
175 **Figure 2.** Footprints of OCO-2 and OCO-3 XCO₂ data on 20th January 2018 and 4th
176 December 2021 (with quality filtering) as examples.
177

178 2.1.2 TCCON XCO₂ data

179 The Total Carbon Column Observing Network (TCCON) is a global network for
180 measuring atmospheric CO₂, methane (CH₄), carbon monoxide (CO) and other trace
181 gases in the atmosphere. The XCO₂ data from TCCON were demonstrated to have high
182 accuracy with $\sim 0.2\%$ of XCO₂ (Wunch et al., 2011). Consequently, the data have been
183 used widely for the validation of satellite observations such as OCO-2, OCO-3 and
184 GOSAT (Deng et al., 2016; Wunch et al., 2017). In this research, we used the GGG2014
185 and GGG2020 datasets from 23 sites (Fig. 3 and Table 1) around the world to validate
186 the reconstructed XCO₂ products.



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 189
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Figure 3. The locations of the TCCON sites.

Table 1. The information on the TCCON *in situ* stations.

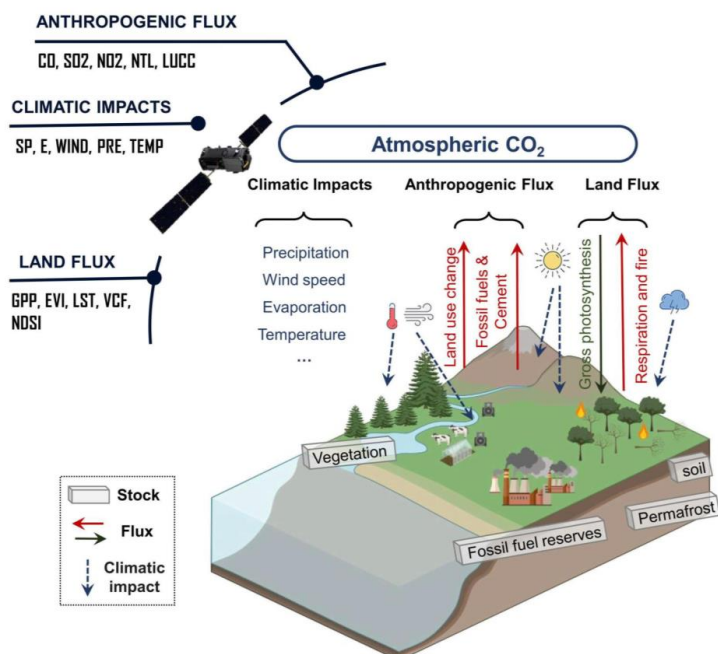
ID	Site name	Longitude	Latitude	Start date	End date
1	saga01 (JP)	130.29	33.24	2011-07-28	2021-06-30
2	xianghe01 (PRC)	116.96	39.80	2018-06-14	2022-04-09
3	burgos01 (PH)	120.65	18.53	2017-03-03	2021-08-20
4	harwell01 (UK)	-1.32	51.57	2021-05-30	2022-05-22
5	bremen01 (DE)	8.85	53.10	2009-01-06	2021-06-24
6	tsukuba02 (JP)	140.12	36.05	2014-03-28	2021-03-31
7	lauder03 (NZ)	-97.49	36.60	2018-10-02	2022-11-14
8	edwards01 (US)	-117.88	34.96	2013-07-20	2022-12-25
9	nicosia01 (CY)	33.38	35.14	2019-09-06	2021-06-01
10	izana01 (ES)	-16.5	28.31	2014-01-02	2022-10-31
11	orleans01 (FR)	2.11	47.96	2009-09-06	2022-04-24
12	hefei01 (PRC)	119.17	31.90	2015-11-02	2020-12-31
13	easttroutlake01 (CA)	-104.99	54.35	2016-10-03	2022-08-13
14	karlsruhe01 (DE)	8.44	49.10	2014-01-15	2023-01-20
15	paris01 (FR)	2.36	48.85	2014-09-23	2022-03-28
16	garmisch01 (DE)	11.06	47.48	2007-07-18	2021-10-18
17	rikubetsu01 (JP)	143.77	43.46	2014-06-24	2021-06-30
18	lamont01 (US)	169.68	-45.04	2011-04-16	2022-12-19
19	reunion01 (RE)	55.48	-20.90	2015-03-01	2020-07-18
20	darwin01 (AU)	130.93	-12.46	2005-08-28	2020-04-30
21	Wollongong (AU)	150.88	-34.41	2008-06-26	2020-06-30
22	Manaus01(BR)	-60.60	-3.21	2014-09-30	2015-07-27
23	parkfalls01 (US)	-90.27	45.94	2004-06-02	2020-12-29



191 JP: Japan, DE: Germany, FI: Finland, FR: French, RE: Réunion Island, AU: Australia,
192 BR: Brazil; US: United States, PRC: People's Republic of China, NO: Norway, CY:
193 Cyprus, NZ: New Zealand, PH: Philippines, UK: United Kingdom, CA: Canada.
194

195 **2.1.3 Environmental variables**

196 In the selection of environmental variables, our primary focus was on processes
197 within the terrestrial carbon cycle. The carbon cycle on land can be conceptualized as
198 two flux exchange processes influenced by the climatic conditions (Fig. 4). The CO₂ in
199 the atmosphere is fixed by vegetation photosynthesis and the carbon is released back
200 into the atmosphere by respiration and disturbance processes (Beer et al., 2010; Pan et
201 al., 2011). The carbon fluxes through these processes we considered as the land flux.
202 Since Industrial Era, anthropogenic carbon from land use change (e.g., deforestation)
203 and fossil fuels and cement become important parts of atmospheric CO₂ (Friedlingstein
204 et al., 2010), which we considered as the anthropogenic flux. Meanwhile, the two
205 processes are directly or indirectly driven by the climatic features (Sitch et al., 2015;
206 Chen et al., 2021). Consequently, we explored the potential drivers of XCO₂ from the
207 perspective of the carbon cycle at atmosphere-land interface. Multiple satellite products
208 and reanalysis data from three aspects (i.e., land flux, anthropogenic flux and climatic
209 impacts) were selected to consider their various effects on the XCO₂.



