

1 **Influence of Meteorology and interrelationship with greenhouse gases (CO₂**
2 **and CH₄) at a suburban site of India**

3 Sreenivas. G, Mahesh. P*, Subin Jose, Kanchana A. L., Rao P.V.N, Dadhwal. V.K

4 Atmospheric and Climate Sciences Group (ACSG),

5 Earth and Climate Science Area (ECSA),

6 National Remote Sensing Center (NRSC),

7 Indian Space Research Organization (ISRO),

8 Hyderabad, India-500037

9
10 *Corresponding author: Mahesh P

11 Mail-Id:mahi952@gmail.com

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29 **Abstract**

30 Atmospheric greenhouse gases (GHGs), such as carbon dioxide (CO₂) and methane (CH₄),
31 are important climate forcing agents due to their significant impacts on the climate system. The
32 present study brings out first continuous measurements of atmospheric GHGs using high precision
33 LGR-GGA over Shadnagar, a suburban site of Central India during the period 2014. The annual
34 mean CO₂ and CH₄ over the study region are found to be 394±2.92 ppm and 1.92±0.07 ppm (μ
35 $\pm 1\sigma$) respectively. CO₂ and CH₄ show a significant seasonal variation during the study period with
36 maximum (minimum) CO₂ observed during Pre-monsoon (Monsoon), while CH₄ recorded
37 maximum during post-monsoon and minimum in monsoon. Irrespective of the seasons, consistent
38 diurnal variations of these gases are observed. Influences of prevailing meteorology (air
39 temperature, wind speed, wind direction and relative humidity) on GHGs have also been
40 investigated. CO₂ and CH₄ show a strong positive correlation during winter, pre-monsoon,
41 monsoon, and post-monsoon with correlation coefficients (Rs) equal to 0.80, 0.80, 0.61, and 0.72
42 respectively; indicating common anthropogenic source for these gases. Analysis of this study
43 reveals the major sources for CO₂ are soil respiration and anthropogenic emissions while
44 vegetation act as a main sink. Whereas the major source and sink for CH₄ are vegetation and
45 presence of hydroxyl (OH) radicals.

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47 Keywords: Carbon dioxide, Methane, OH radical.

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58 **1. Introduction**

59 The Intergovernmental Panel on Climate Change (IPCC, 2013) reported that humankind is
60 causing global warming through the emission of greenhouse gases (GHGs), particularly carbon
61 dioxide (CO₂) and methane (CH₄). CO₂ and CH₄ concentrations have increased by 40% and 150 %
62 respectively since pre-industrial times, mainly from fossil fuel emissions and secondarily from net
63 land use change emissions (IPCC, 2013; Huang et al., 2015). CO₂ measurements at MaunaLoa,
64 Hawaii (Monastersky, 2013) have exceeded the 400 ppm mark several times in May 2013. CH₄ is
65 also receiving increasing attention due to high uncertainty in its sources and sinks (Keppler et al.,
66 2006; Miller et al., 2007; Frankenberg et al., 2008). Stefanie Kirschke et al., (2013) reported that
67 in India, agriculture and waste constitute the single largest regional source of CH₄. Although many
68 sources and sinks have been identified for CH₄, their relative contribution to atmospheric CH₄ is
69 still uncertain (A. Garg et al., 2001; StefanieKirschke et al., 2013). In India, electric power
70 generation contributes to half of India's total CO₂ equivalent emissions (A. Garg et al., 2001).

71 Arid and semi-arid areas comprise about 30% of the Earth's land surface. Climate change and
72 climate variability will likely have a significant impact on these regions (Huang et al., 2008; Huang
73 et al., 2015). The variability of environmental factors may result in significant effects on regional
74 climate and global climate (Wang et al., 2010), especially the radiative forcing; via the
75 biogeochemical pathways affecting the terrestrial carbon cycle. Global climate change has serious
76 impact on humans and ecosystems. Due to this, many factors have been identified that may reflect
77 or cause variations in environmental change (Pielke et al., 2002). Out of these, the Normalized
78 Difference Vegetation Index (NDVI) has become one of the most widely used indices to represent
79 the biosphere influence on global change (Liu et al., 2011). Greenhouse and other trace gases have
80 great importance in atmospheric chemistry and for radiation budget of the atmosphere-biosphere
81 system (Crutzen et al., 1991). Hydroxyl radicals (OH) are very reactive oxidizing agents, which
82 are responsible for the oxidation of almost all gases that are emitted by natural and anthropogenic
83 activities in the atmosphere. Atmospheric CO₂ measurements are very important for understanding
84 the carbon cycle because CO₂ mixing ratios in the atmosphere are strongly affected by
85 photosynthesis, respiration, oxidation of organic matter, biomass and fossil fuel burning, and air-
86 sea exchange process (Machida et al., 2003).

87 The present study brings out first continuous measurements of atmospheric GHGs using high
88 precision LGR-GGA over Shadnagar, a suburban site of Central India during the period 2014. In
89 addition to GHGs observations, we have also made use of an automatic weather station (AWS)
90 data along with model/satellite retrieved observation during the study period. Details about study
91 area and data sets are described in the following sections.

92

93 **2. Study Area**

94 Shadnagar is situated in Mahabubnagar district of newly formed Indian state of Telangana.
95 It is a suburban location situated ~70km away from urban site of Hyderabad (Northern side) with
96 a population of ~0.16 million (Patil et al., 2013). A schematic map of study area is shown in Fig.
97 1a. Major sources of pollutants over Shadnagar can be from small and medium scale industries,
98 biomass burning and bio-fuel as well as from domestic cooking. In the present study sampling of
99 GHGs and related meteorological parameters are carried out in the premises of National Remote
100 Sensing Center (NRSC), Shadnagar Campus (17°02'N, 78°11'E). Sampling site is near (aerial
101 distance ~ 2.25 km) to National highway 7 (NH7) and a railway track (non-electrified) is in the
102 East (E) direction.

103 Mean monthly variations of temperature (°C) and relative humidity (RH %) observed at
104 Shadnagar during 2014 are shown in Figure 1e and 1d respectively. The Indian Meteorological
105 Department (IMD) defined monsoon as June-July-August-September (JJAS), post-monsoon
106 (October-November-December-OND), winter (January-February-JF) and pre-monsoon (March-
107 April-May-MAM) in India. Temperature over Shadnagar varies from ~20°C to ~29°C. Relative
108 humidity (RH) in Shadnagar reached a maximum of ~82 % in monsoon from a minimum of ~48
109 % recorded during pre-monsoon. Surface wind speed (Fig. 1c) varies between 1.3 to 1.6 m s⁻¹ with
110 a maximum observed during monsoon and minimum in pre-monsoon. The air mass advecting (Fig.
111 1b) towards study site is either easterly or westerly. The easterly wind prevails during winter and
112 gradually shifts to south-westerlies in pre-monsoon, and dominates during monsoon.

113 **3. Data set and Methodology**

114 Details about the instrument and data utilized are discussed in this section. The availability and
115 frequency of the observations all data used in present study are tabulated in Table 1.

116

117 3.1 *In-situ observations*

118 3.1.1 Greenhouse Gas Analyser (GGA)

119 The Los Gatos Research's - Greenhouse Gas Analyser (model: LGR-GGA-24EP) is an
120 advanced instrument capable of simultaneous measurements of CO₂, CH₄ and H₂O. This
121 instrument is well known for high precision and accuracy which are crucial towards understanding
122 background concentrations of atmospheric GHGs, with specifications meeting WMO standards of
123 measurement (Berman et al., 2012; Shea et al., 2013; Mahesh et al., 2015). It is based on enhanced
124 Off-Axis Integrated Cavity Output Spectroscopy (OA-ICOS) technology (Paul et al. 2001, Baer et
125 al., 2002), which utilizes true wavelength scanning to record fully resolved absorption line shapes.
126 Considering the nature of the site, flow rate is fixed to be 7 liters per minute (lpm). Ambient air
127 entering the GGA is analysed using two near infrared (NIR) distributed feedback tunable diode
128 lasers (TDL), one for a CO₂ absorption line near 1.60 μm ($\nu_0 = 6250 \text{ cm}^{-1}$) and the other to probe
129 CH₄ and H₂O absorption lines near 1.65 μm ($\nu_0 = 6060.60 \text{ cm}^{-1}$). The concentration of the gases is
130 determined by the absorption of their respective characteristic absorption lines with a high
131 sampling time of 1sec. A detailed explanation regarding the configuration, working and
132 calibration procedure performed for GGA in NRSC can be found elsewhere in Mahesh et al.,
133 (2015). In the present study we used GGA retrieve CO₂ and CH₄ data. High resolution data sets
134 are diurnally averaged and used in further analysis. Due to failure of internal central processing
135 unit (CPU) of the analyzer, data are not recorded from pre-monsoon month of 1st May to 18th June
136 during the study period.

