Seasonal responses of $\delta 13C$ and $\delta 18O$ of atmospheric CO2 over sub-urban region of India

Mahesh Pathakoti¹, Kanchana Asuri¹, Mahalakshmi D¹, Sreenivas Gaddamidi², TANIA GUHA³, Vijay Sagar¹, Raja P⁴, and Sesha Sai MVR¹

¹NRSC ²JNTU, Hyderabad ³IITM ⁴IISWC, ICAR

November 24, 2022

Abstract

Seasonal and diurnal variability of atmospheric CO2 and its driving factors are studied using its continuous monitoring of concentration and isotopic ratios (δ 13C-CO2 and δ 18O-CO2) for the first time over Shadnagar, a sub-urban location of India with high precision in-situ data from November 2018 to October 2019. The annual averaged atmospheric CO2 concentrations, δ 13C-CO2 and δ 18O-CO2 are 414.76±4.26 ppm, -11.19±1.63 atmospheric CO2 was observed in summer monsoon (17.30 +-9.29 ppm) and the minimum was noticed in winter (7.19 +- 0.11 ppm) indicating strong seasonality at the study site. To characterize the atmospheric CO2 sources, an improved model of Miller and Tans was implemented by plotting DCO2 against D(CO2 x δ 13C) respectively during day and night. However, a strong source/sink signature of δ 13C was observed during nighttime of summer monsoon with a slope of -37.42 ± 0.73 using a reduced major axis (RMA) regression. The source identified is attributed to combustion and dominance of C3 ecosystem respiration respectively. The seasonal relationship between δ 18O-CO2 and δ 13C-CO2 is strongly correlated during pre-monsoon ('r' = 0.93-0.95) than post-monsoon ('r' = 0.07-0.13), which might be due to high vapour pressure deficit. A Lagrangian back-trajectory model confirms the influence of the Indian summer monsoon on the variability of atmospheric CO2 concentration during the summer monsoon season.

1	
2	Seasonal responses of δ^{13} C and δ^{18} O of atmospheric CO ₂ over sub-urban region of
3	India
4	
5	Mahesh Pathakoti ^{1*} , A.L. Kanchana ¹ , D.V. Mahalakshmi ¹ , Sreenivas G ² , Tania Guha ^{3,4} ,
6	Vijay Kumar Sagar ¹ , P.Raja ⁵ , and SeshaSai M.V.R ¹
7	
8	¹ Earth and Climate Sciences Area (ECSA), National Remote Sensing Centre (NRSC), Indian Space
9	Research Organisation (ISRO), Hyderabad-500037, India.
10	2
11	² Department of Physics, Jawaharlal Nehru Technological University, Hyderabad, 500085, India
12	2
13	³ Department of Geography, University of Calgary, Calgary, Alberta, Canada
14	4
15	⁴ Previously at Indian Institute of Tropical Meteorology, Pune, India
16	
17	⁵ ICAR-Indian Institute of Soil and Water Conservation, Research Centre, Ooty, The Nilgiris, Tamil
18	Nadu, - 643004,India.
19	
20	
21 22	*Corresponding author: Mahesh Pathakoti (mahi952@gmail.com)

- 23 Key Points:
- The Study focused on diurnal variability of seasonal averaged atmospheric CO₂ concentration and its stable isotopes (δ^{13} C-CO₂ and δ^{18} O-CO₂).
- Determination of δ^{13} C signature of atmospheric CO₂ using improved Miller and Tans model.

• The effect of the Indian summer monsoon circulation on atmospheric CO₂ variation was studied.

30 Abstract

Seasonal and diurnal variability of atmospheric CO₂ and its driving factors are studied using its 31 continuous monitoring of concentration and isotopic ratios (δ^{13} C-CO₂ and δ^{18} O-CO₂) for the first 32 time over Shadnagar, a sub-urban location of India with high precision in-situ data from 33 November 2018 to October 2019. The annual averaged atmospheric CO₂ concentrations, δ^{13} C-34 CO₂ and δ^{18} O-CO₂ are 414.76±4.26 ppm, -11.19±1.63 ‰ and 9.02 ±12.78 ‰. Maximum 35 seasonal diurnal variability of atmospheric CO₂ was observed in summer monsoon (17.30 ± 9.29 36 ppm) and the minimum was noticed in winter $(7.19 \pm 0.11 \text{ ppm})$ indicating strong seasonality at 37 the study site. To characterize the atmospheric CO₂ sources, an improved model of Miller and 38 Tans was implemented by plotting ΔCO_2 against $\Delta (CO_2 \times \delta^{13}C)$ respectively during day and 39 night. However, a strong source/sink signature of δ^{13} C was observed during nighttime of summer 40 monsoon with a slope of -37.42 ± 0.73 %, obtained using a reduced major axis (RMA) 41

- regression. The source identified is attributed to combustion and dominance of C₃ ecosystem respiration respectively. The seasonal relationship between δ^{18} O-CO₂ and δ^{13} C-CO₂ is strongly
- respiration respectively. The seasonal relationship between δ^{18} O-CO₂ and δ^{13} C-CO₂ is strongly correlated during pre-monsoon ('r' = 0.93-0.95) than post-monsoon ('r' = 0.07-0.13), which
- 45 might be due to high vapour pressure deficit. A Lagrangian back-trajectory model confirms the
- influence of the Indian summer monsoon on the variability of atmospheric CO_2 concentration
- 47 during the summer monsoon season.
- 48 Keywords: atmospheric CO₂, reduced major axis,Miller and Tans, Indian summer monsoon.

49 **1. Introduction**

The Intergovernmental Panel for Climate Change (IPCC) reported that carbon dioxide (CO₂) 50 sources from anthropogenic gases in the atmosphere cause more radiative forcing next to water 51 vapor (Smith et al., 1999). CO₂ concentrations are consistently increasing and touched 400 ppm 52 at Mauna Loa, a global reference site during May 2013 (Monastersky, 2013). Globally, CO, 53 54 concentrations are increasing, which could be due to land use land cover changes (LULCC) and 55 progress in industrial activities (Ballantyne et al., 2012) especially fossil fuel combustion, cement manufacturing etc.. Due to fossil fuels burnings and LULC emissions, an increase of 40 56 and 150 % in CO₂ and CH₄ concentration respectively is observed since the pre-industrial period 57 (Huang et al., 2015). Emissions of CO₂ by different processes are controlled by varied 58 environmental conditions, in which about half of the CO₂ levels are released into the atmosphere 59 as source and remaining are absorbed by the processes of the terrestrial biosphere and ocean 60 uptake (Andres et al., 1996) as sink. Hence monitoring and maintaining long-term records of 61 atmospheric CO₂ measurements are very important to understand the carbon cycle and to assess 62 the CO, mixing ratios in the atmosphere by controlling factors namely photosynthesis, 63 64 respiration, biomass, fossil fuel burning and air-sea exchange processes (Machida et al., 2003).

Globally, systematic high precision atmospheric CO₂ observations are accelerated to understand 65 the global carbon cycle. Over the Indian subcontinent, spatio-temporal variability in atmospheric 66 CO₂ concentrations are characterized by the terrestrial biosphere and seasonal weather patterns 67 68 which brings long-range air-masses (Valsala et al., 2013; Tiwari et al., 2014). To understand the seasonal, inter and intra annual variations of atmospheric CO₂ over the Indian subcontinent, high 69 precision CO₂ measurements are being generated across the country from different research 70 institutes (Bhattacharya et al., 2009; Sharma et al., 2014; Mahesh et al., 2016; Nalini et al., 71 2019). A literature survey on atmospheric CO₂ variability over the Indian region is mainly 72 focused on local sources and transport (Sreenivas et al., 2016). However, a need was felt to 73 74 understand the reasons and causes for uncertainty in surface fluxes. Hence an advanced studies of stable isotopic measurements of carbon and oxygen are gaining momentum to understand CO₂ 75 levels, source and sinks of CO₂, both on regional and global levels. 76

Stable carbon and oxygen isotopes of atmospheric CO₂ can be used as tracers in the carbon cycle, 77 which are affected by the anthropogenic and biogenic CO₂ components. The δ^{13} C-CO₂ and δ^{18} O-78 CO₂ are stable isotopes of CO₂ molecules are widely used for source apportionments in the 79 atmosphere, hydrosphere and geosphere as well as interaction between them (Guillon et al., 80 2015). The δ^{13} C-CO₂ concentration in the atmosphere has been decreasing since pre-industrial 81 times, which indicates the more addition of CO₂ to the atmosphere by fossil fuel burning (Yakri, 82 2011). Many research activities have been carried on atmospheric CO₂ and its stable isotopes 83 $(\delta^{13}C-CO_2 \text{ and } \delta^{18}O-CO_2)$ by various groups (Bhattacharya et al., 2009; Pataki et al., 2003; 84