210

211 **Figure 4.** Simplified illustration of the global carbon cycle on land (referring to IPCC
212 2023). Noting that the carbon cycle in the ocean was not considered in our study and
213 we only focused on the fast exchange fluxes. The slow carbon exchanges (e.g., chemical
214 weathering, volcanic emissions) which are generally assumed as relatively constant
215 over the last few centuries (Sundquist, 1986), were not included here.

216
217 The key factors selected related to the land flux included gross primary
218 productivity (GPP), enhanced vegetation index (EVI), land surface temperature (LST),
219 vegetation continuous fields (VCF), and normalized difference snow index (NDSI).
220 These products are all obtained from the Moderate Resolution Imaging
221 Spectroradiometer (MODIS), which has been operated for over 20 years and produced
222 various satellite products with fine spatial resolution and accuracy. The EVI and NDSI
223 were converted to monthly data using the maximum value composite (MVC) method.
224 The GPP and LST were converted to monthly data by the averaging method.

225 The rising anthropogenic activities have greatly influenced atmospheric CO₂
226 (Friedlingstein et al., 2022). In this study, five anthropogenic factors, including land
227 use/cover change (LUCC), nighttime lights (NTL), and three trace gases (i.e., sulfur
228 dioxide (SO₂), nitrogen dioxide (NO₂), and carbon monoxide (CO)) were selected. The
229 LUCC was obtained from MODIS MCD12Q1 with a spatial resolution of 500 m. The
230 monthly Suomi National Polar-orbiting Partnership-Visible Infrared Imaging



231 Radiometer Suite (NPP-VIIRS) day/night band (DNB) NTL products (spatial
 232 resolution of 15 arc-second, ~500 m) were obtained from the Earth Observation Group
 233 (EOG) of the Colorado School of Mines. We also used the SO₂, NO₂ and CO products
 234 from the TROPOspheric Monitoring Instrument (TROPOMI) onboard Sentinel-5
 235 Precursor (S5P), a global air monitoring satellite for the Copernicus mission. The data
 236 were also converted to the same temporal resolution (i.e., monthly).

237 The selected climatic factors affecting XCO₂ were surface pressure (SP), 10 m
 238 wind speed (WS), precipitation flux (PRE), 2 m air temperature (Temp), and total
 239 evaporation (E). These data are from the reanalysis products (Hersbach et al., 2020)
 240 developed at the European Center for Medium Weather Forecasting (ECMWF,
 241 <https://www.ecmwf.int/>). The WS is calculated using the products of 10 m wind
 242 components of U and V. All data were converted to monthly time-series. The bilinear
 243 interpolation approach was used to convert the data at different spatial resolutions to 1
 244 km resolution. The data preprocessing was conducted on GEE, R and ArcGIS 10.3.
 245 Details of these products are listed in Table 2.

246 **Table 2.** Ancillary variables selected in this study.

Variables	Spatial resolution	Temporal resolution	Product names	Category
GPP	500 m	8-day	MOD17A2H	Land flux-related
EVI	1 km	16-day	MOD13A2	
LST	1 km	daily	MOD11A1	
VCF	250 m	annual	MOD44B	
NDSI	500 m	daily	MOD10A1	
LUCC	500 m	annual	MCD12Q1	
NTL	15 arc-second	monthly	VNP46A2	Anthropogenic flux-related
SO ₂			OFFL/L3_SO ₂	
NO ₂	~1 km	daily	OFFL/L3_NO ₂	
CO			OFFL/L3_CO	
SP				Climatic impacts
E				
Wind-v	~10 km	monthly	ERA5-Land	
Wind-u				
Pre				
Temp				



247 2.2 Deep learning-based XCO₂ reconstruction

248 Given the complexity temporal dependence and nonlinear relationship between
249 XCO₂ and the environmental variables, we selected the At-BiLSTM model to relate the
250 XCO₂ data with the 16 response variables affecting atmospheric CO₂, and further
251 reconstruct the XCO₂ data at a fine spatial resolution. The equation to reconstruct XCO₂
252 data in this research can be denoted as:

$$XCO_{2(i)} = f_{bilstm}([LF_{i,j}], [AF_{i,j}], [CI_{i,j}]) \quad (1)$$

$$= f_{bilstm}([GPP_t, EVI_t, LST_t, VCF_t, NDSI_t], [LUCC_t, NTL_t, SO_{2t}, NO_{2t}, CO_t], [SP_t, E_t, WU_t, WV_t, PRE_t, TEMP_t]) \quad (2)$$

253 where $XCO_{2(i)}$ denotes the aggregated monthly atmospheric CO₂ concentration at grid
254 cell i , $[LF_{i,j}]$, $[AF_{i,j}]$, and $[CI_{i,j}]$ denote the variables from land flux, anthropogenic flux,
255 and climatic impacts, respectively. And j denotes the number of variables in each
256 category. The f_{bilstm} represents the estimation model of each grid cell at the spatial
257 resolution of 0.05°.