137 3.1.2 O₃ and NO_x analyzer

138 Surface concentrations of O₃ and NO_x have been measured continuously using on-line
139 analyzers (Model No.s: 49i and 42i for O₃ and NO_x respectively), procured from Thermo
140 Scientific, USA since July 2014. The trace gases (O₃ and NO_x) sampling inlet is installed on the
141 top of a 2 m mast fixed on the roof of an 8 m high building, and ambient air flow is supplied to the
142 instruments. The inlet prevents the ingress of rain water, and is equipped with 0.5 μm filter to
143 prevent accumulation of dust within the instrument. The ozone analyzer is based on Beer-Lambert-
144 Baugher law which relates absorption of light to the concentration of species as its operating
145 principle and has an in-built calibration unit for conducting periodical span and zero checks. The
146 NO_x analyzer utilizes a molybdenum converter to convert NO₂ into NO and estimates the NO_x

147 concentration by the intensity of light emitted during the chemiluminescent reaction of NO with
148 O₃ present in the ambient air. The analyzer is integrated with zero and span calibrations which are
149 performed twice monthly.

150 Simultaneous observations of meteorological parameters are obtained from an automatic
151 weather station (AWS) installed in NRSC, Shadnagar campus as a part of Calibration and
152 Validation (CAL/VAL) project in March 2012 is equipped with nine sensors to measure fifteen
153 weather parameters. Weather parameters measured are at surface level and height of the AWS
154 mast is ~10 meters. ~~Wind speed and direction measurements are collected at the maximum height~~
155 ~~(3m) and all others are at 1–1.5m height.~~

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157 **3.2 Satellite and Model observations**

158 3.2.1 MODIS

159 Moderate-resolution Imaging Spectrometer (MODIS) is launched in December 1999 on the
160 polar-orbiting NASA-EOS Terra platform (Salomonson et al. 1989; King et al. 1992). It has 36
161 spectral channels and acquires data in 3 spatial resolutions of 250 m, 500 m, and 1 km (channels
162 8–36), covering the visible, near-infrared, shortwave infrared, and thermal-infrared bands. In the
163 present study we used monthly Normalised Difference Vegetation Index (NDVI) data obtained
164 from Terra/MODIS at 5 km spatial resolution. The NDVI value is defined as following ratio of
165 albedos (α) at different wavelengths:

$$166 \quad \text{NDVI} = \frac{\alpha_{0.86\mu\text{m}} + \alpha_{0.67\mu\text{m}}}{\alpha_{0.86\mu\text{m}} - \alpha_{0.67\mu\text{m}}} \quad (1)$$

167 NDVI values can range from -1.0 to 1.0 but typical ranges are from 0.1 to 0.7, with higher values
168 associated with greater density and greenness of plant canopies. More details of the processing
169 methods used in generating the data set can be found in James and Kalluri (1994).

170 3.2.2 COSMIS-RO

171 COSMIC (Constellation Observation System for Meteorology, Ionosphere and Climate) is a
172 GPS (Global Positioning System) radio occultation (RO) observation system (Wang et al., 2013).
173 It consists of six identical microsatellites, and was launched successfully on 14 April 2006. GPS
174 radio occultation observation has the advantage of near-global coverage, all-weather capability,
175 high vertical resolution, high accuracy and self-calibration (Yunck et al., 2000). Geophysical

176 parameters (such as, temperature and humidity profiles) have been simultaneously obtained from
177 refractivity data using one-dimensional variational (1DVAR) analysis. Further COSMIC-RO
178 profiles are used to estimate planetary boundary layer height (BLH). BLH is defined to be the
179 height at which the vertical gradient of the refractivity or water vapor partial pressure is minimum
180 (Ao et al., 2012), explained detail methodology for calculating the BLH from refractivity (N). The
181 planetary boundary layer (PBL) is part of the atmosphere closest to the Earth's surface where
182 turbulent processes often dominate the vertical redistribution of sensible heat, moisture,
183 momentum, and aerosols/pollution (AO et al., 2012).

184 3.2.3 Hysplit model

185 The general air mass pathway reaching over Shadnagar is analysed using HYSPLIT model
186 (Draxler and Rolph, 2003) [<http://ww.arl.noaa.gov/ready/hysplit4.html>]. We computed 5 day
187 isentropic model backward air mass trajectory for all study days with each trajectory starting at
188 00:60 UTC and reaching study site, (Shadnagar) at different altitudes(1 km,2 km,3km and 4 km).
189 Even though the trajectory analysis have inherent uncertainties (Stohl, 1998), they are quite useful
190 in determining long range circulation.

191 4. Results and Discussion

192

193 4.1 Seasonal variations of CO₂ and CH₄

194 Temporal variations of CO₂ and CH₄ during the study period are shown in Figure. 2a and 2b.
195 The circles indicate the daily mean, while triangular markers represent weekly averages and
196 monthly mean by square markers. Annual mean of CO₂ over study region is found to be 394 ± 2.92
197 (mean (μ) \pm standard deviation (1σ)) ppm with an observed minimum in monsoon and maximum
198 in pre-monsoon. Seasonal mean values of CO₂ observed during different seasons are 393 ± 5.60 ,
199 398 ± 7.60 , 392 ± 7.0 , and 393 ± 7.0 ppm in winter, pre-monsoon, monsoon, and post-monsoon
200 respectively. Minimum CO₂ during winter (dry season) can be due to respiratory loss of carbon
201 (Gilmanov et al., 2004; Aurela et al. 2004) as decreased temperature and solar radiation during
202 this period inhibit increases in local CO₂ assimilation (Thum et al., 2009). A steady increase in
203 CO₂ concentration is observed as season changes from winter to pre-monsoon months.
204 Enhancement in Pre-monsoon is due to higher temperature and solar radiation prevailing during
205 these months which stimulate the assimilation of CO₂ in the daytime and respiration in the night
206 (Fang et al., 2014). The enhanced soil respiration during these months also compliments the

207 increase in CO₂ concentration during this period. In addition to these natural causes, biomass
208 burning over Indian region can also have a significant effect on pre-monsoon CO₂ concentration.
209 More detailed explanation of biomass burning influence on pre-monsoon GHGs concentration is
210 discussed in section 4.6. Surface CO₂ concentration recorded a minimum during monsoon months
211 can be mainly because of enhanced photosynthesis processes with the availability of greater soil
212 moisture. A decrease in CO₂ concentration is also observed as the monsoon progress. The
213 decreases in temperature (due to cloudy and overcast conditions prevailing during these months)
214 reduce leaf and soil respiration which contributes to the enhancement of carbon uptake (Patil et
215 al., 2013; Jing et al., 2010). Further increase during post-monsoon CO₂ is associated with high
216 ecosystem productivity (Sharma et al., 2014) also an enhancement in soil microbial activity
217 (Stefanie Kirschke et al., 2013).

218 CH₄ concentration in the troposphere is principally determined by a balance between surface
219 emission and destruction by hydroxyl radicals (OH). The major sources for CH₄ in the Indian
220 region are rice, paddies, wetlands and ruminants (Schneising et al., 2009). Annual CH₄
221 concentration over study area is observed to be 1.92 ± 0.07 ppm, with a maximum (2.02 ± 0.01
222 ppm) observed in post-monsoon and minimum (1.85 ± 0.03 ppm) in monsoon. Seasonal mean
223 (average) values of CH₄ observed during different seasons are 1.93 ± 0.05 , 1.89 ± 0.05 , 1.85 ± 0.03 ,
224 and 2.02 ± 7 ppm with respectively winter, pre-monsoon, monsoon, and post-monsoon. The highest
225 concentration appears during post-monsoon and may be associated with the Kharif season
226 (Goroshiet al., 2011). Hayashida et al. (2013) reported that the seasonality of CH₄ concentration
227 over monsoon Asia is characterized by higher values in the wet season and lower values in the dry
228 season; possibly because of the effects of strong emissions from rice paddies and wetlands during
229 the wet season. Low mixing ratios of CH₄ observed during monsoon season were mainly due to
230 the reduction in atmospheric hydrocarbons because of the reduced photochemical reactions and
231 the substantial reduction in solar intensity (Abhishek Gaur et al 2014). The rate of change of CH₄
232 was found to be high during post-monsoon. Both biological and physical processes control the
233 exchange of CH₄ between rice paddy fields and the atmosphere (Nishanth et al., 2014; Goroshiet
234 al., 2011). Due to this, enhanced CH₄ observed during post-monsoon at present study area..

235 ***4.2 Influence of vegetation on GHGs.***

236 In India cropping season is classified into (i) Kharif and (ii) Rabi based on the onset of
237 monsoon. The Kharif season is from July to October during the south-west (SW) monsoon and
238 Rabi season is from October to March (Koshal Avadhesh, 2013). NDVI being one of the indicators
239 of vegetation change, monthly variations of CO₂ and CH₄ against NDVI is studied to understand
240 the impact of land use land cover on mixing ratios of CO₂ and CH₄. Monthly mean changes in
241 NDVI, CO₂ and CH₄ are shown in Figure 2c and 2d. Monthly mean of GHGs represented in this
242 analysis is calculated from daily mean in day time (10-16 LT). Analysis of the figure reveals that
243 an inverse relationship exists between NDVI and CO₂; while a positive relation is observed w.r.t
244 CH₄. Generally over this part of the country vegetation starts during the month of June with the
245 onset of SW monsoon and as vegetation increases a decrease in CO₂ concentration is observed,
246 due to enhancement in photosynthesis. Further a decline in NDVI is observed as the season
247 advances from post monsoon to winter and then to pre-monsoon, and it is associated with an
248 increase in CO₂ concentration. Similarly, the main source for CH₄ emissions are soil microbial
249 (Stefanie Kirschke et al., 2013) activity which are more active during monsoon and post monsoon
250 seasons.