Clark-Thorne and yapp, 2003; Francey and Tans, 1987; Murayama et al., 2010; Newman et al, 85 2003; Wada et al., 2003; Pataki et al., 2006; Zhou et al., 2005; Zhou et al., 2006; Sturm et al., 86 2006; Djuricin et al., 2010; Guha and Ghosh, 2010, 2013, 2015; Gorka and Lewicka-Szczebak, 87 2013; Liu et al., 2014; Pang et al., 2016). Liu et al. (2014) studied the atmospheric CO₂, δ^{13} C-88 CO₂ composition and their relationship to understand the sources and sinks at two stations viz., 89 Waliguan and Shangdianzi in China using observational data for the period from 2007 to 2010. 90 The result of this study indicates that CO₂ and δ^{13} C-CO₂ composition possesses long-term trends 91 and seasonal cycles that correlate with each other. An improved model by Miller and Tans 92 (2003) is widely used to determine the source or sink that causes CO_2 variability. Pang et al. 93 (2016) used Keeling plot intercept method for isotopic composition of CO₂ and found 94 increased value in vegetative season and depleted value in heating season. A few studies were 95 carried on atmospheric CO₂ and its stable isotopes in India, which are limited to discrete sample 96 analysis (Bhattacharya et al., 2009; Guha and Ghosh, 2013, 2015). The present study on 97 atmospheric CO₂ and its stable isotopes was carried using laser based Isotope ratio Infrared 98 spectrometer analyser. Measurements of the δ^{18} O-CO₂ also play an important role in the carbon 99 cycle to distinguish the photosynthesis and respiration process of CO₂ fluxes (Farquhar et al., 100 101 1993; Kato et al., 2004).

The objective of the present study is to understand diurnal variability of seasonal 102 atmospheric CO₂ and its stable isotopic composition at the sub-urban region, of Telangana, India. 103 104 Since the study site is surrounded by multiple sources for CO₂ (e.g., biospheric respiration and fossil fuel emissions), thus an improved model by Miller and Tans (2003) was used to 105 characterize the CO₂ sources/sinks. Here, we report continuous high precision CO₂ isotopic 106 measurements, first of their kind from sub-urban region, of Telangana. This work has been 107 carried out as part of the Atmospheric CO₂ Retrieval and Monitoring (ACRM) of the National 108 Carbon Project (NCP) funded by the Climate and Atmospheric Processes of ISRO-Geosphere 109 Biosphere Programme (CAP-IGBP). 110

111 2. Materials and Methods

Observations of CO₂ and its isotopic composition are measured during November 2018 to 112 113 October 2019 by laser based Isotope Ratio Infrared Spectrometer (IRIS) analyser. This instrument is installed at 8 m height from the surface of ASL, NRSC, Shadnagar (Latitude: 17.09 114 °N; Longitude: 78.21 °E and Elevation: 648 m above mean sea level), a sub-urban region, 115 northern side of Hyderabad(~ 60 km away). Thus our study site is near to the highly populated 116 city of Hyderabad. 117 Surface meteorological data at the study site are collected from an automatic weather station. An 118 hourly Boundary Layer Height (BLH) were obtained from European Centre for MediumRange 119 (ECMWF-ERA, 120 Weather Forecasts https://cds.climate.copernicus.eu/cdsapp#!/dataset/reanalysis-era5-single-levels?tab=form). 121 In addition to the above datasets, fire counts with confidence interval of > 70% are considered from 122 Moderate Resolution Imaging Spectroradiometer(MODIS). Normalized Difference vegetation 123 Index (NDVI) is obtained from an open data archival of Bhuvan site (https://bhuvan-124 app3.nrsc.gov.in/data/download/index.php), which is derived from Oceansat-2 Ocean color 125 monitor sensor. Figure 1 shows that the study area is overlaid with NDVI as well as time series 126 of air temperature, relative humidity and wind speed recorded at the study site. 127 128

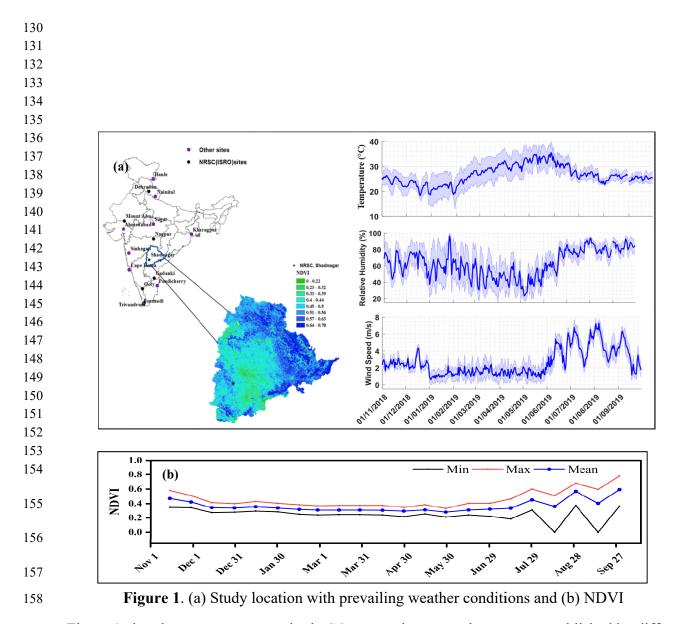


Figure 1 also shows *in-situ* atmospheric CO_2 networks across the country established by different 159 research institutes of India. A few of the station's information is obtained from Nalini et al. 160 (2019). Black solid circles are the atmospheric CO₂ measurement stations installed under ACRM 161 of NCP project by NRSC, ISRO. At the present study site (solid black star) long-term CO₂, CH₄ 162 and H₂O measurements are complemented by CO₂ and N₂O isotopic observations. During the 163 study period, air temperature is observed to be high in pre-monsoon (March-May) with 164 165 maximum air temperature of 43°C and low in winter (January-February) with mininum temperature of 9°C. Relative humidity is observed to be high in monsoon (June-September) with 166 maximum value reaching 100% and low in pre-monsoon with minimum of 12.18%. However, 167 wind speed is observed to be ranging from 0.005 m s^{-1} to 6.0 m s^{-1} during the study period. 168 Seasonal NDVI during the study period for post-monsoon, winter, pre-monsoon and monsoon 169 are 0.39, 0.33, 0.30 and 0.41 respectively. The study site is about 60 km away from the urban 170 region of Hyderabad (fifth largest city in India) associated with 75% air pollution from traffic 171 sector alone (significant anthropogenic impacts) due to increase in population and related factors 172

173 (Mahalakshmi et al., 2014). Thus the present study site doesn't fall under single source contributor of atmospheric CO_2 .

- 175
- 176
- 177

180

178

179 **2.1. Continuous stable isotopic measurements**

In this study, we used commercial laser based IRIS CO₂ Carbon Isotope analyzer Enhanced 181 Performance (CO₂-CCIA-EP), procured from Los Gatos Research, U.S.A in November 2018. 182 Details of the instrument functioning are given in Baer et al. (2002). The CO₂-CCIA-EP is 183 capable of simultaneous measurements of dry CO₂, ¹³C, ¹⁸O and H₂O are measured using the 184 absorption line at 2.05 µm with 1 Hz frequency and uses a performance enhancing off-axis 185 cavity ring down spectroscopy. Mole fraction of isotopic composition also depends on internal 186 cavity pressure and temperature hence maintained constant at 119.14 Torr and 45.36 °C 187 respectively (Mahesh et al., 2015). To keep the moisture as constant and as low as possible in the 188 analyzer, sample and reference gases are passed through a Neflon drying unit. The dry-air mole 189 fractions of CO₂ where the measured mole fraction of H₂O, which is also in ppm has been 190 removed as shown in Equation 1. 191

$$CO_{2}(ppm)_{dry} = \left(\frac{CO_{2} (ppm)_{wet}}{(1 - \frac{H_{2}O (ppm)}{10^{6}})}\right)$$
(1)

The ¹³C-CO₂ and ¹⁸O-CO₂ compositions are reported as δ^{13} C and δ^{18} O respectively versus VPDB (Vienna Pee Dee Belemnite) and reported in per mil (‰) as shown in Equation (2) and (3)

$$\delta^{13}C(\%_{0}) = \left(\frac{C_{s}}{C_{r}} - 1\right) \times 1000$$

$$\delta^{18}O(\%_{0}) = \left(\frac{O_{s}}{O_{r}} - 1\right) \times 1000$$
(2)
(3)

195 Where C_s, C_r, O_s , and O_r defined as follows; $C_{s (sample)} = ({}^{13}C/{}^{12}C)_s$; $C_{r (VPDB)} = ({}^{13}C/{}^{12}C)_r$; 196 $O_{s (sample)} = ({}^{18}O/{}^{16}O)_s$ and $O_{r (VPDB)} = ({}^{18}O/{}^{16}O)_r$

197

198 **2.2. Calibration of CO₂ isotope analyzer**

199

The CO₂-CCIA-EP is calibrated using National Oceanic and Atmospheric Administration 200 (NOAA) supplied CO₂ isotope calibration reference gases (ID: CC718409) towards reporting 201 data, eliminating instrument drifts and generating high quality data during the period of study. 202 The precision and accuracy of the instrument are computed by performing the internal 203 calibration(4 times) at frequent intervals (24th February 2019, 11th June 2019, 6th August 2019 204 and 8th January 2020). The resultant calibration showed a variation of 404.38±0.14 ppm of CO₂, 205 -8.63±0.04 % VPDB of δ^{13} C-CO₂ and -1.41±0.07 % VPDB of δ^{18} O-CO₂ respectively. An 206 inflow of reference gas has been passed through the Neflon drying unit for 5 minutes and 207 collected data at 1 Hz frequency. 208

- 209
- 210
- 211

213

214 The 100 sec (1 σ) average precision of CO₂, δ^{13} C-CO₂ and δ^{18} O-CO₂ are 0.20 ppm, 1.1 ‰ and

215 7.19 ‰ and their respective drift are 0.074 ppm, 0.085 ‰ and 0.485 ‰ respectively. The result

of the calibration report is summarized in table 1.