258 The LSTM model is a variant of RNN that excels in processing time-series data
259 (Hochreiter and Schmidhuber, 1997; Graves et al., 2005). It has been utilized
260 extensively for prediction of remote sensing data. Each LSTM cell includes an input
261 gate, a forget gate and an output gate. The forget gate f_t determines which information
262 from the previous time step to forget, the input gate i_t governs the selective storage of
263 the data in current time step, and the output from forget gate f_t and input gate i_t are
264 combined in the cell state C_t . Lastly, the output gate o_t controls the flow of
265 information from the cell state to the next time step. These gate structures effectively
266 manage the flow of information within the LSTM, enabling it to capture the temporal
267 dependencies present in the data (Yuan et al., 2020; Su et al., 2021). Bidirectional
268 LSTM consists of two directional LSTM, in which the data flows forward and
269 backward. Then we defined a multi-dimensional attention layer behind the BiLSTM to
270 focus on more critical variables and give them higher weights (Bahdanau et al., 2016).
271 In the attention layer, we adopted the full connection layer and softmax activation
272 function to calculate the attention weight of each time step.

273 The At-BiLSTM consists of one input layer, three Bidirectional LSTM layers, one
274 attention layer, one dropout layer to prevent overfitting, and one fully connected layer
275 (i.e., dense layer) for the final output. The mean square error is used as the loss function.
276 The number of network units, batch size, learning rate and activation function



277 hyperparameters were all tuned in model fitting. All data were normalized using the
278 mean and standard deviation of the dataset. The model was built using the deep learning
279 API Keras in Python.

280 In this study, we adopted the sample-based cross-validation (CV) method to
281 evaluate the model performance and the in-situ validation to assess the accuracy of
282 reconstructed XCO₂ products. We also compared the reconstructed XCO₂ products with
283 the original OCO XCO₂ products and the CAMS-EGG4 GHGs data. Four metrics,
284 including coefficient of determination (R^2), root mean squared error (RMSE), mean
285 absolute error (MAE) and mean bias, were calculated as follow, to assess the model
286 performance.

$$R^2 = 1 - \frac{\sum_{i=1}^n (y_i - f_i)^2}{\sum_{i=1}^n (y_i - \bar{y})^2} \quad (3)$$

$$RMSE = \sqrt{\frac{\sum_{i=1}^n (y_i - f_i)^2}{n}} \quad (4)$$

$$MAE = \frac{\sum_{i=1}^n |f_i - y_i|}{n} \quad (5)$$

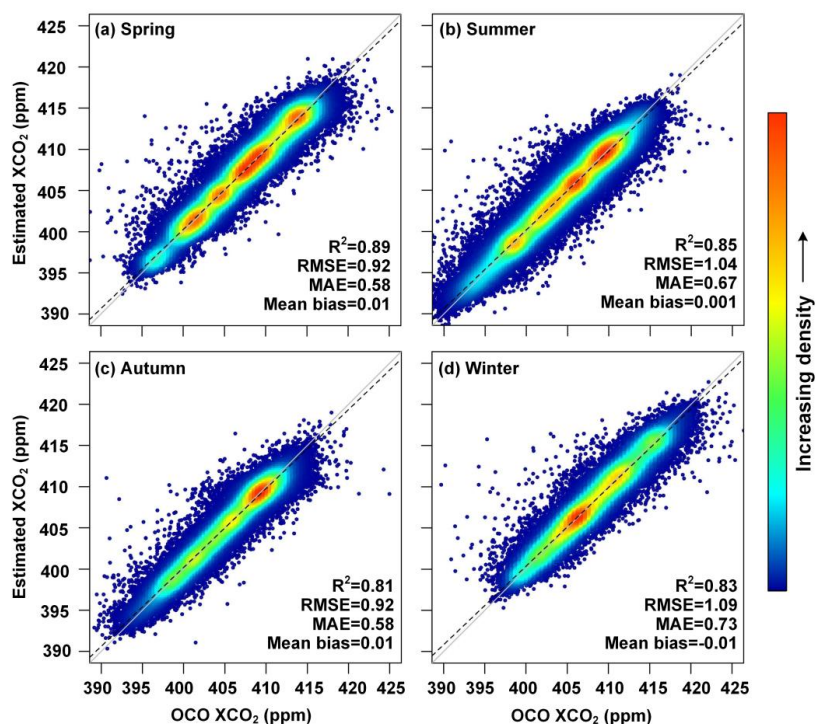
287 where n is the total number of data samples, and f_i, y_i are the observed results and
288 model-estimated results, respectively.

289 3. Results

290 3.1 Validation of the reconstructed XCO₂ product

291 3.1.1 Model validation results

292 Given the distinct seasonal variation in XCO₂ concentrations, we conducted the
293 sample-based CV to evaluate the model performance during different seasons (Fig. 5).
294 The model demonstrated high accuracy across all seasons, with R^2 values exceeding
295 0.81, MAE less than 0.73 ppm, and RMSE less than 1.09 ppm. The model performed
296 better in spring and summer, as indicated by the densest cluster of points being closest
297 to the 1:1 line. Conversely, the model performed worst in winter, when photosynthesis
298 is weakest, leading to greater estimation deviation. These variations are likely
299 influenced by the ecosystem CO₂ exchange during different seasons. Overall, the model
300 effectively captured the seasonal variation of XCO₂ and provided unbiased XCO₂
301 estimations.



302

303 **Figure 5.** (a) Density scatterplots of sample-based CV results during different seasons.
304 The proportion of the number of points is represented as the color of the points. The
305 black dashed lines and grey solid lines denote the linear regression fitted lines and the
306 1:1 line, respectively. The R^2 , RMSE (ppm), MAE (ppm), and mean bias (ppm) are
307 provided.