251 Biomass burning (forest fire and crop residue burning) is one of the major sources of gaseous
252 pollutants such as carbon monoxide (CO₂), methane (CH₄), nitrous oxides (NO_x) and
253 hydrocarbons in the troposphere (Crutzen et al., 1990, 1985; Sharma et al., 2010). In order to study
254 the role of biomass burning on GHGs a case study is discussed. Figure 3c shows the spatial
255 distribution of MODIS derived fire counts over Indian region during 14-21 April 2014 with
256 air mass trajectories ending over study area overlaid on it at different altitudes viz. 1000 m,
257 2000 m and 4000 m respectively. Analysis of the figure shows a number of potential fire
258 locations on the north-western and south-eastern side of study location and trajectories indicate
259 its possible transport to study area. Daily mean variation of GHGs during the month of April
260 2014 (Figure 3b) indicates an enhancement in GHGs during the same period (14-21 April
261 2014). Analysis reveals that CO₂ and CH₄ have increased by ~2% and ~0.06% respectively
262 during event days with respect to monthly mean. This analysis reveals that long range / regional
263 transported biomass burning have a role in enhancement of GHGs over study site. Further to
264 understand the seasonal variation of biomass burning contribution to GHGs we analysed long
265 term (2003-2013) Fire Energetics and Emissions Research version 1.0 (FEER v1) data over study

266 area. Emission coefficient (C_e) products during biomass burning is developed from coincident
267 measurements of fire radiative power (FRP) and AOD from MODIS Aqua and Terra satellites
268 (Ichoku and Ellison, 2014). Figure 3a shows seasonal variation of CO_2 emission due to biomass
269 burning over the study site. Enhancement in CO_2 emission is seen during pre-monsoon months;
270 which also supports earlier observation (Figure 2a). This analysis reveals that biomass burning has
271 a role in pre-monsoon enhancement of CO_2 over study site. For a qualitative analysis of this long
272 range transport, we have analysed air mass trajectories ending over study site during different
273 seasons.

274 **4.3 Correlation between CO_2 and CH_4**

275 A correlation study is carried out between hourly averaged CO_2 and CH_4 during all season for
276 the entire study period. The statistical analysis for different seasons is shown in Table 2. Fang et
277 al., (2015) suggest the correlation coefficients (R_s) value higher than 0.50 indicates a similar
278 source mechanism of CO_2 and CH_4 . Also a positive correlation dominance of anthropogenic
279 emission on carbon cycle. Our study also reveals a strong positive correlation observed between
280 CO_2 and CH_4 during winter, pre-monsoon, monsoon, and post-monsoon with R equal to 0.80, 0.80,
281 0.61, and 0.72 respectively. Seasonal regression coefficients (slope) and their uncertainties (ψ_{slope} ,
282 ψ_{y-int}) are computed using Taylor (1997) which showed maximum during winter, pre-monsoon, and
283 minimum in a monsoon that figure out the hourly stability of the mixing ratios between CO_2 and
284 CH_4 . This can be due to relatively simple source/sink process of CO_2 in comparison with CH_4 .
285 Figure 4 shows the seasonal variation of $\Delta CH_4/\Delta CO_2$. Dilution effects during transport of CH_4
286 and CO_2 can be minimized to some extent by dividing the increase of CH_4 over time by the
287 respective increase in CO_2 (Worthy et al., 2009). In this study, background concentrations of
288 respective GHGs are determined as mean values of the 1.25 percentile of data for monsoon, post-
289 monsoon, pre-monsoon and winter (Pan et al., 2011; Worthy et al., 2009). Annual $\Delta CH_4/\Delta CO_2$
290 over the study region during the study period is found to be 7.1 (ppb/ppm). This low value clearly
291 indicates the dominance of CO_2 over the study region. The reported $\Delta CH_4/\Delta CO_2$ values from some
292 of the rural sites viz Canadian Arctic and Hateruma Island (China) are of the order 12.2 and ~10
293 ppb/ppm respectively (Worthy et al., 2009; Tohjima et al., 2014). Average $\Delta CH_4/\Delta CO_2$ ratio
294 during winter, pre-monsoon, monsoon and post-monsoon are 9.40, 6.40, 4.40, and 8.20 ppb
295 respectively. Monthly average, of $\Delta CH_4/\Delta CO_2$, is relatively high from late post-monsoon to

296 winter, when the biotic activity is relatively dormant (Tohjima et al., 2014). During pre-monsoon
297 decrease in $\Delta\text{CH}_4/\Delta\text{CO}_2$ ratio indicates the enhancement of CO_2 relative to that of CH_4 .

298

299 ***4.4 Diurnal variations of CO_2 and CH_4***

300 Figure 5a to 5d shows the seasonally averaged diurnal cycle of CO_2 and CH_4 over Shadnagar
301 during study period. The vertically bar represents the standard deviation from respective mean.
302 Irrespective of seasonal variation GHGs showed a similar diurnal variation, with maximum mixing
303 ratios observed during early morning (06:00 hrs) as well as early night hours (20:00 hrs) and
304 minimum during afternoon hours. However the difference observed in the maximum diurnal
305 amplitudes can be attributed to seasonal changes. The observed diurnal cycle of GHGs is closely
306 associated with diurnal variation of planetary boundary layer height (PBLH). For better
307 understanding of the diurnal behavior of CO_2/CH_4 , we used European Centre for Medium-range
308 Weather Forecasting (ECMWF) Interim Reanalysis (ERA) PBL data set which gives the data for
309 every three hours viz. 00:00, 03:00, 06:00, 09:00, 12:00, 15:00, 18:00, and 21:00 UTC with a
310 resolution of $0.25^\circ \times 0.25^\circ$ (<http://data-portal.ecmwf.int>). Figure 5a to 5d portrays the diurnal
311 evolution of CO_2/CH_4 during different season along with the evolution of Boundary Layer Height
312 (m) on secondary y axis. The morning peak arises due to combined influence of fumigation effect,
313 (Stull 1988) and morning build-up of local anthropogenic activities (household and vehicular
314 transport). Low value of GHGs as the day progress can be attributed to increased photosynthetic
315 activity during day time and destruction of stable boundary layer and residual layer due convective
316 activity. In the evening hours, surface inversion begins and form a shallow stable boundary layer
317 (Nair et al., 2007) causing the enhancement in GHGs concentration near the surface. Similar trend
318 in diurnal variation of GHGs is reported from other parts of the country (Patil et al., 2013; Mahesh
319 et al., 2014; Sharma et al., 2014; Nishanth et al., 2014).

320 ***4.5 Influence of prevailing meteorology***

321 Redistribution (both horizontal and vertical) of GHGs also plays a role in their seasonal
322 variation, as it controls transport and diffusion of pollutants from one place to another (Hassan
323 2015). A good inverse correlation between wind speed and GHGs suggest the proximity of sources
324 near measurement site, while a not so significant correlation suggests the influence of regional

325 transport (Ramachandran and Rajesh, 2007). Figure 6a and 6b shows scatter plot between GHGs
326 and wind speed during different seasons. Analysis of Figure 6 shows that there exists an inverse
327 correlation between daily mean wind speed and GHGs. Correlation coefficients (R_s) between wind
328 speed and CO_2 during pre-monsoon, monsoon, post-monsoon, and winter is 0.56, 0.32, 0.06, and
329 0.67 respectively. While for CH_4 it is found be 0.28, 0.71, 0.21, and 0.60 respectively. Negative
330 correlation indicates that the influence of local sources on GHGs, however, poor correlation
331 coefficients during different seasons suggest the role of regional/local transport (Mahesh et al
332 2014). Also an understanding of prevailing wind direction and its relationship with GHGs helps in
333 determining their probable source regions. Table 3 shows the monthly mean variation of CO_2 and
334 CH_4 with respect to different wind direction. Enhancement in CO_2 and CH_4 level over Shadnagar
335 are observed to mainly come from NW and NE while the lowest is from the S and SW. This can
336 be associated to some extent with industrial emissions located in western side of sampling site,
337 and the influence of emission and transport from nearby urban center on the NW side of the study
338 site.

339 The influence of meteorological parameters (temperature and relative humidity) on trace gases
340 is also examined. Figure 7a and 7b (top panel corresponds to CO_2 and bottom panel represents
341 CH_4) show the scatter plot of temperature versus relative humidity as a function of GHGs during
342 different seasons. Daily mean data is used instead of hourly mean data, to avoid the influence of
343 the diurnal variations on correlations. CO_2 showed a positive correlation with temperature during
344 all season except during winter. This negative correlation can be attributed to different response
345 of photosynthesis rate to different air temperature. IPCC (1990) reports that many mid-latitude
346 plants shows an optimum gross photosynthesis rate when temperature varied from of 20 to 35 °C.
347 The rate of plant respiration tends to be slow below 20°C. However, at higher temperatures, the
348 respiration rate accelerates rapidly up to a temperature at which, it equals the rate of gross
349 photosynthesis and there can be no net assimilation of carbon. While CH_4 showed a weak positive
350 correlation with temperature during pre-monsoon and post-monsoon, while a weak negative
351 correlation is observed during monsoon and winter. This could be due to the rate of chemical loss
352 reaction with OH is faster in summer and minimum in other seasons. A case study on CH_4 sink
353 mechanism has discussed in section 4.6. Seasonal variation of GHGs also showed an
354 insignificantly negative correlation with relative humidity. One of the supporting argument can be
355 in humid conditions, these stoma can fully open to increase the uptake of CO_2 without a net water

356 loss. Also, wetter soils can promote decomposition of dead plant materials, releasing natural
357 fertilizers that help plants grow (Abhishek et al., 2014).