Cylinder ID	$CO_2(ppm, ref) \pm 1\sigma$	δ^{13} C of CO ₂ (‰, ref) ± 1 σ	δ^{18} O of CO ₂ (‰, ref) ± 1 σ	
	$CO_2 (ppm, M) \pm 1\sigma$	δ^{13} C of CO ₂ (‰, M) ± 1 σ	δ^{18} O of CO ₂ (‰, M) ± 1 σ	
NOAA,CC718409	404.53 ± 0.21 (ppm)	-8.45 ± 0.01 (‰, VPDB)	-1.28 ± 0.03 (‰, VPDB)	
	404.38 ± 0.14 (ppm)*	$-8.63 \pm 0.04 \ (\%, VPDB)^*$	$-1.41 \pm 0.07 \ (\%_0, \text{VPDB})^*$	
Bias (Ref-M)	0.15 (ppm)	0.18 (‰)	0.13 (‰)	
Precision	0.07%	0.54%	5.4%	
Accuracy	0.04%	3.20%	16.4 %	
Ref: Reference; M:	Measured	*Indicates means of 100sec of 5 min calibration for 4		
	12	times calibration		

217 **Table 1** Calibration report of CO₂, δ^{13} C-CO₂ and δ^{18} O-CO₂.

218

Bias between reference and measured values of CO₂, δ^{13} C-CO₂ and δ^{18} O-CO₂ are 0.15 ppm, 0.18 % and 0.13 % respectively. The precision and accuracy of CO₂, δ^{13} C-CO₂ and δ^{18} O-CO₂ are deduced with an averaging time of 100 sec. Results of the calibration in precision term are 0.07 %, 0.54% and 5.4% for CO₂, δ^{13} C-CO₂ and δ^{18} O-CO₂ respectively. The precision of δ^{18} O-CO₂ is

coarse compared δ^{13} C-CO₂, which may be improved by performing calibration for longer averaging time. However, one needs to be compromised for the precision averaging time (Guillon et al., 2015). The second level quality was applied to the raw data by adjusting the respective biases.

227

228 **2.3. Isotopic fraction using Improved model by Miller and Tans**

229 Present study implemented Thoning et al. (1989) to compute the curve fitting and smoothing of 230 231 the recorded time series data. To account, the strong atmospheric mixing of CO_2 during daytime, active convective boundary layer, the entrainment of background air in the free troposphere to 232 surface layer, the Miller and Tans et al. (2003) proposed an improved model to compute the 233 biases (Δ) between the recorded and smoothed values. In the present study, the Thoning et al. 234 235 (1989) curve fitting function consist of a 3 polynomial and 4 harmonics terms as described in equation 4, which approximates the long-term trend, short-term variations due to local influence 236 and a non-sinusoidal annual cycle respectively (https://gml.noaa.gov/ccgg/mbl/crvfit/index.html, 237 accessed on 20 July, 2021). 238

$$f(x) = a_0 + a_1 x + a_2 x^2 + a_3 \sin 2\pi x + a_4 \cos 2\pi x + a_5 \sin 4\pi x + a_6 \cos 4\pi x \quad (4)$$

239

here x is time stamp for the input data and f(x) is (CO₂ or δ^{13} C) time dependent variable. The smoothed data, trend, detrended seasonal cycle, seasonal amplitude and growth rate are computed using the equation 4.

Subsequently, we computed biases (Δ) of CO₂, δ^{13} C-CO₂ and δ^{18} O-CO₂ respectively in order to calculate the isotopic signature as explained by Miller and Tans et al. (2003). Following equation is the improved model by Miller and Tans, which was implemented in the present study.

247

$$(\delta^{13}C \times CO_2)_{obs} - (\delta^{13}C \times CO_2)_{smooth} = \delta_s (CO_{2_{obs}} - CO_{2_{smooth}})$$
(5)

248

 $\delta_{s,CO_{2_{obs}}}$ and $CO_{2_{smooth}}$ in equation 5 are slope representing multiple sources, observational CO_{2} 249 and smoothed CO₂ derived from the equation 4 respectively. A linear relationship between ΔCO_2 250 and $\Delta(\delta^{13}C \times CO_2)$ which is popularly known as Miller Tans plot (Miller and Tans, 2003), was 251 further studied. This linear relation is fitted with a ordinary least square (OLS) regression and the 252 slope of the regression line is the carbon isotopic ratio (δ^{13} C) of source CO₂. The Miller Tans 253 model was applied on both day (10:00 Indian Standard Time (IST) till 18:00 IST) and night time 254 (22:00 IST till 06:00 IST) hours during the study period. The OLS is more commonly known as 255 linear regression which may be simple regression or multiple depending on number of 256 explanatory variables. Generic model of the OLS is defined as shown in Equation (6). 257

$$y = a_0 + \sum_{i=1}^{N} \delta_i x_i + \epsilon$$
(6)

258

where y is $\Delta(\delta^{13}C \times CO_2)$, the dependent variable, a_0 is the intercept of the model, δ_i and x_i are the 259 slope and ΔCO_2 corresponds to the ith explanatory variable of the model (i = 1 to n), and ϵ is the 260 random error. Similarly, equations 5 and 6 are implemented for computing the slope between 261 ΔCO_2 versus $\Delta(\delta^{18}O \times CO_2)$. Both the variable in the Miller Tans plot, ΔCO_2 and $\Delta(\delta^{13}C \times CO_2)$, is 262 associated with a measurement error. Moreover, the OLS regression generally yields slopes with 263 larger error (Miller and Tans, 2003), so reduced major axis regression (RMA) (Cantrell, 2008) 264 was further used to estimate the slope. In RMA regression the error in both the variable was 265 accounted while estimating the source value and the standard error obtained was reported. With 266 these methods, further results are discussed in the following sections. 267

268

269 270

271 272

- **3.Results and Discussion**

3.1. Seasonal and Diurnal variation of CO₂, δ^{13} C-CO₂ and δ^{18} O-CO₂

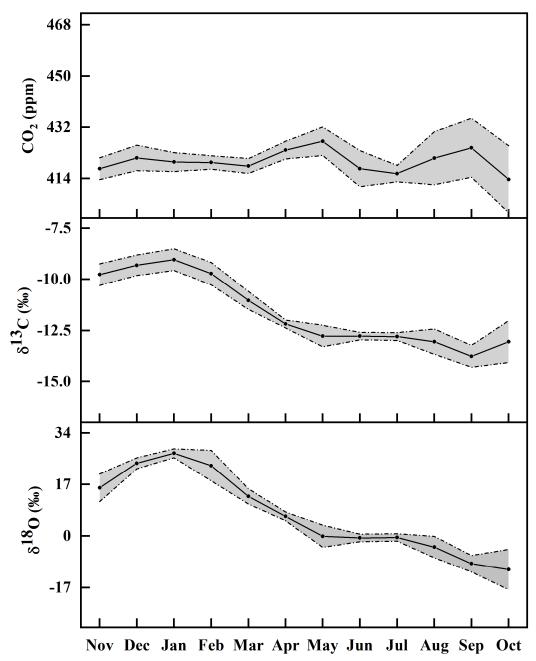




Figure 2. Seasonal variation of atmospheric CO₂ (top), δ^{13} C-CO₂ (middle) and δ^{18} O-CO₂ (bottom). Grey shaded region represents the 1 standard deviation (STD).