308 We further validated the model performance across different continents. Table 3
309 presents the validation results for six continents. The model performance varied across
310 continents. Notably, the model achieved the highest accuracy in Africa and Europe,
311 with R^2 of 0.80 and 0.81, and RMSE values of 1.02 and 1.14 ppm, respectively. In
312 contrast, the model demonstrated relatively low accuracy in Oceania and South
313 America, both located in the southern hemisphere. Despite this, the RMSE of the model
314 in these continents were 1.21 and 0.66 ppm, respectively, indicating that the model
315 maintained acceptable estimation accuracy in these regions.

316



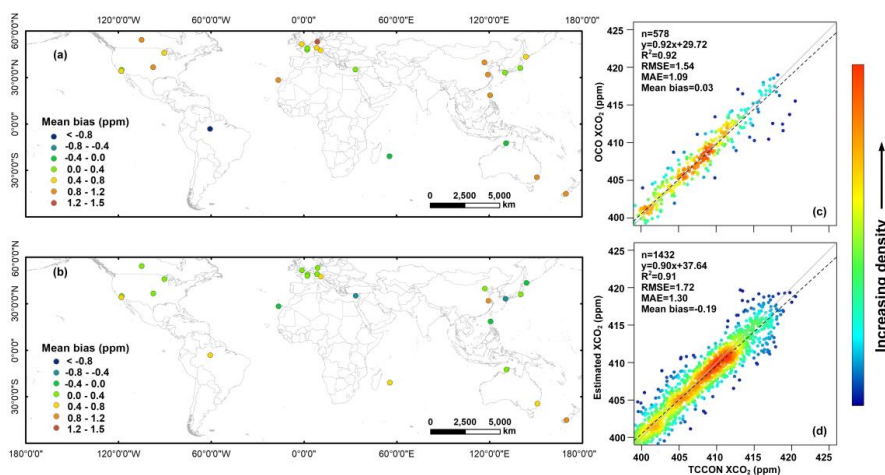
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Table 3. Model performance in different continents.

	R ²	RMSE (ppm)	MAE (ppm)	Mean bias (ppm)
Africa	0.80	1.02	0.70	-0.009
Asia	0.73	1.27	0.85	0.002
Europe	0.81	1.14	0.77	-0.030
North America	0.73	1.26	0.83	-0.020
South America	0.59	1.22	0.86	-0.012
Oceania	0.67	0.66	0.4	0.051

318 **3.1.2 In situ validation results**

319 The TCCON in situ XCO₂ data were adopted for validating the accuracy of the
320 reconstructed XCO₂ over the globe. The validation results for our reconstructed XCO₂
321 and the origin OCO-2/3 XCO₂ are displayed in Fig. 6. The two XCO₂ data showed
322 similar precision with the R² value of 0.91 and 0.92, respectively (Fig. 6c-d). While the
323 reconstructed XCO₂ greatly increases the data coverage with the validation sample
324 increasing from 578 to 1432. Meanwhile, the reconstructed XCO₂ has a smaller RMSE
325 and MAE with values of 1.72 and 1.3 ppm, respectively, compared with the OCO XCO₂.
326 These results indicate that the reconstructed XCO₂ had a closer agreement with TCCON
327 XCO₂. We also displayed the mean bias of OCO and reconstructed XCO₂ in each
328 TCCON site (Fig. 6a-b). As shown in Fig. 6a, the OCO-2/3 observation tend to
329 overestimate the XCO₂, while the reconstructed XCO₂ could amend the underestimation
330 of OCO XCO₂. Over 68% of the validation sites of reconstructed XCO₂ had a mean
331 bias less between ± 0.4 ppm. Given the orbital constraints of the ISS (Eldering et al.,
332 2019), OCO-3 measurements were restricted to latitudes below ± 52°. Consequently,
333 substantial missing values of OCO XCO₂ data were shown around 50°N, introducing a
334 potential bias. In contrast, the reconstructed XCO₂ effectively solves this problem and
335 demonstrates markedly enhanced performance.



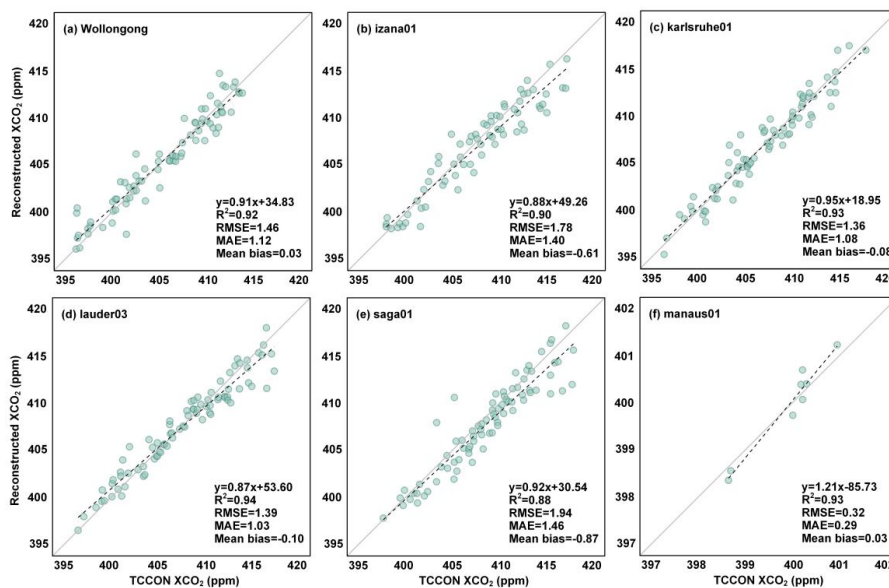
336

337 **Figure 6.** The mean bias of the (a) OCO observed XCO₂, and (b) reconstructed XCO₂
338 against global TCCON XCO₂; (c) density scatterplots of the validation results for OCO
339 observed XCO₂, and (d) reconstructed XCO₂ against the TCCON XCO₂. The
340 proportion of the number of points is represented as the color of the points. The number
341 of samples n , linear regression relation, R^2 , RMSE (ppm), MAE (ppm), and mean bias
342 are provided.