358 Figure 8a and 8b illustrates the daily mean variation of GHGs with respect to soil moisture and
359 soil temperature (Top panel represent the seasonal variation of CO₂ w.r.t soil moisture and soil
360 temperature, while bottom panel represent the seasonal variation of CH₄ against the same
361 parameters). It's quite interesting to observe that GHGs behave differently w.r.t soil moisture
362 during different seasons. CH₄ shows a positive relationship during monsoon and post-monsoon
363 and an inverse relationship exist during pre-monsoon and winter; while a reverse relationship exist
364 for CO₂. During wet season aeration is restricted (Smith et al. 2003) hence soil respiration is
365 limited, which decrease CO₂ flux. This can be one of the factors for low values of CO₂ during
366 monsoon months, during dry months soil may act as sink of CH.

367 ***4.5.1 Influence of boundary layer height on GHGs mixing ratios***

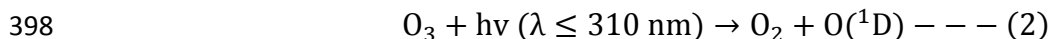
368 The planetary boundary layer is the lowest layer of the troposphere where wind speed as a
369 function of temperature plays major role in its thickness variation. It is an important parameter for
370 controlling the observed diurnal variations and potentially masking the emissions signal (Newman
371 et al., 2013). Since complete set of COSMIC RO data is not available during the study period, in
372 this analysis we have analysed RO data from July 2013 to June 2014, along with simultaneous
373 observations of GHGs. Monthly variations (Figure not show) of BLH computed from high vertical
374 resolution of COSMIC-RO data against CO₂ and CH₄ concentrations. Monthly BLH is observed
375 to be minimum (maximum) during winter and monsoon (pre monsoon) seasons and it closely
376 resembles with the air temperature pattern. The highest (lowest) BLH over study region was
377 identified 3.20 km (1.50 km). A monthly average air temperature is maximum (minimum) of 29°C
378 (20°C) during the summer (winter) months.

379 Seasonal BLH during winter, pre-monsoon, monsoon and post monsoon are 2.10 km, 3.15 km,
380 1.74 km and 2.30 km respectively. Its influence on CO₂ and CH₄ mixing ratios are shown in Figure
381 9a and 9b. X axis represents the seasonal transition i.e. monsoon to post monsoon (M-PM) etc and
382 y axis indicates seasonal difference of BLH and GHGs concentration respectively. As seasonal
383 BLH increase, mixing ratios of CO₂ (CH₄) decreased from 8.68 ppm to 5.86 ppm (110 ppb to 40
384 ppb). This effect clearly captured by seasonal diurnal averaged BLH data sets used from ECMWF-

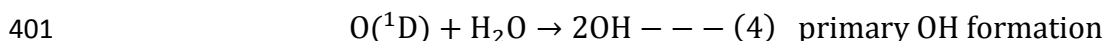
385 ERA. The amount of biosphere emissions influence on CO₂ and CH₄ can be estimated through
386 atmospheric boundary layer processes. Since the study region being a flat terrain, variations in
387 CO₂ and CH₄ were mostly influenced by BLH through convection and biosphere activities.

388 ***4.6 Methane (CH₄) sink mechanism***

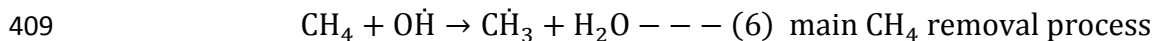
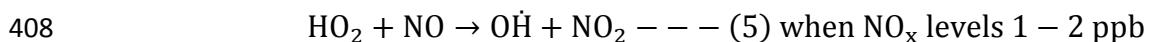
389 Methane (CH₄) is the most powerful greenhouse gas after CO₂ in the atmosphere due to its
390 strong positive radiative forcing (IPCC, AR5). Atmospheric CH₄ is mainly (70-80%) from
391 biological origin produced in anoxic environments, by anaerobic digestion of organic matter
392 (Crutzen and Zimmermann, 1991). The major CH₄ sink is oxidation by hydroxyl radicals (OH),
393 which accounts for 90 % of CH₄ sink (Vaghjiani and Ravishankara, 1991; Kim et al., 2015). OH
394 radicals are very reactive and are responsible for the oxidation of almost all gases in the
395 atmosphere. Primary source for OH radical formation in the atmosphere is photolysis of ozone
396 (O₃) and water vapor (H₂O). Eisele et al., (1997) defined primary and secondary source of OH
397 radicals in the atmosphere. Primary source of OH radical is as follows;



399 where O(^1D) is electronically excited atom



402 Removal of CH₄ is constrained by the presence of OH radicals in the atmosphere. A 1 min time
403 series analysis of CH₄, NO_x, O₃ and H₂O and associated wind vector for August 2014 to understand
404 the CH₄ chemistry is shown in Figure 10a and Figure 10b. Low NO_x (1-2 ppb) values are shown
405 in horizontal elliptical region of Figure 10a and observed corresponding low CH₄ (1.80 ppm)
406 concentrations. The low NO_x in turn produces high OH radicals in the atmosphere due to
407 conversion of HO₂ radical by NO, which removes CH₄ through oxidation process as shown below.



410 $\text{NO}_2 + \text{OH} + \text{M} \rightarrow \text{HNO}_3 + \text{M} - - - (7) \text{ if } \text{NO}_x > 2 \text{ ppb } (\text{OH} \downarrow, \text{CH}_4 \uparrow)$

411 Crutzen and Zimmermann, (1991) and Eisele et al., (1997) observed that at low NO_x (0.5-2.0 ppb)
412 levels most HO_x family radicals such as HO_2 and peroxy radicals (RO_2) react with NO to form OH
413 radicals. Therefore OH radicals are much higher in the case of low NO_x . When NO_x levels increase
414 more than 2 ppb, most of the OH radicals react with NO_2 to form nitric acid (HNO_3). In first order,
415 the levels of CH_4 in the atmosphere depend on the levels of NO_x though the production of OH
416 radicals in the atmosphere is still uncertain. Figure 10a and 10b showed high CH_4 , H_2O , O_3 and
417 NO_x during a few days in August 2014. High concentrations of CH_4 , NO_x and other gases are
418 observed in the eastern direction of study site. Very high NO_x levels above 10 ppb are observed
419 and subsequently CH_4 concentrations also increased to 2.40 ppm from 1.80 ppm. In the eastern
420 direction of study site a national highway and single line broad gauge railway network are present
421 which act as possible sources of NO_x , CH_4 and CO_2 . Increase in emissions of NO_x causes decline
422 in the levels of OH radicals and subsequently observed high CH_4 over the study region.

423

424 ***4.7 Long range circulations***

425 To understand the role of long range circulation, we separated the trajectory into 4 clusters
426 based on their pathway, namely North-East (N-E), North-West (N-W), South-East (S-E), South-
427 West (S-W). The main criterion of trajectory clustering is to minimize the variability among
428 trajectories and maximize variability among clusters. Cluster mean trajectories of air mass and
429 their percentage contribution to the total calculated for each season over the study period at 3 Km
430 altitude are depicted in Figure 11. Majority of air mass trajectories during winter (~44%), pre-
431 monsoon (~64%), monsoon (~80%) and post-monsoon (~41%) are originating from NW parts of
432 the study site. For a comprehensive analysis, percentage occurrences of cluster mean trajectories
433 of air mass over study area during different season at different altitudes are also tabulated in Table
434 4. During post-monsoon to early pre-monsoon periods which are generally the post-harvest period
435 for some of the crops agriculture residue burning which are quite common in the NW and NE
436 regions part of India (Sharma et al, 2010). Our analysis reveals that during this period majority of
437 air mass reaching the study site at different altitudes come from this part of the country.