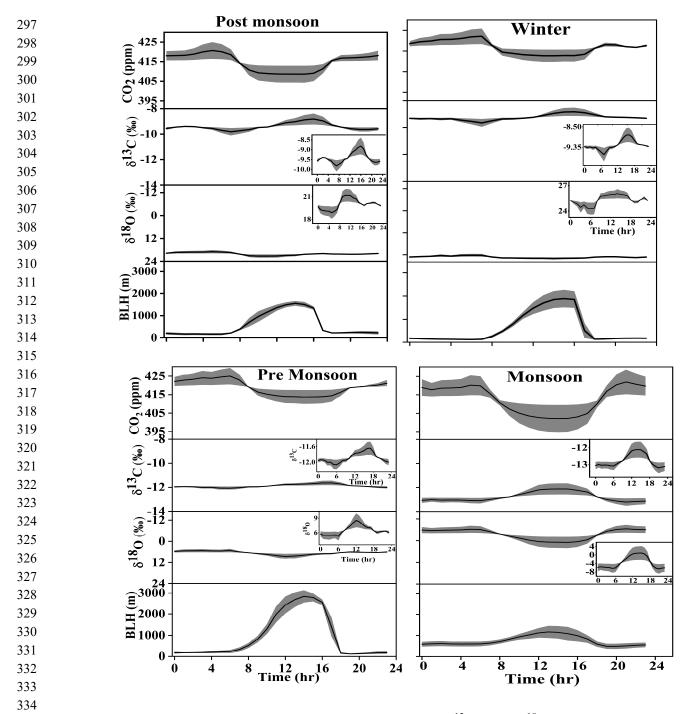


Figure 3. Diurnal variation of seasonally averaged CO₂, δ^{13} C-CO₂, δ^{18} O-CO₂ and boundary layer height during the study period. Grey shaded region represents the 1STD.

The monthly averaged with 1STD of atmospheric CO₂, δ^{13} C-CO₂ and δ^{18} O-CO₂ during the study period are shown in figure 2. At the study site, the annual averaged atmospheric CO₂ concentrations and ¹³C-CO₂ and ¹⁸O-CO₂ are 414.76±4.26 ppm, -11.19±1.63 ‰ and 9.02±12.78 ‰.

The study site experience 4 seasons namely post-monsoon (November-December, vegetation-II), 342 winter (January-February), pre-monsoon (March-May, also known as summer months-dry 343 season) and summer monsoon (June-September; vegetation-I). The NDVI in figure 1 show 344 maximum during monsoon and minimum during rest of the seasons. Seasonal averages of CO₂ 345 $(\delta^{13}C-CO_2 \& \delta^{18}O-CO_2)$ during post-monsoon, winter, pre-monsoon and summer monsoon are 346 414.70±4.49 ppm (-9.41±0.27 ‰ & 20.03±0.73 ‰), 416.84±3.25 ppm (-9.25±0.21 ‰, & 347 25.95±0.57 ‰), 418.88±4.07 ppm (-11.88±0.13 ‰ & 6.51±0.98 ‰), 412.61±7.59 ppm (-348 12.76±0.38 ‰ & -3.49±2.87 ‰) respectively. Site specific δ^{13} C-CO₂ seasonal values with other 349 study sites in India and across the world are summarized in table 2. Liu et al. (2014) showed 350 annual means of δ^{13} C-CO₂ varying from -8.30 ‰ to -8.35 ‰ at Waliguan station in China. 351 Further, the annual means of δ^{13} C-CO₂ are -8.27 ‰ and -8.36 ‰ respectively for 2009 and 2010 352 at Shangdianzi station which is located in a small village about 100 km northeast of Beijing 353 (second populated urban city in China). This site is influenced by strong pollution events from 354 Beijing and surrounding urban areas in the presence of southwesterly winds. Also, the annual 355 mean of δ^{13} C-CO₂ were -8.55‰, -8.52‰, -8.46‰ and -8.61‰ during 2007-2010 at Tae-ahn 356 Peninsula which is located on a small Peninsula on the western coast of Korea. Irrespective of 357 the sites background influence, results of worldwide sites mentioned above indicates δ^{13} C-CO₂ 358 values are in close proximity with our present observations. An increase and decrease of ¹⁸O/¹⁶O 359 $(\delta^{18}O)$ ratio of CO₂ in the atmosphere across the seasons indicates dominance of photosynthesis 360 and ecosystem respiration. During the study period, the atmospheric $\delta^{18}O-CO_2$ varies seasonally 361 from -3.49‰ to + 25.95‰ with large scatter compared to seasonal CO₂ and δ^{13} C. The scatter 362 could be due to vegetation cover, leafwater content, retention of soil water and ocean (Zhou et 363 al., 2006). However, interpretation of seasonal δ^{18} O-CO₂ is not that straight as CO₂ and δ^{13} C-364 CO_2 due to the varying fluxes of biospheric CO_2 and prevailing meteorology at the study site. 365

High atmospheric CO₂ in pre-monsoon followed by winter and subsequent increase of δ^{18} O-CO₂ 366 in the atmosphere in the absence of relatively low photosynthesis could be attributed to long-367 range air mass transport with enhanced δ^{18} O-CO₂ (Murayama et al., 2010) and fossil fuel burning 368 activities Through isotopic exchange, δ^{18} O-CO₂ in the atmosphere may affect due to variation in 369 the ¹⁸O from precipitation and soil respiration (Kato et al., 2004). The range of δ^{18} O-CO₂ at Los 370 Angeles basin, Southern California during 1972 -1973 (1998-2003) is -3.56 ‰ to + 0.21 ‰ (-371 3.99 % to +0.45 %) and average value during 1972-1973 (1998-2003) of δ^{18} O-CO₂ is -1.28 % (-372 1.07‰) (Newman et al., 2008). 373

Thus, figure 3 shows diurnal variations of seasonally averaged CO₂, δ^{13} C-CO₂ and δ^{18} O-CO₂ 374 against BLH. The height of the BL is the vertical extent of air column driven by convection 375 processes associated with the earth's surface heating (Stull, 1988). Due to strong convection and 376 associated surface temperature, the BLH attains maximum height in the afternoon, which 377 modulates the dispersion of air pollutats in the mixed layer. In contrast to this, the BLH quickly 378 dissipates and reaches stable form during night hours due to the absence of convection processes. 379 380 Irrespective of seasons, diurnal variation of atmospheric CO₂ showed minimum (maximum) concentration against maximum (minimum) BLH between 12:00 IST to 16:00 IST (6:00 IST and 381 20:00 IST). The diurnal peak of the BLH is maximum (minimum) in pre-monsoon (summer 382 383 monsoon) with 2839 m (1167 m) respectively, which indicates strong convection in the dry season. Subsequently, observed low (high) CO₂ concentration during strong (absence) 384 convective hours. The observed diurnal cycle of CO₂ is closely associated with diurnal variation 385 of the BLH. The maximum mean value of CO_2 is 418.88 ± 4.07 ppm in pre-monsoon during the 386 study period. An afternoon drop of CO₂ during the study period is associated with the destructed 387

stable BLH and active photosynthetic process in daytime. In contrast, large variability in diurnal atmospheric CO₂ is observed during the summer monsoon with low BLH (1167 m). Thus, exhibits prominent diurnal variations of CO₂ and δ^{13} C-CO₂ against BLH during all the seasons which is also attributed to isotopic fractionation processes during biological activity (Demény and Haszpra 2002). Except for monsoon, diminished diurnal seasonal variability observed with δ^{18} O-CO₂ (Murayama et al., 2010) compared to CO₂ and δ^{13} C-CO₂ at present site. Besides the influence of BLH, minimum CO₂ and maximum δ^{13} C-CO₂ and δ^{18} O-CO₂ during afternoon hours are due to the uptake of CO₂ by the plants through photosynthesis process. During night hours, the concentration CO₂ is maximum with low δ^{13} C-CO₂ and δ^{18} O-CO₂ due to active terrestrial respiration (Sreenivas et al., 2016). Therefore, diurnal patterns of CO₂ and δ^{13} C-CO₂ are anti-correlated during all the seasons with minimum CO_2 in daytime and maximum $\delta^{13}C-CO_2$ respectively (figure 3). Similarly, Pang et al. (2016) observed a negative relationship of δ^{13} C-CO₂ diurnal cycle with CO₂ mixing ratio at Beijing, in Northern China.

402	Study sites	δ^{13} C-CO ₂ values (‰)	References
403		Seasonal	
404	Dallas, USA	-12.0 to -8.1	Clark-Thorne and
405			Yapp, 2003
	Bern, Switzerland	-14 to -8	Sturm et al.,2006
106	Salt Lake City, USA	-18 to -8	Pataki et al., 2006
407		(Dec 2004 - Jan 2005)	
408		-9.3 to -7.5 (Oct)	Djuricin et al., 2010
109	Los Angeles, USA	-12.5 to -8.8 (Dec)	
410		-12.2 to -9.2 (Feb)	
411		-12.5 to -10.2 (April)	
412	Nagoya, Japan	-13.4 to -8.5(May)	Wada et al., 2011
413		-15.0 to -8.5	
114		(Dec from 2008 - 2009)	
415	Krakow, Poland	-11 to -9.5	Zimnoch et al., 2012
416	Wroclaw (SW Poland)	-16.4 to -8.2	Gorka et al., 2013
417	Cabo de Rama	-8.4 to -7.8	Bhattacharya et al.,
	(West Coast of India)		2009
418	Bangalore, India	-9.31 to -8.04	Guha and Ghosh, 2015
419	NRSC, Shadnagar,	-12.8 to -9.3	Present study
420	India	(Nov 2018 - Oct 2019)	

Table 2. δ^{13} C-CO₂ values from different study sites.