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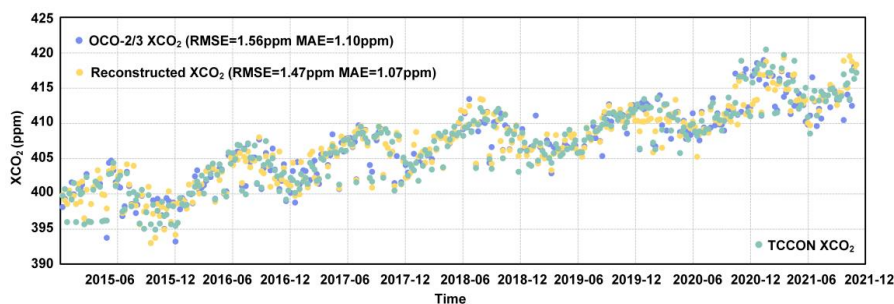
345 Fig. 7 shows the individual in situ validation results of the reconstructed XCO₂
346 against TCCON site in different continents (except Antarctica). The sample numbers
347 are varying in different sites due to the observation constraints, while the validation
348 results from all sites showed satisfying performance. The R^2 for all sites are over 0.88
349 and the MAE are less than 1.46 ppm. The reconstructed XCO₂ data performs the best
350 in sites lauder03 and karlsruhe01, which located in North America and Europe,
351 respectively. While the reconstructed XCO₂ performed worst in saga01 which located
352 in Asia, potentially due to the high CO₂ concentrations in these regions. Overall, the
353 reconstructed XCO₂ showed high consistency with the in situ XCO₂ observation in



354

355 **Figure 7.** Scatterplots of the TCCON in situ validation results of the reconstructed
 356 XCO₂ on different TCCON sites over the globe.
 357

358 To assess the performance of our reconstructed XCO₂ in temporal analysis, we
 359 compared the time series for monthly OCO-2/3, reconstructed and TCCON XCO₂ data
 360 from December 2014 to December 2021. As depicted in Fig. 8, the reconstructed XCO₂
 361 exhibits similar temporal patterns compared to the TCCON data, with the mean RMSE
 362 and MAE of 1.47 and 1.07 ppm. While the OCO-2/3 XCO₂ exhibits some
 363 overestimation for high values and underestimation for low values compared with
 364 TCCON data. In contrast, the reconstructed XCO₂ provided more stable estimate results.



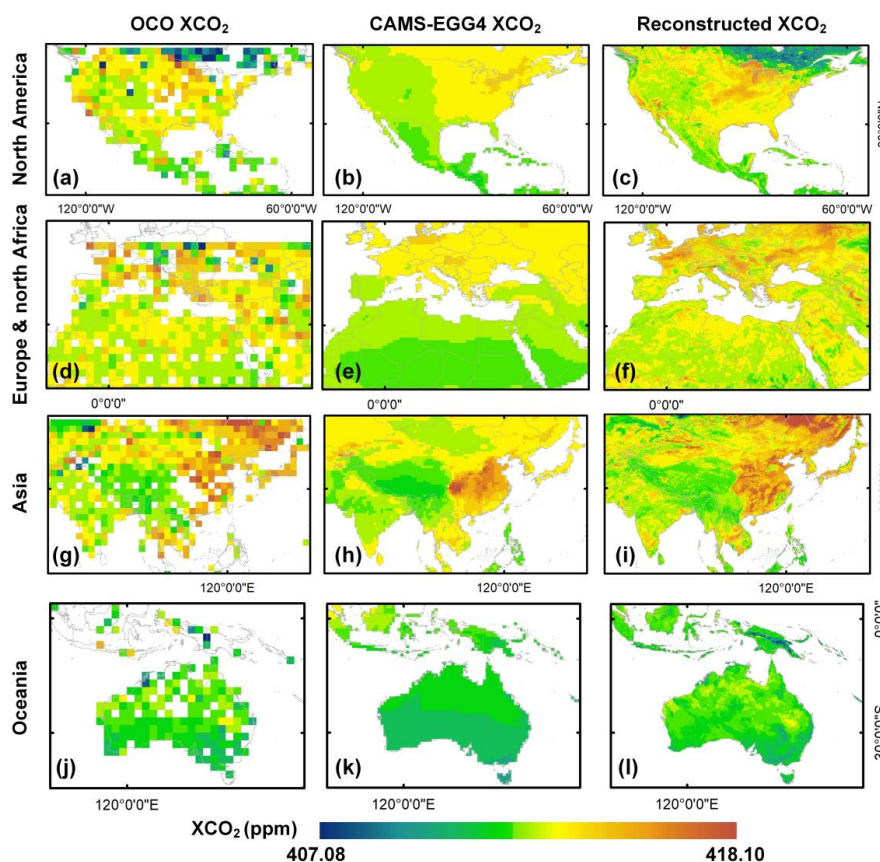
365

366 **Figure 8.** Comparison of the temporal variation of XCO₂ data from OCO-2/3 (blue
 367 dots), TCCON (green dots), and the reconstructed products (yellow dots).
 368