438 5. Conclusions

439 The present study analysed the seasonal variations of atmospheric GHGs (CO₂ and CH₄) and
440 associated prevailing meteorology over Shadnagar, a suburban site of Central India during the
441 period 2014. The salient findings of the study are the following:

- 442 • Irrespective of seasons, major sources for CO₂ are soil respiration and anthropogenic
443 emissions while vegetation acts as a main sink. Whereas the major source and sink for
444 CH₄ are vegetation and presence of hydroxyl (OH) radicals. In addition, boundary layer
445 dynamics and long range transport also plays a vital role on GHGs mixing ratios.
- 446 • The annual mean of CO₂ and CH₄ over the study region are found to be 394±2.92 ppm
447 and 1.92±0.07 ppm ($\mu\pm 1\sigma$) respectively. CO₂ and CH₄ show a significant seasonal
448 variation during the study period. Maximum (Minimum) CO₂ is observed during Pre-
449 monsoon (Monsoon), while CH₄ recorded maximum during post-monsoon and minimum
450 in monsoon. Seasonal analysis of FEER data also showed maximum emission of CO₂
451 due to biomass burning during pre-monsoon months which indicates the influence of
452 biomass burning on local emissions.
- 453 • CO₂ and CH₄ showed consistent diurnal behavior in spite of their significant seasonal
454 variations, with an observed morning (06:00 IST) maxima, followed by afternoon
455 minima (14:00 IST) and enhancing in the late evening (~22:00 IST).
- 456 • Correlation coefficient (Rs) between wind speed and CO₂ during pre-monsoon, monsoon,
457 post-monsoon and winter is 0.56, 0.32, 0.06 and 0.67 respectively. While for CH₄ it is
458 found be 0.28, 0.71, 0.21, and 0.60 respectively. Negative correlation indicates that the
459 influence of local sources on GHGs, however, poor correlation coefficients during
460 different seasons suggest the role of regional/local transport.
- 461 • CO₂ showed a positive correlation with temperature during all seasons except during
462 winter. Whereas CH₄ showed a weak positive correlation with temperature during pre-
463 monsoon and post-monsoon, while showing a weak negative correlation during monsoon
464 and winter.
- 465 • CO₂ and CH₄ showed a strong positive correlation during winter, pre-monsoon, monsoon
466 and post-monsoon with Rs equal to 0.80, 0.80, 0.61 and 0.72 respectively. This clearly
467 indicates common anthropogenic sources for these gases.

468

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Table 1 Data used

Sensor	Period	Parameter	resolution	Source
GGA-24EP	Jan-2014 to Dec 2014	CO ₂ ,CH ₄ and H ₂ O	1 Hz time	ASL,NRSC
42i-NO-NO ₂ -NO _x	Jul-2014 to Sep-2014	NO _x (=NO+N O ₂)	1 min time	ASL,NRSC
49i-O ₃	Jul-2014 to Sep-2014	O ₃	1 min time	ASL,NRSC
AWS	Jan-2014 to Dec-2014	WS,WD,AT,R H	60 min time	NRSC
Terra/MODIS	Jan-2014 to Dec-2014	NDVI	5 Km horizontal	http://ladsweb.nascom.nasa.gov/data/search.html
COSMIC-1DVAR	Jul-2013 to Jun-2014	Refractivity (N)	0.1 Km vertical	
HYSPLIT	Jan-2014 to Dec-2014	Backward trajectory	5 day isentropic model (1km to 4 km)	http://ww.arl.noaa.gov/ready/hysplit4.html
FEER v1	Jan-2013 to Dec-2013	fire radiative power (FRP)		http://ladsweb.nascom.nasa.gov/data/search.html

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709 **Table 2** Statistical correlation between CO₂ and CH₄

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S.No	Seasons	Correlation coefficient (R)	Slope $\left(\frac{Y_{CH_4} (ppm)}{X_{CO_2} (ppm)}\right)$	Ψ_{slope} (ppm)	Ψ_{y-int} (ppm)
1	Monsoon (JJAS)	0.61	0.005	0.00015	1.91
2	Post-monsoon (OND)	0.72	0.0065	0.00014	1.52
3	Winter (JF)	0.80	0.0085	0.00018	9.13
4	Pre-monsoon (MAM)	0.80	0.0059	0.00021	2.73

721 **Table 3** Seasonal amplitudes of CO₂ and CH₄ over study region arriving from different directions

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Wind Direction	Winter $\frac{CO_2}{CH_4}$ (ppm)	Pre-monsoon $\frac{CO_2}{CH_4}$ (ppm)	Monsoon $\frac{CO_2}{CH_4}$ (ppm)	Post-monsoon $\frac{CO_2}{CH_4}$ (ppm)
0-45	399.85/1.98	410.37/1.94	400.72/1.91	395.13/2.02
45-90	391.66/1.94	399.59/1.89	388.82/1.91	390.23/1.98
90-135	391.57/1.93	397.79/1.87	388.99/1.87	389.06/1.97
135-180	389.34/1.89	393.87/1.85	391.81/1.86	387.69/1.97
180-225	391.14/1.89	396.75/1.85	390.28/1.82	392.30/2.02
225-270	389.13/1.88	394.81/1.86	390.26/1.82	384.40/1.94
270-315	388.68/1.87	398.68/1.89	389.58/1.82	384.99/1.93
315-360	390.87/1.91	401.17/1.89	387.58/1.83	389.32/1.98

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724 **Table 4** Cluster analysis of air mass trajectories reaching Shadnagar at various heights during different
725 seasons

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Seasonal Backward trajectory (%)	NW				NE				SE				SW			
	1 km	2 km	3 km	4 km	1 km	2 km	3 km	4 km	1 km	2 km	3 km	4 km	1 km	2 km	3 km	4 km
Winter	54	32	2	0	32	24	44	52	10	25	11	7	4	19	42	41
Pre-monsoon	24	9	8	1	26	31	64	78	36	46	2	10	14	14	26	11
Monsoon	0	1	7	19	12	34	80	70	4	4	4	6	84	61	9	5
Post- monsoon	42	15	11	14	47	53	41	49	8	30	32	26	3	2	16	11

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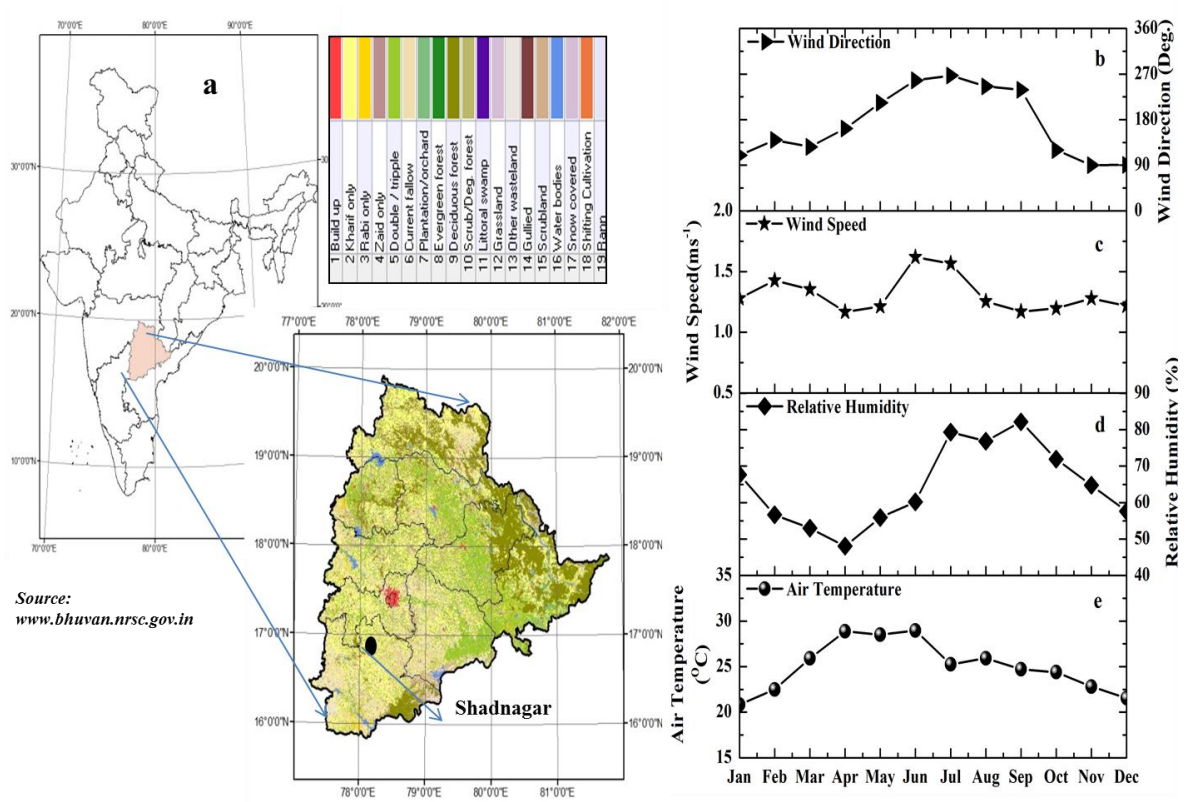
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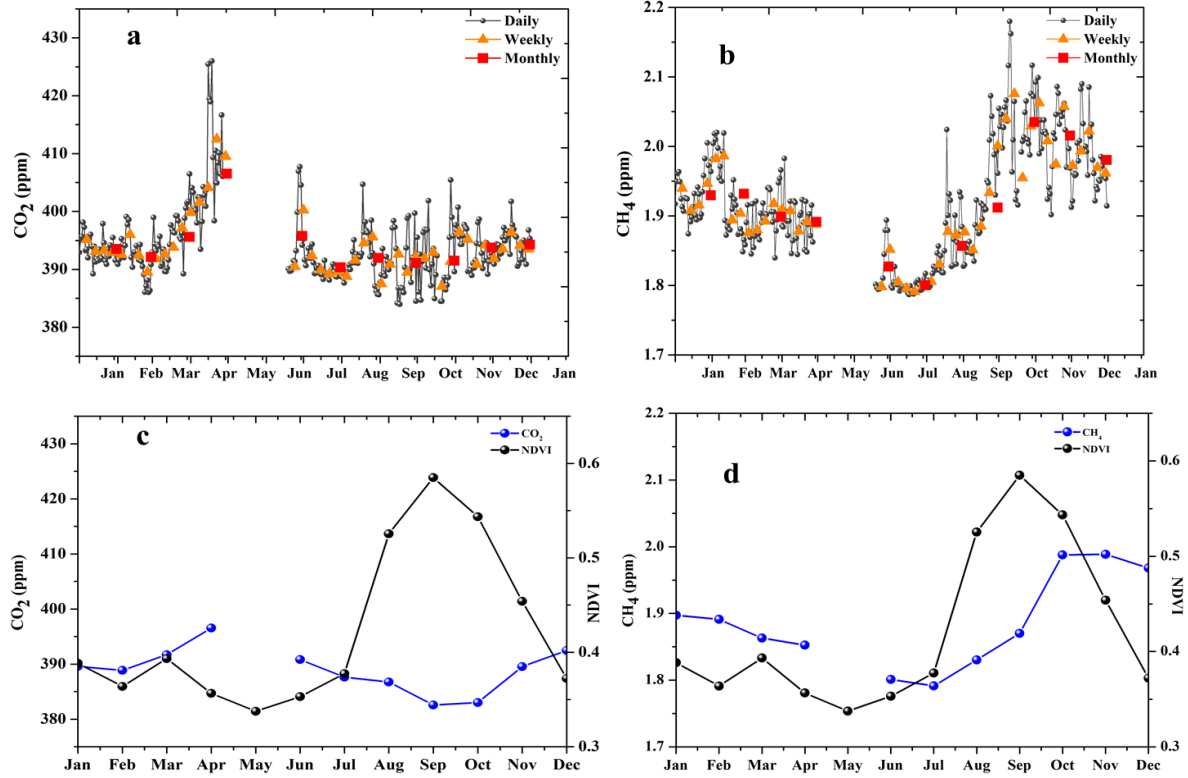
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741 **Figure 1** a) Schematic representation of study area; b-e) Seasonal variation of prevailing meteorological
742 conditions during 2014