Seasonal variability of diurnal atmospheric CO₂ was 9.82 ± 1.39 ppm, 7.19 ± 0.11 ppm, 434 9.23 ±2.1 ppm and 17.30 ±9.29 ppm in post-monsoon, winter, pre-monsoon and summer 435 monsoon respectively during the study period (figure 3). Weak and strong seasonality was 436 observed during winter and monsoon seasons repectively reflecting enrichment of atmospheric 437 CO_2 in low vegetation months and depleting CO_2 in high vegetationseason. The diurnal 438 variability of seasonal δ^{13} C-CO₂ (δ^{18} O-CO₂) are 0.48 ± 0.06 ‰ (1.91±0.005 ‰), 0.44 ±0.02 ‰ 439 $(1.60 \pm 0.13 \text{ }), 0.32 \pm 0.09 \text{ }$ $(2.53 \pm 1.003 \text{ })$ and $0.87 \pm 0.73 \text{ }$ $(6.58 \pm 4.10 \text{ })$ in post-440 monsoon, winter, pre-monsoon and summer monsoon respectively. These amplitudes at our site 441 are lower as compared to urban region listed in Table 2. The Sinhagad (located over the Western 442 Ghats mountains) and Cape Rama, Goa (located close to the shoreline) of western India showed 443 CO₂ seasonal amplitudes between 8-10 ppm during monsoon and >15ppm for remaining seasons 444 (Tiwari et al., 2014), which could be due to the influence of monsoon circulation and strong local 445 -regional biospheric activity (Metya et al., 2021). Thus, modulation in the seasonal amplitude of 446 CO_2 in the atmosphere is attributed to the rate of photosynthesis and respiration besides the 447 impact of local and long-range prevailing meteorology. For better understanding of the 448 relationship between atmospheric CO₂ against its stable isotopic composition, improved model 449 450 by Miller and Tans was adopted by fitting linear regression using the OLS.

451

452 **3.2.** Seasonal correlation between CO₂, and δ^{13} C-CO₂ using the improved model by Miller 453 and Tans

454

Since the study site is not a single sourced location and is closed proximity of Hyderabad, a 455 metropolitan city of India, the Miller-Tans (2003) method was applied on the day and night time 456 observations to characrterize the effective $\delta^{13}C({}^{13}C/{}^{12}C)$ source ratios at the study site. In order 457 to estimate the slope (δ_s) for δ^{13} C-CO₂, the OLS curve fitting was applied to Δ CO₂ against 458 $\Delta(CO_2 \times \delta^{13}C)$ and results shown in Figure 4. The modified Keeling method is applied to capture 459 the processes of day and night which are driving the atmospheric CO₂. During the day (night) 460 time δ^{13} C-CO₂ slopes in post-monsoon, winter, pre-monsoon and monsoon are -10.43±5.60 % (-461 27.08±4.21 ‰), -36.43±8.11 ‰ (-25.11±5.55 ‰), -32.02±4.57 ‰ (-23.79±2.08 ‰) and -462 52.50±4.60 ‰ (-27.30±0.36 ‰) respectively. During day and nighttime hours, the correlation 463 coefficient ('r') was poor (-0.24 to -0.78) in postmonsoon, winter and premonsoon, which is 464 small to estimate the source value. However, a strong 'r' value of -0.91 was found during the 465 nighttime hours in monsoon season (Röckmann et al., 2016; Vardag et al., 2016), thus estimated 466 the source signature (δ_s) using OLS (-27.30±0.36 ‰) and RMA (-37.42±0.73 ‰) methods. The 467 δ^{13} C slope during daytime is largely varied in all the seasons representing the contribution of 468 mixed source emissions at the regional scale and suppressed local sources (Xu et al., 2017). An 469 average value of daytime δ_s of δ^{13} C during all seasons is -32.84 ‰ indicating the source of 470 atmospheric CO₂ is related to gasoline and natural gas combustion in and around the study site 471 (Clark-Thorne and Yapp, 2003). The estimated slopes are evaluated statistically and found 472 significant (p-value < 0.05) with 95 % confidence interval during all the seasons. The δ^{13} C 473 slopes in nighttime during all the seasons are between -25.11 % to -27.39 %, with an average 474 value of -26.72 ‰. In general, the average value of δ_s of δ^{13} C-CO₂ is -26.20 ‰ for C₃ ecosystem 475 respiration (Pataki et al., 2003). Thus, the present study also confirm the dominance of C₃ 476 vegetation at the study site contributes to the emissions of atmospheric CO₂ mixing ratio during 477 night time. The varied δ_s of δ^{13} C-CO₂ in different study locations are attributed to the local 478 anthropogenic activities and terrestrial biospheric pathways. 479

480

Large variability in atmospheric δ^{18} O-CO₂ (Yakir et al., 2011) was noticed during the study 481 period. However, a strong correlation of δ^{18} O with 'r' value of -0.86 was observed (Figure not 482 shown) in night hours of monsoon, which coincidence with the $\delta^{13}C$ during the same season, 483 indicating the active role of biospheric –atmospheric interactions in the exchange of CO₂. With 484 the present study, we understand that interpretation of the seasonal δ^{18} O variation is complicated 485 compared to δ^{13} C and CO₂ mixing ratio. However, current knowledge on δ^{18} O seasonality can be 486 improved with great understanding of varying fluxes of bisopheric CO₂ and role of prevailing 487 meteorology at the study site. Since poor correlation between ΔCO_2 and $\Delta (CO_2 \times \delta^{13}C)$ during day 488 and nighthours in all seasons except nighthours in monsoon, we further implemented OLS and 489 RMA methods only on monsoon data to estimate the source/sink signature (δ_s) of atmospheric 490 CO₂. Figure 4 show improved Miller and Tans method applied on monsoon data during 491 nighthours and fitted with the OLS and RMA linear regression. The deduced slopes from the 492 OLS and RMA are -27.30±0.36 ‰ and -37.42±0.73 ‰ respectively indicating probable 493 contribution from fossil fuel combustion and dominance of C₃ ecosystem respiration. While 494 swapping the x and y axis in the Miller Tans plots, the slopes obtained from the OLS and RMA 495 methods remain statistically same with 'r' value of -0.91. This further confirms the efficacy of 496 RMA regression on the Miller Tans plot in identifying the source CO₂. 497

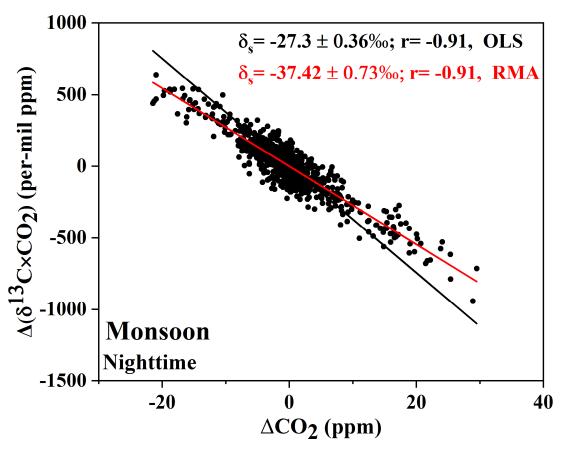
The δ_s of δ^{13} C analysis at the study site confirms the prominence of C₃ ecosystem activities viz. photosynthesis and respiration in the variability of atmospheric CO₂ during vegetative seasons. During dry seasons (winter and pre-monsoon), the δ_s values indicates the important role of local and long-range combustion processes that enrich atmospheric CO₂ at the study site. However, this information is pertained to the total source contributions and individual source apportionment was not carried out in this study.

504	
505	

	$\begin{array}{c} \text{Daytime} \\ \Delta \text{CO}_2 \text{vs} \Delta (\text{CO}_2 \times \delta^{13} \text{C}) \end{array}$		Nighttime $\Delta CO_2 vs \Delta (CO_2 \times \delta^{13}C)$	
Season				
	ʻr'	Slope (‰)	ʻr'	Slope(‰)
Postmonsoon	-0.24	-10.43 ± 5.60	-0.64	-27.08±4.21
Winter	-0.51	-36.43±8.11	-0.51	-25.11±5.55
Premonsoon	-0.61	-32.02±4.57	-0.78	-27.39±2.08
Monsoon	-0.75	-52.50 ± 4.60	-0.91	-27.30±0.36
				(-37.42±0.73)*
*indicates the slope calculated using RMA method for the nighttime da during monsoon			the nighttime data	

517 **Table 3.** Seasonal correlation coefficient (r), and slope (δ s) derived from the improved Miller 518 and Tans method fitted with the OLS and RMA.

519



522

Figure 4. Improved Miller and Tans method applied to the monsoon seasonal $\Delta CO_2 vs\Delta(CO_2 \times \delta^{13}C)$ during nighttime. The solid line black line and red coloar line represents the regression fit using ordinary least square method and reduced major axis method respectively.