369 3.1.3 Product assessment with previous XCO₂ products

370 Fig. 9 shows the comparison between different XCO₂ products (i.e., original OCO
371 XCO₂, CAMS-EGG4 GHGs data, and our reconstructed XCO₂) in four specific regions.
372 The original OCO XCO₂ data were aggregated into 2°×2° latitude/longitude bins
373 following Taylor et al. (2020). The CAMS-EGG4 GHGs data are a global reanalysis
374 dataset at a spatial resolution of 0.75° released by the European Centre for Medium-
375 range Weather Forecasts (Agustí-Panareda et al., 2023). The North America, Europe
376 and part of north Africa, Asia and Oceania were chosen as examples. As shown in Fig.
377 9, the spatial coverage of CAMS-EGG4 GHGs data and our reconstructed XCO₂ is
378 significantly increased compared to the original OCO XCO₂ data. However, the
379 CAMS-EGG4 GHGs data is at a coarse spatial resolution and miss much of the detailed
380 information on XCO₂ change. In comparison, our reconstructed seamless XCO₂
381 product can provide much more information on the global and regional spatial patterns
382 of XCO₂. Due to the limited coverage, the original OCO XCO₂ data failed to capture
383 the variation of XCO₂ in the Midwestern United States, northern part of the United
384 Kingdom, and central China. In contrast, these regions are well represented in the
385 reconstructed XCO₂ product. Furthermore, the reconstructed XCO₂ with a fine spatial
386 resolution can offer a more accurate spatial distribution of carbon sources and sinks.
387 For example, lower XCO₂ concentrations are clearly observed in the forests in eastern
388 Canada and Papua New Guinea (Fig. c and l), indicating the great potential carbon sink
389 of these areas. While the CAMS-EGG4 GHGs data cannot capture this change due to
390 their coarse spatial resolutions. In general, reconstructed XCO₂ products with complete
391 coverage and finer resolution provide valuable support for analyzing atmospheric CO₂
392 variation and accurate monitoring of carbon sources and sinks.



393

394 **Figure 9.** Comparison between the OCO observed XCO₂ data aggregated into 2°×2°
 395 latitude/longitude bins, the CAMS-EGG4 GHGs data, and our reconstructed XCO₂ data
 396 in four regions, using the products of December of 2020 as an example.

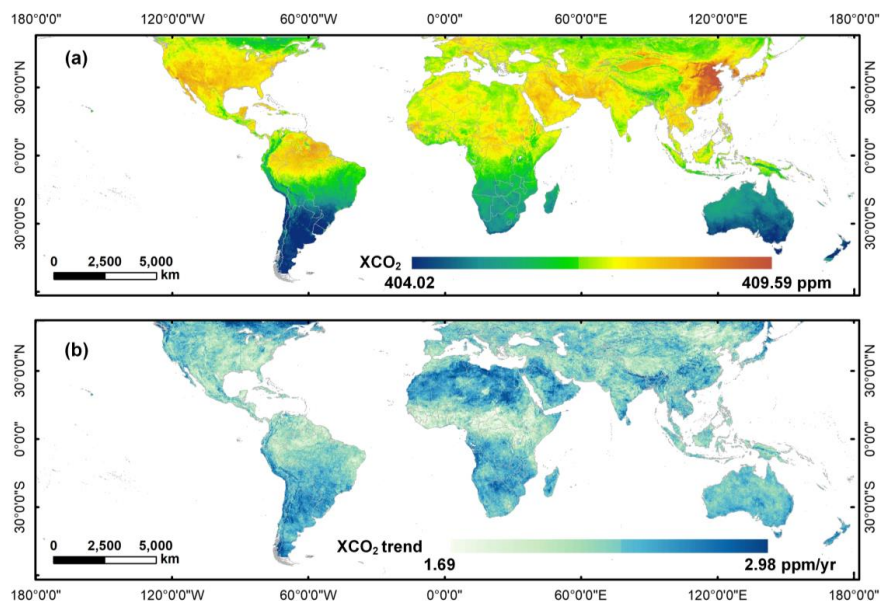
397 3.2 Spatiotemporal pattern of global XCO₂

398 The average XCO₂ concentration from 2015 to 2021 was 406.90 ± 0.80 ppm across
 399 the globe. The atmospheric CO₂ exhibited an apparent spatial variation, with higher
 400 concentrations typically observed in the Northern Hemisphere and lower
 401 concentrations in the Southern Hemisphere (Fig. 10a). The highest concentration of
 402 XCO₂ mainly occurs in the northern low-to-mid-latitudes (10°N-45°N), such as East
 403 Asia, southern Northern America, and the Middle East. More frequent human activities
 404 and carbon emissions contributed to higher atmospheric CO₂ concentrations in the
 405 Northern Hemisphere. In contrast, the lowest XCO₂ concentration was 404.02 ppm,
 406 occurring in the Southern Hemisphere where 81% of the area is ocean. The oceans act
 407 as a vital carbon sink and absorb most atmospheric CO₂. For the continent scale, the



408 XCO₂ concentrations showed a slight variation (± 1 ppm) between different continents.
409 The largest XCO₂ were mainly occurred in Asia and North America over years, while
410 the lowest XCO₂ concentration all presented in Oceania (Table 4).

411 Fig. 10b presents the spatial distribution of the 7-year (2015-2021) XCO₂ trend
412 over the globe. In terms of temporal trend, the atmospheric CO₂ exhibited a distinct
413 increasing trend over time, with the mean growth rate of 2.32 ppm yr⁻¹. The large
414 growth rate meanly occurs in the northern low latitudes (0°N-30°N), especially the
415 Middle East and North Africa (growth rate over 2.5 ppm yr⁻¹). Globally, the XCO₂
416 increased by 14.16 ppm over seven years (Table 4), especially in 2021, with increased
417 values of up to 3 ppm. This result is consistent with the Global Carbon Budget 2022
418 (Friedlingstein et al., 2022), which reported that the global average atmospheric CO₂
419 increased sharply in 2021 and reached 414.71 ppm.