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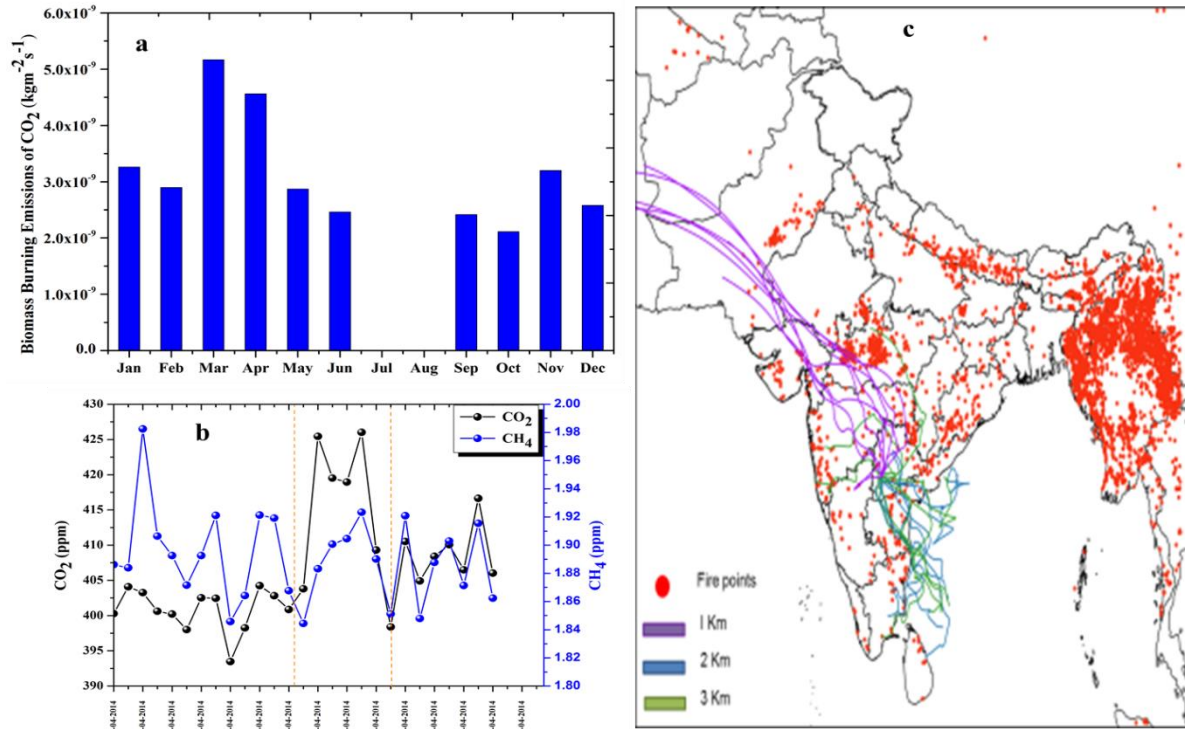
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753 **Figure 2** a-b) Temporal variations of CO₂ and CH₄; c-d) Seasonal variations of CO₂ and CH₄ in conjunction
 754 with NDVI (Normalized Difference Vegetation Index) during 2014

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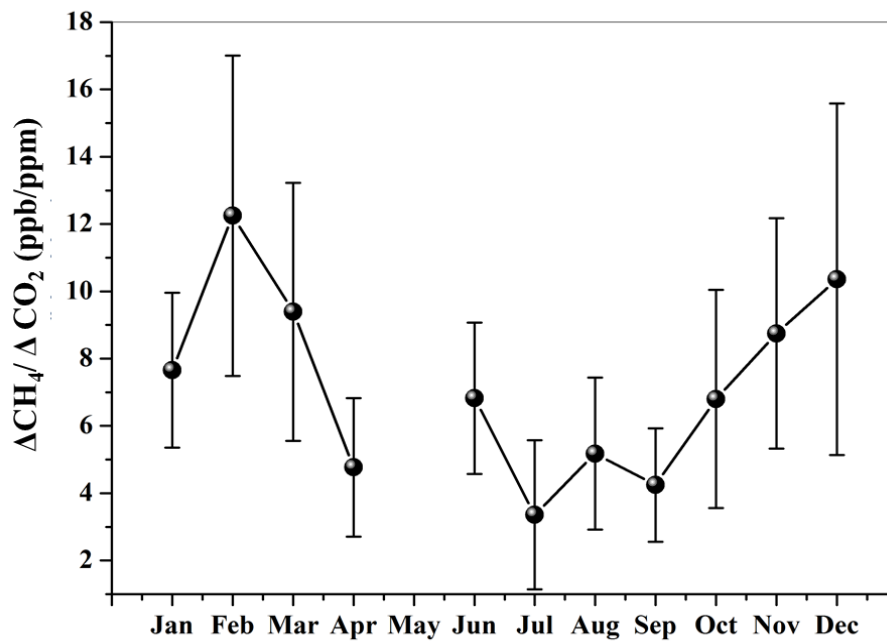
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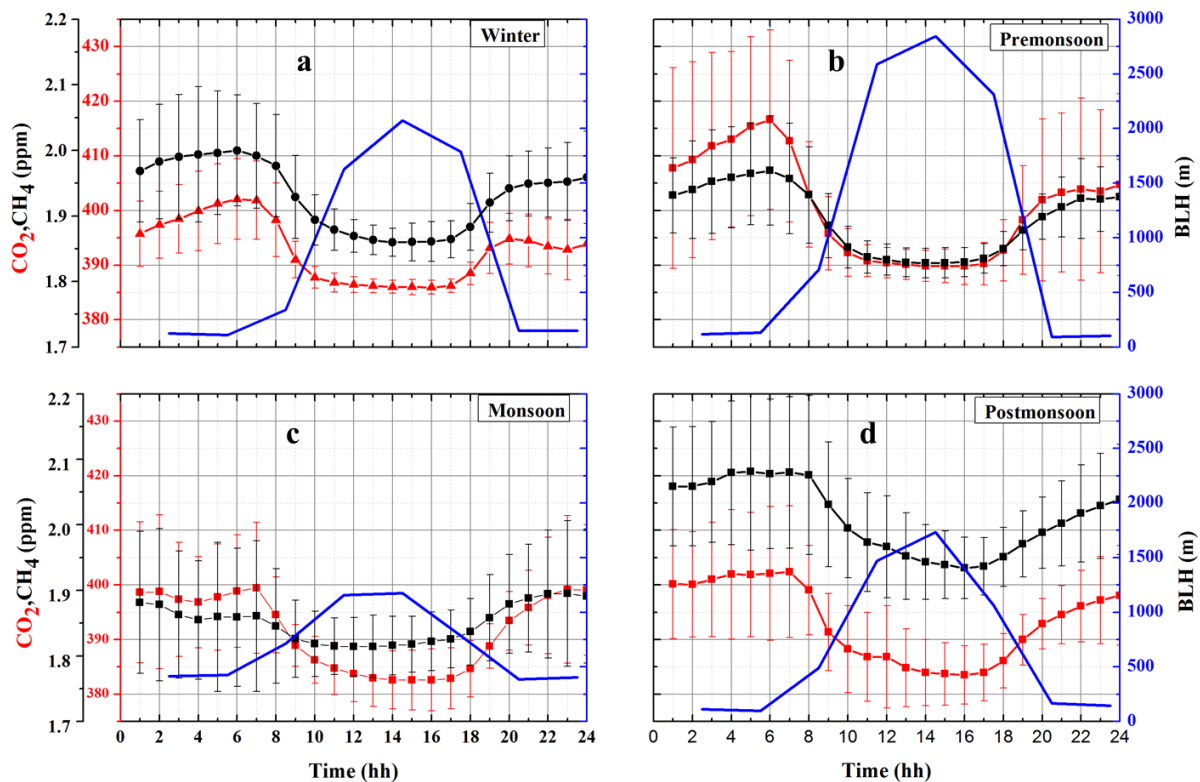
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759 **Figure 3** a) Long term analysis of CO₂ biomass burning emissions over study region b) Biomass signatures
 760 on CO₂/CH₄ during 14-21 April 2014, a case study c) Spatial distribution of MODIS derived fire counts
 761 over Indian region during 14-21 April 2014.