Table 3 shows seasonal 'r' and slope values derived between $\Delta CO_2 vs \Delta (CO_2 \times \delta^{13}C)$ and 527 $\Delta CO_2 vs \Delta (CO_2 \times \delta^{18}O)$ during day and night hours respectively. During nighttime the average 528 value of slope at the study site is close to C₃ ecosystem respiration and during daytime the 529 average value of slope is close to sources which are combustion, photosynthesis processes and 530 long range transport. The differences in slope value at various study sites is due to photosynthesis 531 pathways and local anthropogenic sources prevailing at the study areas (Pang et al., 2016). In 532 533 similar studies carried by various authors, reported slope values at their study site which is as follows. Liu et al. (2014) observed annual mean value of slope which is -25.44‰ and -21.70 ‰ 534 at Waliguan and Shangdianzi stations in China respectively. Zhou et al. (2006) studied the 535 isotopic fractionation at 11 stations of Northern Hemisphere and found slope (δ_s) ranging from -536 28.85‰ to -26.50‰ with improved Miller-Tans method. Murayama et al.(2010) observed δ_s 537 value of -28.7‰ averaged over the study period at Takayama site in central Japan, which is also 538 539 comparable to the present study site.

The correlation coefficient (r) between δ^{18} O-CO₂ and δ^{13} C-CO₂ during daytime (night time) are 0.13(0.07), 0.64(0.60), 0.95(0.93), 0.24(0.92) for post-monsoon, winter, pre-monsoon and summer monsoon seasons respectively. A very strong positive correlation (0.93 to 0.95) in day and night hoursduring pre-monsoon between δ^{18} O-CO₂ and δ^{13} C-CO₂ might be due to high vapour pressure deficit (VPD) than the post monsoon (low VPD). The high VPD and summer

conditions are supportive for high transpiration as well as photosynthetic activities as sunshine 545 hours are more during pre-monsoon than post monsoon (approaching towards winter). Low δ^{13} C-546 CO_2 and $\delta^{18}O$ - CO_2 correlation during post-monsoon indicates high stomatal conductance in 547 vegetation, ensuring enhanced release of water to the atmosphere (Cullen et al., 2008; Liu et al., 548 2014). Our study showed clear evidence of an increase or decrease of atmospheric CO₂ is 549 associated with the changes of its isotopic composition during photosynthesis and biogenic 550 respiration besides local and long-range anthropogenic influences. Also, Zimnoch et al. (2004) 551 observed good correlation between δ^{18} O-CO₂ and δ^{13} C-CO₂ with 'r' ranging from 0.84 to 0.94 in 552 their observations while studying their diurnal variability from Poland. Further to understand the 553 influence of transport pathways on atmospheric CO₂ via long range airmass is examined in the 554 present study using Hybrid Single Particle Lagrangian Integrated Trajectory Model (HYSPLIT) 555 along with the forest fire count. 556

557

558 **3.3. Influence of local and long-range airmass on atmospheric CO₂**

Study site being a suburban region and about 60 km away from the Hyderabad city, the analysis 559 of improved Miller-Tan model depicts atmospheric CO₂ concentration at the study site is mainly 560 controlled by terrestrial biosphere activities during the study period. During the dayhours, the 561 derived δ_s of δ^{13} C indicating mixed source contribution at the study site, which may be possibly 562 due to combustion activities, biomass burning and the transportation of airmass. Seasonal wind 563 vector obtained from the ECMWF at 850 hPa over the Indian region are shown in figure 5a. The 564 long-range airmass circulation that is reaching the study site has been analyzed using Lagrangian 565 back-trajectory model along with the firecounts during all the seasons (Figure 5b). 566

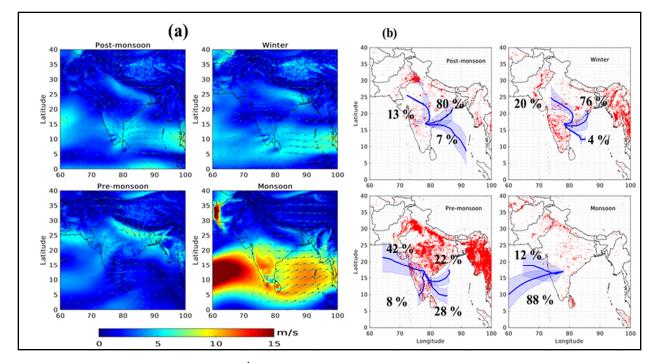


Figure 5. (a) Mean winds (m s⁻¹) at 850 hPa (b) Seasonal long-range air mass circulation at 2
 km altitude using HYSPLIT overlaid by forest fire count.

During the Indian summer-monsoon season, the south-westerly(SW) winds as shown in Figure 570 5a are in general dominate, which brings maritime airmass (Bhattacharya et al., 2009; Tiwari et 571 al., 2014; Guha and Ghosh, 2015) over the Indian region. The atmospheric CO₂ concentration at 572 the study site is observed minimum during the summer-monsoon period indicating the 573 contribution of monsoon circulation through scavenging effect (Tiwari et al., 2014; Mahesh et 574 al., 2014). The maritime airmass during summer-monsoon is relatively pristine due to the 575 absence of anthropogenic sources thus observed low atmospheric CO_2 concentration at the study 576 site. 577

578

A Lagrangian back trajectory analysis as shown in Figure 5b also confirms the observed low 579 atmospheric CO₂ due to influence of maritime airmass (88 %) during the summer-monsoon 580 period. During winter and pre-monsoon seasons, enrichment of atmospheric CO₂ at the study site 581 could be due to the predominant continental sources reaching from the north-east (NE, 76 %) and 582 north-west (NW, 42 %) respectively. Figure5b shows seasonal long-range air mass circulation 583 over laid with fire count obtained from Moderate Resolution Imaging Spectroradiometer 584 (MODIS) during the study period. During pre-monsoon and post-monsoon, agriculture residue 585 burning is commonly observed in Punjab, Haryana and Indo Gangetic Plains (IGBP) areas of 586 India (Liu et al., 2019), which are in NW and NE directions of the study site. Thus, the present 587 study reveals that during these seasons, the biomass burning is one of continental sources 588 589 contributing to the elevated CO₂ concentration. Therefore the variability of atmospheric CO₂ concentration at the study site is greatly influenced by the maritime airmass during summer-590 591 monsoon and continental sources during other seasons.

592

593 **4. Conclusion**

594

The present study examined the diurnal and seasonal variations of atmospheric CO₂ and its stable isotopes δ^{13} C-CO₂ and δ^{18} O-CO₂ at the sub-urban site of India during November 2018 to October 2019 using high precision CO₂ isotopic analyzer.

598 Following are the salient findings of the present study

599 600

601

602

1. The atmospheric CO₂ concentration ranged from 405 ppm to 450 ppm, with its stable isotopic composition ranging from -7.72 to -14.55 % VPDB for δ^{13} C and from -24.63 to 28.91 % VPDB for δ^{18} O respectively.

- 603 2. The CO₂ and δ^{13} C-CO₂ exhibit clear diurnal variation with opposite patterns during all 604 the seasons with minimum CO₂ and maximum δ^{13} C-CO₂ in daytime and vice versa.
- Buring strong convective hours (peak BLH) low atmospheric CO₂ concentration was observed. High CO₂ concentration was observed in weak convective period which indicates strong atmospheric mixing.
- 6084. Diurnal variability of seasonal atmospheric CO_2 at the study site was 9.82 ± 1.39 ppm,609 7.19 ± 0.11 ppm, 9.23 ± 2.1 ppm and 17.30 ± 9.29 ppm in post-monsoon, winter, pre-610monsoon and summer monsoon respectively. Large seasonality in summer-monsoon at611the study site due to strong influence of monsoonal winds, precipitation and vegetation612cover.
- 5. To capture the driving processes of atmospheric CO₂ during day and night hours, we thus applied improved Miller-Tans model on seasonal ΔCO_2 , $\Delta (CO_2 \times \delta^{13}C)$ and $\Delta (CO_2 \times \delta^{13}C)$