420
421 **Figure 10.** The global spatial distribution of (a) reconstructed annual mean XCO₂
422 concentration, and (b) its trend from 2015 to 2021 (ppm yr⁻¹ denotes parts per million
423 per year).
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Table 4. The reconstructed XCO₂ concentrations at different continents from 2015 to 2021.

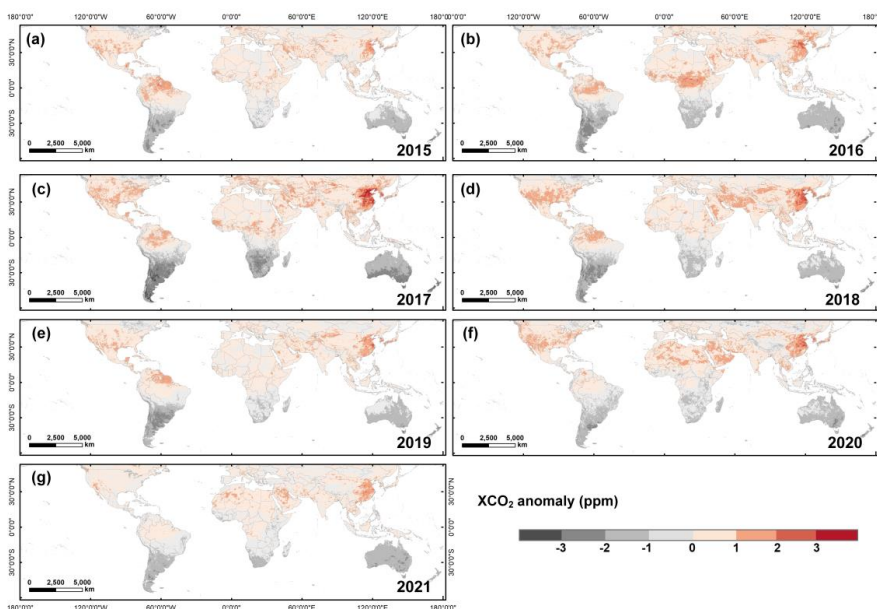
Continents	XCO ₂ concentrations (ppm)							
	2015	2016	2017	2018	2019	2020	2021	Increase
Africa	399.26	402.66	404.98	406.71	409.26	411.13	414.11	14.85
Asia	399.57	403.03	405.80	407.37	409.68	411.39	414.38	14.81
Europe	399.55	402.88	405.77	406.96	409.48	411.30	414.17	14.62
North America	399.60	402.95	405.76	407.32	409.70	411.61	414.28	14.68
South America	398.94	401.96	404.27	406.17	408.78	410.47	413.57	14.63
Oceania	398.03	401.04	403.31	405.53	408.13	409.82	412.55	14.52
Global	399.84	401.56	405.16	407.50	409.21	411.07	414.00	14.16

434

435 3.3 The distribution of XCO₂ anomaly

436 To better explore the dynamics of global carbon change, we further calculated the
437 XCO₂ anomalies based on the full-coverage XCO₂ products and presented their global
438 distributions from 2015 to 2021 (Fig. 11). The XCO₂ anomalies were calculated by the
439 statistical filtering method, that is, subtracting the global median XCO₂ value from the
440 global XCO₂ distribution (Hakkarainen et al., 2016). The spatial pattern of XCO₂
441 anomalies were relatively consistent over seven years with no significant variations.
442 From the global perspective, high XCO₂ anomalies mainly occurred in the Northern
443 Hemisphere. East Asia has the largest XCO₂ anomalies with values ranging from 2 to
444 3 ppm, such as the east part of China. The Middle East, North Africa and the southern
445 part of Northern America also experienced high XCO₂ anomalies. Nevertheless,
446 negative XCO₂ anomalies were also identified in the Northern Hemisphere, specifically
447 in regions such as Tibet in China, eastern Canada, and southern Russia. Most negative
448 XCO₂ anomalies were observed in the Southern Hemisphere, which behaves as a
449 carbon sink. However, some positive XCO₂ anomalies are also observed in the tropical
450 regions (e.g., Amazonia), which indicates the Amazonia has changed into a carbon
451 source due to the deforestation and fire occurrence in recent years (Hubau et al., 2020;
452 Gatti et al., 2021).

453



454
455 **Figure 11.** The global spatial distribution of annual XCO₂ anomaly from 2015 to 2021.