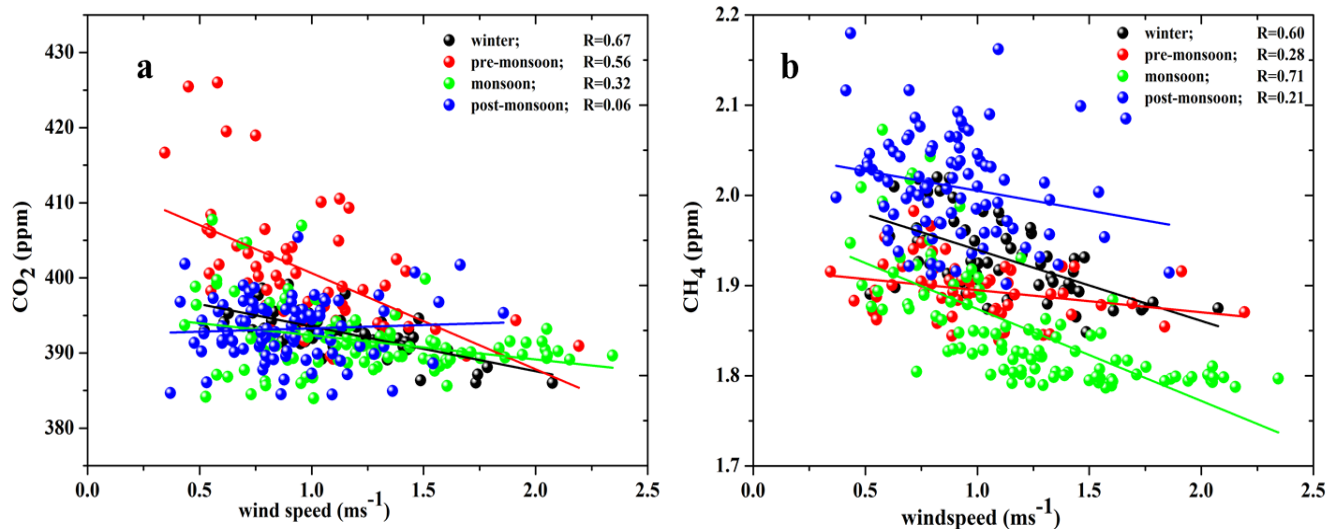


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776 **Figure 4** Monthly variation of $\Delta\text{CH}_4/\Delta\text{CO}_2$ during study period



777 **Figure 5** a-d) Seasonal variations of diurnal averaged CO_2/CH_4 against boundary layer height during 2014

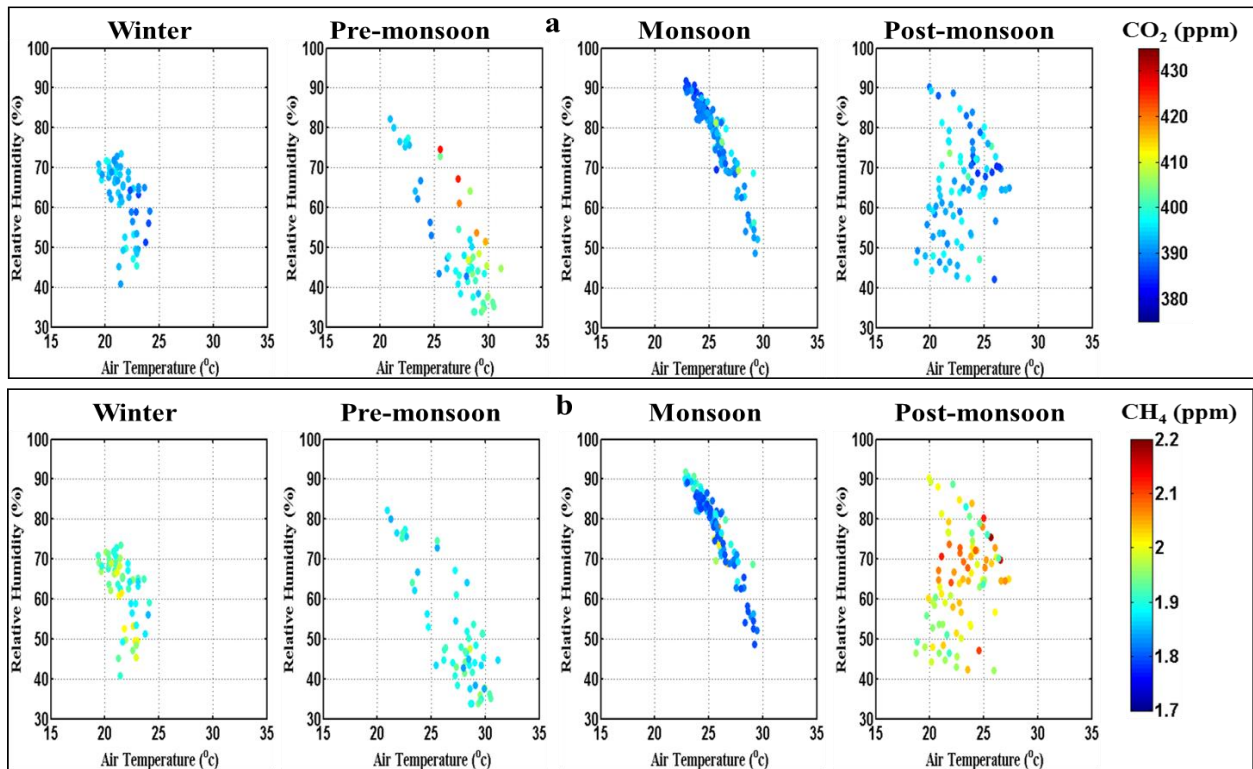


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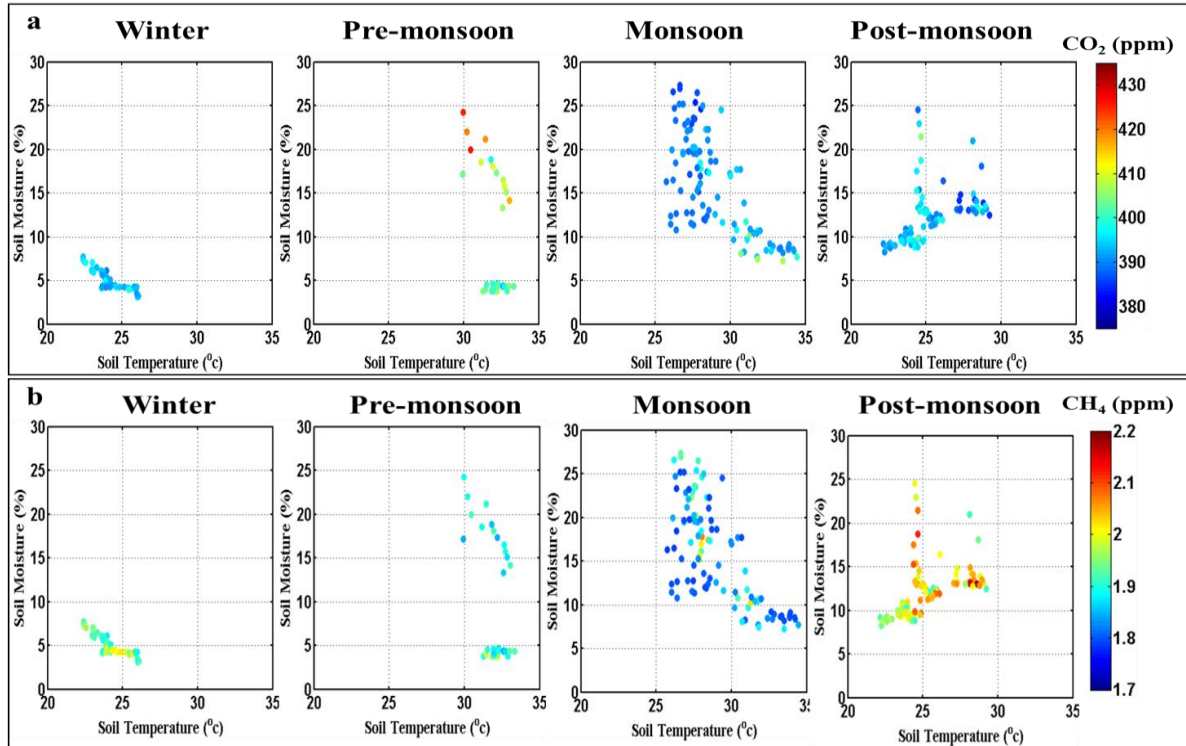
779 **Figure 6** a-b) Daily mean scatterplot between wind speed and GHGs (CO_2 and CH_4).