615	$\times \delta^{18}$ O) data. During all seasons, the nighttime δ_s values were between -25.11 ‰ to -
616	27.39 ‰, with an average value of -26.72 ‰ with moderate 'r' value.
617	6. However, a strong source/sink signature of $\delta^{13}C$ was observed during nighttime of
618	summer monsoon with the slope of -37.42±0.73 ‰, obtained using a reduced major axis
619	(RMA) regression. The source identified is attributed to combustion and dominance of
620	C_3 ecosystem respiration respectively.
621	7. Seasonal 'r' value between δ^{18} O-CO ₂ and δ^{13} C-CO ₂ during day and night time are varied
622	0.13 to 0.95 and 0.07 to 0.93 respectively. A very strong positive correlation (0.93 to
623	0.95) in day and night hours during pre-monsoon between $\delta^{18}O-CO_2$ and $\delta^{13}C-CO_2$
624	might be due to high VPD than the post monsoon (low VPD).
625	8. Upwind transport confirms the influence of biomass burning on enriched atmospheric
626	CO_2 during pre-monsson and post-monsoon seasons.
627	9. A lagrangian back-trajectories confirms the variability of atmospheric CO ₂ concentration
628	at the study site is largely influenced by the maritime airmass during summer-monsoon
629	and continental sources in other seasons.
630	
631	Our study showed clear evidence of an increase or decrease of atmospheric CO ₂ is
632	associated with the changes of its isotopic composition during photosynthesis and
633	biogenic respiration besides local and long-range anthropogenic influences. The
634	variability in atmospheric CO ₂ during monsoon season is strongly associated with the
635	ISM. However, the individual source apportionment of different sources is not discussed
636	in this present study.
637	Acknowledgements
638	Authors sincerely thank Dr Raj Kumar, Director NRSC for his kind encouragement and support
639	to carry out this work. This work was part of the Atmospheric CO_2 Retrievals and Monitoring
640	(ACRM) of the National Carbon Project (NCP) funded by CAP-IGBP. We greatly thank Dr. V.K
641	Dadhwal, Former Director, Indian Institute of Space Science and Technology, Trivandrum,
642	India and Project Director, NCP for reviewing the manuscript. Authors thank Dr. M.V.R
643	SeshaSai, Deputy Director, ECSA for his encouragement to carry this study. We greatly
644	acknowledge the HYSPLIT, ECMWF-ERA and MODIS fire teams for providing the scientific
645	data sets used in this study.
646	
647	Declaration of competing interest
648	Declaration of competing interest
649	The authors declare no competing interests.
650	
651 (52	Data Availabilty Statement
652	Data Availabilty Statement
653 654	The in situ data may be available publicly once archival is completed
654	The <i>in- situ</i> data may be available publicly once archival is completed.
655	

- **References**

Andres, R.J., Marland, G., Fung, I., & Matthews, E. (1996). A 1°× 1° distribution of carbon
dioxide emissions from fossil fuel consumption and cement manufacture, 1950–1990. *Global Biogeochem Cycles*, 10(3), 419-429.https://doi.org/10.1029/96GB01523.

- Baer, D. S., Paul, J. B., Gupta, M., & O'Keefe, A. (2002). In Diode Lasers and Applications in
 Atmospheric Sensing; Fried, A., Ed. In SPIE-The International Society for Optical Engineering:
 Bellingham, WA, Vol. 4817, pp. 167-176.
- Ballantyne, A.P., Alden, C.B., Miller, J.B., Tans, P.P., & White, J.W.C. (2012). Increase in
 observed net carbon dioxide uptake by land and oceans during the past 50 years. *Nature*, 488,
 70-72, doi:10.1038/nature11299.
- 670
- Bhattacharya, S.K., Borole, D.V., Francey, R.J., Allison, C.E., Steele, L.P., Krummel, P., et al.
 (2009). Trace gases and CO₂ isotope records from Cabo de Rama, India. *Current Science* 97(9),
 1336-1344.
- 674
- Chakraborty S., Tiwari Y.K., Deb Burman P.K., Baidya Roy S., Valsala V. (2020) Observations
 and Modeling of GHG Concentrations and Fluxes Over India. In: Krishnan R., Sanjay J.,
 Gnanaseelan C., Mujumdar M., Kulkarni A., Chakraborty S. (eds) Assessment of Climate
 Change over the Indian Region. Springer, Singapore. https://doi.org/10.1007/978-981-15-43272_4
- 680
- ⁶⁸¹ Cullen, L. E., Adams, M. A., Anderson, M. J., & Grierson, P. F. (2008). Analyses of δ^{13} C and ⁶⁸² δ^{18} O in tree rings of Callitriscolumellaris provide evidence of a change in stomatal control of ⁶⁸³ photosynthesis in response to regional changes in climate. Tree Physiology, 28(10), 1525-1533. ⁶⁸⁴ doi: 10.1093/treephys/28.10.1525.
- 685
- Djuricin, S., Pataki, D.E., &Xu, X. (2010). A comparison of tracer methods for quantifying
- 687 CO₂ sources in an urban region. *Journal of Geophysical Research: Atomspheres* 115, 1–13, 688 https://doi.org/10.1029/2009JD012236.
- 689
- Demény A, Haszpra L (2002) Stable isotope compositions of CO₂ in background air and at
 polluted sites in Hungary. Rapid Commun Mass Spectrom 16:797–804
- 693 Farquhar, G. D., Lloyd, J., Taylor, J.A., Flanagan, L.B., Syvertsen, J. P., et al. (1993).
- Vegetation effects on the isotope composition of oxygen in atmospheric CO₂. *Nature*, 363(6428),
 439-443,doi: 10.1038/365368b0
- 696
- Francey, R. J., & Tans, P. P. (1987). Latitudinal variation in oxygen-18 of atmospheric CO₂.
 Nature, 327(6122), 495-497.
- 699
- 700 Górka, M., &Lewicka-Szczebak, D. (2013). One-year spatial and temporal monitoring of
- concentration and carbon isotopic composition of atmospheric CO₂ in a Wrocław (SW Poland)
- city area. Applied geochemistry, 35, 7-13. https://doi.org/10.1016/j.apgeochem.2013.05.010.
- 703

Guillon, S., Agrinier, P., &Pili, E., (2015). Monitoring CO₂ concentration and δ^{13} C in an underground cavity using a commercial isotope ratio infrared spectrometer. *Applied Physics B-Lasers and optics*, 119(1), 165-175. doi: 10.1007/s00340-015-6013-4.

- ⁷⁰⁷ Guha T, Ghosh P (2010) Diurnal variation of atmospheric CO₂ concentration and δ^{13} C in an urban atmosphere during winter—role of the Nocturnal Boundary Layer. *J AtmosChem* 65:1–12. doi:10.1007/s10874-010-9178-6.
- 711
- 712 Guha, T, and Ghosh P (2013) An experimental set-up for carbon isotopic analysis of 713 atmospheric CO_2 and an example of ecosystem response during solar eclipse 2010. *Journal of* 714 *Earth System Science* 122.3 (2013): 623-638.
- 715
- Guha, T.and Ghosh, P. (2015). Diurnal and seasonal variation of mixing ratio and δ^{13} C of air CO₂ observed at an urban station Bangalore, India. Environmental Science and Pollution Research, 22(3), 1877-1890. https://doi.org/10.1007/s11356-014-3530-3.
- 719
- Huang, J., Yu, H., Guan, X., Wang, G., &Guo, R., (2016). Accelerated dryland expansion under
 climate change, *Nature Climate Change*, 6, 166–171, http://dx.doi.org/10.1038/nclimate2837.
- 722
- Kato, T., Nakazawa, T., Aoki, S., Sugawara, S., & Ishizawa, M. (2004). Seasonal variation of the
 oxygen isotopic ratio of atmospheric carbon dioxide in a temperate forest, Japan. *Global Biogeochem Cycles*, 18(2),https://doi.org/10.1029/2003GB002173.
- 726
- Liu, L., Zhou, L., Vaughn, B., Miller, J. B., Brand, W. A., Rothe, M., & Xia, L. (2014). Background variations of atmospheric CO₂ and carbon-stable isotopes at Waliguan and Shangdianzi stations in China. *Journal of Geophysical Research: Atmospheres*, 119(9), 5602-5612.
- 731
- Liu, X., An, W., Leavitt, S. W., Wang, W., Xu, G., Zeng, X., & Qin, D. 2014. Recent strengthening of correlations between tree-ring δ^{13} C and δ^{18} O in mesic western China: Implications to climatic reconstruction and physiological responses. Global and Planetary Change, 113, 23-33.http://dx.doi.org/10.1016/j.gloplacha.2013.12.005
- 736
- Liu, T., Marlier, M. E., Karambelas, A., Jain, M., Singh, S., Singh, M. K., Gautam, R.,
 &DeFries, R.S. (2019). Missing emissions from post-monsoon agricultural fires in northwestern
 India: regional limitations of MODIS burned area and active fire products.*Environmental Research Communications* 1(1), 011007,https://doi.org/10.1088/2515-7620/ab056c.
- 741
- Machida, T., Kita, K., Kondo, Y., Blake, D., Kawakami, S., Inoue, G., & Ogawa, T. (2002).
 Vertical and meridional distributions of the atmospheric CO₂ mixing ratio between northern midlatitudes and southern subtropics. *Journal of Geophyical Research: Atmospheres* 107(D3),
- 745 BIB 5-1-BIB 5-9,https://doi.org/10.1029/2001JD000910.
- Mahesh, P., Sreenivas, G., Rao, P. V. N., Dadhwal, V. K., Sai Krishna, S. V. S., & Mallikarjun,
 K. (2015). High-precision surface-level CO₂ and CH₄ using off-axis integrated cavity output

spectroscopy (OA-ICOS) over Shadnagar, India.*International Journal of Remote Sensing* ,36(22), 5754-5765,https://doi.org/10.1080/01431161.2015.1104744.

Mahesh, P., Sreenivas, G., Rao, P. V. N., &Dadhwal, V. K. (2016). Atmospheric CO₂ retrieval
 from ground based FTIR spectrometer over Shadnagar, India.*Atmospheric Measurement Techniques Discussions*, https://doi.org/10.5194/amt-2016-177, 2016.