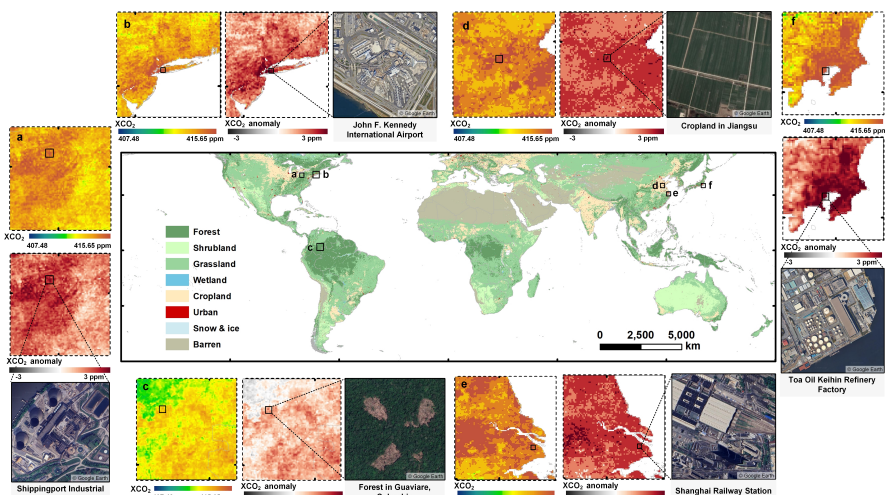
456

457 Fig. 12 illustrates the detailed spatial distribution of XCO₂ concentrations and
458 anomalies over six regions with high XCO₂ retrievals in 2020. High concentrations of
459 XCO₂ were typically associated with energy-intensive heavy industrial activities, such
460 as Toa Oil Keihin Refinery Factory located in Kawasaki City, Japan (Fig. 12f), and the
461 Shippingport Industrial Park in Pennsylvania, United States (Fig. 12a). Moreover,
462 certain metropolitan transport hubs also exhibited elevated CO₂ anomalies attributable
463 to dense populations and intensive activities. Examples included Shanghai Station in
464 China (Fig. 12e) and John F. Kennedy International Airport in New York, USA (Fig.
465 12b). Attention has also been drawn to natural sources of emissions. Driven by the
466 significant impact of agricultural mechanization and agro-industrial activities on
467 cropland (Lin and Xu, 2018), the XCO₂ anomalies also occurred in the agricultural
468 areas northwestern Jiangsu, China (Fig. 12d). Additionally, we also observed the high
469 XCO₂ anomalies in Amazonia forest in Colombia, which have been suffered from
470 deforestation (Gatti et al., 2023). In conclusion, our products could successfully capture
471 the XCO₂ anomalies from different sources over the globe.

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475
476 **Figure 12.** Examples of XCO₂ hotspots in six regions for 2020 detected using the
477 reconstructed products. The subplots present the spatial distribution of XCO₂
478 concentrations, anomalies (the red panels), and the emission sources (the true color
479 images from © Google Earth), respectively. The global map in the middle presents the
480 land use and land cover types over the globe.

481 4. Discussion

482 In this study, we utilized deep learning and multi-source satellite data (i.e.,
483 OCO2/3, MODIS, VIIRS and TROPOMI) to reconstruct global XCO₂ products at a
484 spatial resolution of $0.05^\circ \times 0.05^\circ$. Though our products achieved full spatial coverage
485 and high accuracy, there are still several limitations need further improvement. In terms
486 of sensors, OCO-2 and OCO-3 data provide different spatiotemporal coverages.
487 Analyzing OCO-2 and OCO-3 data simultaneously may introduce several uncertainties
488 due to these differences. However, OCO-3 has a similar sensor and inherits the retrieval
489 algorithms of OCO-2. According to Taylor et al. (2023), the mean differences between
490 OCO-3 and OCO-2 are around 0.2 ppm over land. Therefore, we suppose that the
491 discrepancies between their datasets are minimal, and the combined analysis of data
492 from these two satellites will have a negligible impact on our results.

493 Moreover, despite OCO is considered to be one of the most accurate carbon
494 satellite datasets to date, it still encounters some retrieval errors due to the influence of
495 retrieval methods and meteorology conditions, which may be introduced by using the
496 data as a target for XCO₂ reconstruction. However, the validation results against
497 TCCON suggested that the RMSEs of the two OCO XCO₂ datasets are both less than



498 1 ppm, which is sufficient for monitoring changes in atmospheric CO₂ (Taylor et al.,
499 2023).

500 Additionally, prediction uncertainty may also arise from the model and covariates
501 used (Chen et al., 2022). While our deep learning model achieved high accuracy in
502 general, its performance in the Southern Hemisphere could be further improved. This
503 is attributed to the distribution of data features and the complex non-linear relationship
504 between XCO₂ and the environmental covariates. Selecting more relevant
505 environmental factors may help mitigate these issues and enhance model performance
506 in this region.

507 5. Data availability

508 The XCO₂ dataset produced in this paper is available at
509 <https://doi.org/10.5281/zenodo.12706142> (Wang et al., 2024). It includes monthly
510 global XCO₂ data at 0.05° resolution, covering the period from December 2014 to
511 December 2021. The dataset is archived in netCDF4 format, with units in parts per
512 million (ppm).

513 6. Conclusion

514 The launch of carbon satellites offers a significant advancement for CO₂
515 monitoring. However, their limited spatial coverage restricts the effectiveness of XCO₂
516 data. To address this issue, we reconstructed a global full-coverage XCO₂ product at a
517 fine spatial resolution using multi-component satellite data. The advanced deep learning
518 method was adopted to model time-series XCO₂ and incorporate terrestrial flux,
519 anthropogenic flux and climatic impacts into the parameterization process. Our
520 reconstructed XCO₂ products showed a strong agreement with TCCON XCO₂, with R²,
521 RMSE, and MAE values of 0.92, 1.54 ppm, and 1.09 ppm, respectively. The products
522 provided accurate information on the global and regional spatial pattern of XCO₂. The
523 global XCO₂ exhibited a distinct increasing trend over time, reaching 414.00 ppm in
524 2021. Higher XCO₂ concentrations were primarily observed in the northern low-to-
525 mid-latitudes (10°N-45°N) such as Asia and North America. Utilizing the reconstructed
526 products, we further detected XCO₂ anomalies globally and identified intensive carbon
527 emission sources across different land use types. Our study presents a viable method
528 for global-scale, high-resolution XCO₂ mapping based on carbon satellites and



529 demonstrates the feasibility of applying this methodology to explore global and
530 regional carbon dynamics.

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538 data, the Copernicus Climate Data Store for providing the ERA5 reanalysis data and
539 CAMS-EGG4 XCO₂ data, and the Total Carbon Column Observing Network for their
540 dedication in providing the in situ XCO₂ observations.

541 **Author contributions**

542 ZW developed the overall workflow, processed the data and wrote the manuscript. CZ
543 and BH revised the manuscript. KS, YS, and XC compiled the data. SC conceptualized
544 and revised the manuscript. PA and QZ supervised this study. All the authors
545 contributed to the study.

546 **Competing interests**

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

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