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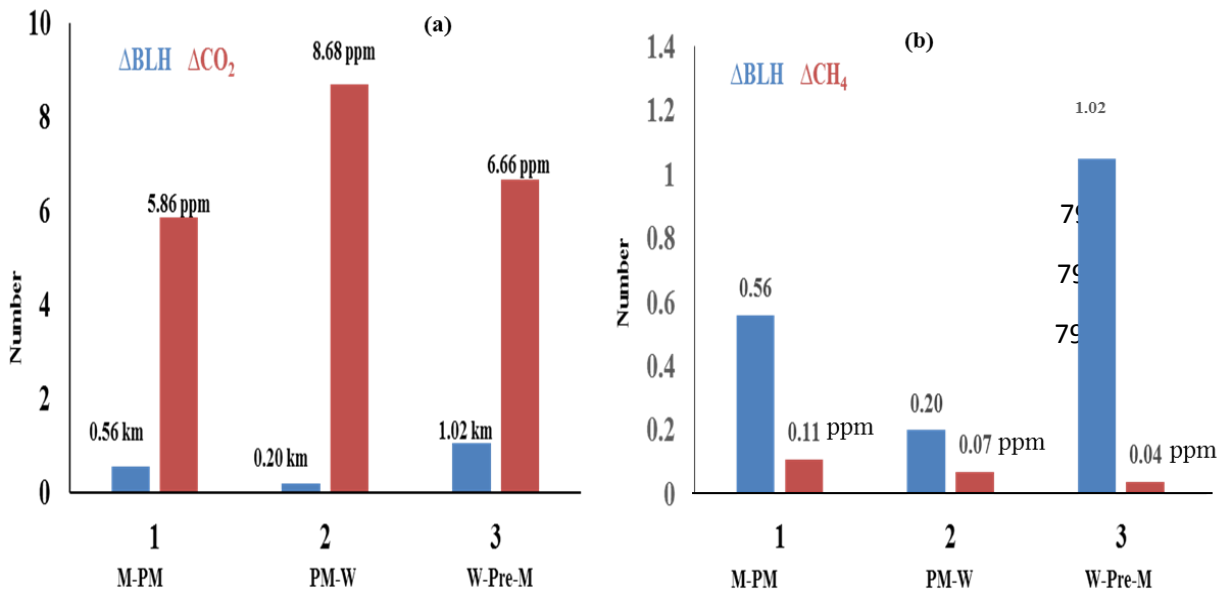
783 Figure 7 a-b) Daily mean seasonal variation of CO_2 and CH_4 as function of humidity and air temperature
784 during 2014



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786 Figure 8 a-b) Daily mean seasonal variation of CO₂ and CH₄ as function of soil temperature and soil
 787 moisture during 2014

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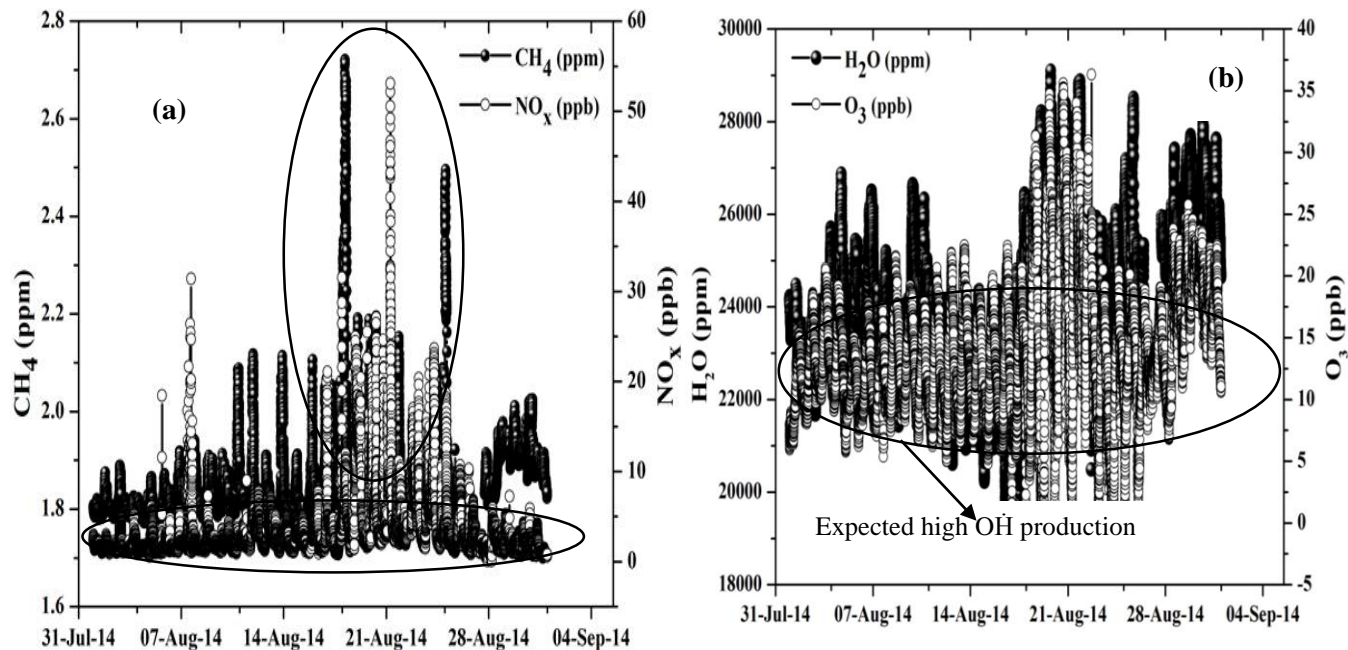
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796 **Figure 9** Seasonal difference in BLH against respective change in a) CO₂ and b) CH₄

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799 **Figure 10** Time series analysis of a) CH₄ vs. NO_x, b) H₂O vs. O₃

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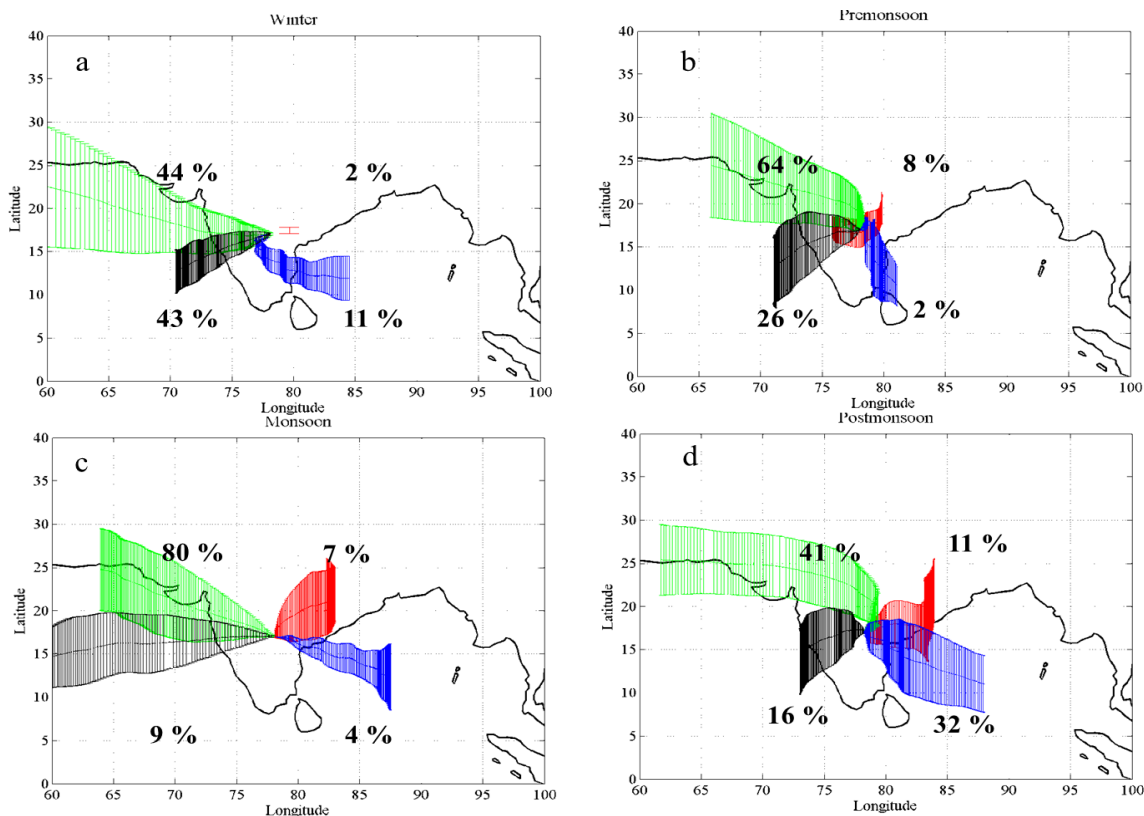
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814 **Figure 11** a-d) Long range circulation of air mass trajectories ending over Shadnagar at 3 km during
815 winter, pre-monsoon, monsoon and post-monsoon