- 753
- Mahalakshmi, D.V., Sujatha, P., Naidu , CV., and Chowdary, V.M. (2014). Contribution of
- vehicular emissions on urban air quality: results from public strike in Hyderabad.Indian Journal
- 756 of Radio & Space Physics (IJRSP), 43(6), 340-348.
- 757

761

- Metya, A., Datye, A., Chakraborty, S., Tiwari, Y. K., Sarma, D., Bora, A., and Gogoi, N. (2021).
 Diurnal and seasonal variability of CO₂ and CH₄ concentration in a semi-urban environment of
 western India. Scientific reports, 11(1), 1-13.
- Miller JB and Tans PP (2003) Calculating isotopic fractionation from atmospheric measurements
 at various scales. Tellus Series B: Chemical and Physical Meteorology 55: 207–214.
- Monastersky, R. (2013). Global carbon dioxide levels near worrisome milestone. *Nature*, 497 (7447), 13–14, https://doi.org/10.1038/497013a.
- 767
- ⁷⁶⁸ Murayama, S., C. Takamura, S. Yamamoto, N. Saigusa, S. Morimoto, H. Kondo, T. Nakazawa, ⁷⁶⁹ S. Aoki, T. Usami, and M. Kondo (2010), Seasonal variations of atmospheric CO₂, δ^{13} C, and ⁷⁷⁰ δ^{18} O at a cool temperate deciduous forest in Japan: Influence of Asian monsoon, J. Geophys. ⁷⁷¹ Res., 115, D17304, doi:10.1029/2009JD013626.
- 772
- Nalini, K., Sijikumar, S., Valsala, V., Tiwari, Y. K., & Ramachandran, R. (2019). Designing
 surface CO₂ monitoring network to constrain the Indian land fluxes. *Atmospheric Environment*218(1), 117003, https://doi.org/10.1016/j.atmosenv.2019.117003
- 776
- Newman, S., Xu, X., Affek, H. P., Stolper, E., & Epstein, S. (2008). Changes in mixing ratio and
 isotopic composition of CO₂ in urban air from the Los Angeles basin, California, between 1972
 and 2003. *Journal of Geophysical Research: Atmospheres*,
 113(D23).https://doi.org/10.1029/2008JD009999
- 781
- Pang, J., Wen, X., & Sun, X. (2016). Mixing ratio and carbon isotopic composition investigation
 of atmospheric CO₂ in Beijing, China. *Science of Total Environment* 539, 322-330,DOI:
 10.1016/j.scitotenv.2015.08.130.
- 785
- Pataki, D. E., Ehleringer, J. R., Flanagan, L. B., Yakir, D., Bowling, D. R., Still, C. J., ... &
 Berry, J. A. (2003). The application and interpretation of Keeling plots in terrestrial carbon cycle
 research. Global biogeochemical cycles, 17(1).
- Pataki, D.E., Bowling, D.R., Ehleringer, J.R., & Zobitz, J.M. (2006). High resolution atmospheric monitoring of urban carbon dioxide sources. *Geophysical Research Letters* 33, 1–
- 792 5,https://doi.org/10.1029/2005GL024822.

793

Röckmann, T., Eyer, S., Veen, C. van der, Popa, M.E., Tuzson, B., Monteil, G., Houweling, S.,
Harris, E., Brunner, D., Fischer, H., Zazzeri, G., Lowry, D., Nisbet, E.G., Brand, W.A., Necki,
J.M., Emmenegger, L., Mohn, J. (2016). In situ observations of the isotopic composition of
methane at the Cabauw tall tower site. Atmospheric chemistry and physics, 16(16), 1046910487.https://doi.org/10.5194/acp-16-10469-2016.

- Sharma, N., Dadhwal, V. K., Kant, Y., Mahesh, P., Mallikarjun, K., Gadavi, H., Sharma, A., &
 Ali, M. M. (2014). Atmospheric CO₂ variations in two contrasting environmental sites over
- India. Air soil and water Research, 7, ASWR-S13987, https://doi.org/10.4137/ASWR.S13987.
 803
- Smith, H. J., Fischer, H., Wahlen, M., Mastroianni, D., & Deck, B. (1999). Dual modes of the
 carbon cycle since the Last Glacial Maximum. *Nature*, 400(6741), 248-250.
- Sreenivas, G., Mahesh, P., Subin, J., Kanchana, A. L., Rao, P. V. N., &Dadhwal, V. K. (2016). 807 Influence of meteorology and interrelationship with greenhouse gases (CO₂ and CH₄) at a 808 809 suburban site of India. *Atmospheric* Chemistry and **Physics** 16, 3953-3967, doi.org/10.5194/acp-16-3953-2016, 2016. 810
- 811
- Sturm, P., Leuenberger, M., Valentino, F.L., Lehmann, B., &Ihly, B. (2006). Measurements of
- CO₂, its stable isotopes, O₂/N₂, and Rn-222 at Bern, Switzerland. *Atmospheric Chemistry and Physics*. 6, 1991-2004, https://doi.org/10.5194/acp-6-1991-2006, 2006.
- 815
- Thorne, S. T., & Yapp, C. J. (2003). Stable carbon isotope constraints on mixing and mass
 balance of CO₂ in an urban atmosphere: Dallas metropolitan area, Texas, USA. *Applied Geochemistry*, 18(1), 75-95, https://doi.org/10.1016/S0883-2927(02)00054-9.
- 819
- Thoning, K. W., Tans, P. P., &Komhyr, W. D. (1989). Atmospheric carbon dioxide at Mauna
 Loa Observatory: 2. Analysis of the NOAA GMCC data, 1974–1985. Journal of Geophysical
 Research: Atmospheres, 94(D6), 8549-8565.
- 823
- Tiwari, Y. K., Vellore, R. K., Kumar, K. R., van der Schoot, M., & Cho, C. H. (2014). Influence
- of monsoons on atmospheric CO₂ spatial variability and ground-based monitoring over India.
- *Science of The Total Environment*, 490, 570-578, https://doi.org/10.1016/j.scitotenv.2014.05.045
- Valsala, V., Tiwari, Y. K., Pillai, P., Roxy, M., Maksyutov, S., &Murtugudde, R. (2013).
 Intraseasonal variability of terrestrial biospheric CO₂ fluxes over India during summer
 monsoons. *Journal of Geophysical Research-Biogeosciences*, 118(2), 752-769,
 https://doi.org/10.1002/jgrg.20037.
- 832
- Vardag, S. N., Hammer, S., & Levin, I. (2016). Evaluation of 4 years of continuous $\delta^{13}C$ (CO₂) data using a moving Keeling plot method. Biogeosciences, 13(14), 4237-4251.https://doi.org/10.5194/bg-13-4237-2016.
- 836
- 837 Wada, R., Pearce, J.K., Nakayama, T., Matsumi, Y., Hiyama, T., Inoue, G., & Shibata, T.(2011).
- 838 Observation of carbon and oxygen isotopic compositions of CO₂ at an urban site in

- Nagoya using mid-IR laser absorption spectroscopy. *Atmospheric Environment*, 45, 1168–
 1174,https://doi.org/10.1016/j.atmosenv.2010.10.015.
- 841

Xu, W., Ruhl, M., Jenkyns, H.C., Hesselbo, S.P., Riding, J.B., et al. (2017). Carbon sequestration
in an expanded lake system during the Toarcian oceanic anoxic event. *Nature Geoscience*, 10,
129-134, doi: 10.1038/NGEO2871

- Yakir, D. (2011). The paper trail of the ¹³C of atmospheric CO₂ since the industrial revolution
 period. *Environmental Research Letters* 6(3), 034007, http://dx.doi.org/10.1088/17489326/6/3/034007.
- 849

845

Zhou, L., Conway, T. J., White, J. W., Mukai, H., Zhang, X., Wen, Y., Li, Jinlon.,&MacClune,
K. (2005). Long-term record of atmospheric CO₂ and stable isotopic ratios at Waliguan
Observatory: Background features and possible drivers, 1991–2002. *Global Biogeochemical Cycles*, 19(3), https://doi.org/10.1029/2004GB002430.

854

Zhou, L., J. W. C. White, T. J. Conway, H. Mukai, K. MacClune, X. Zhang, Y. Wen, and J. Li
(2006), Long-term record ofatmospheric CO₂ and stable isotopic ratios at Waliguan Observatory:
Seasonally averaged 1991–2002 source/sink signals, and acomparison of 1998–2002 record to
the 11 selected sites in the Northern Hemisphere, Global Biogeochem. Cycles, 20,
GB2001,doi:10.1029/2004GB002431.

860 861

Zimnoch, M., Florkowski, T., Necki, J. M., &Neubert, R. E. (2004). Diurnal variability of δ^{13} C and δ^{18} O of atmospheric CO₂ in the urban atmosphere of Kraków, Poland. Isotopes in Environmental and Health Studies, 40, 129-143. https://doi.org/10.1080/10256010410001670989.