



## RESEARCH PAPER

## OPEN ACCESS

**Geo-spatial assessment of temporal variation of greenhouse gases emission using GIS in Jahangirnagar University, Bangladesh****Mohammad Amzad Hosain, Md. Bodrud-Doza, Fahad Ahmed\*, Md. Shawon Zoarddar, Mohammad Amir Hossain Bhuiyan***Fahad Ahmed, Department of Environmental Sciences, Jahangirnagar University, Dhaka, 1342, Bangladesh***Key words:** Greenhouse gases (GHGs), GIS, Temporal variation, Jahangirnagar University.

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**Abstract**

To assess the present scenario of GHGs (CO<sub>2</sub> and CH<sub>4</sub>) emission and its distribution, day time emissions of CO<sub>2</sub> and CH<sub>4</sub> in August 2013 and 2014 in Jahangirnagar University area were studied using statistical analysis and GIS interpretation. In 2013, maximum value of CO<sub>2</sub> was 500 ppm which was found in Bishmile bio gas plant, Prantic gate and Dairy gate characterized by solid waste dumping site and high way road areas. Similarly in 2014, maximum value of CO<sub>2</sub> was 600 ppm which was found around the high way road site area. In 2013 and 2014 maximum value of CH<sub>4</sub> was 3500 ppm and 4000ppm respectively which was found in Bishmile situated in the Northern site of study area. These sites are characterized by solid waste dumping site and biogas plant area. The results of the geospatial calculations show that in 2013, 43.355% of total area covers 357.143 to 385.714 ppm concentration of CO<sub>2</sub> while in 2014, 45.896% of the total area covers 385.714 to 428.571 ppm concentration of CO<sub>2</sub>. Apart this in 2013, 38.604% of total area covers 621.418 to 1042.18ppm concentration of CH<sub>4</sub> while in 2014, 39.168 % of the total area covers 856.583 to 1434.818 ppm concentration of CH<sub>4</sub>. From the spatial and temporal analysis it can be said that concentration of CO<sub>2</sub> and CH<sub>4</sub> in this area in a rising trend. This study would also implicitly provide a mechanism to monitor potential sources of CO<sub>2</sub> and CH<sub>4</sub> as well as their adverse contribution to enhance GHGs emissions.

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## Introduction

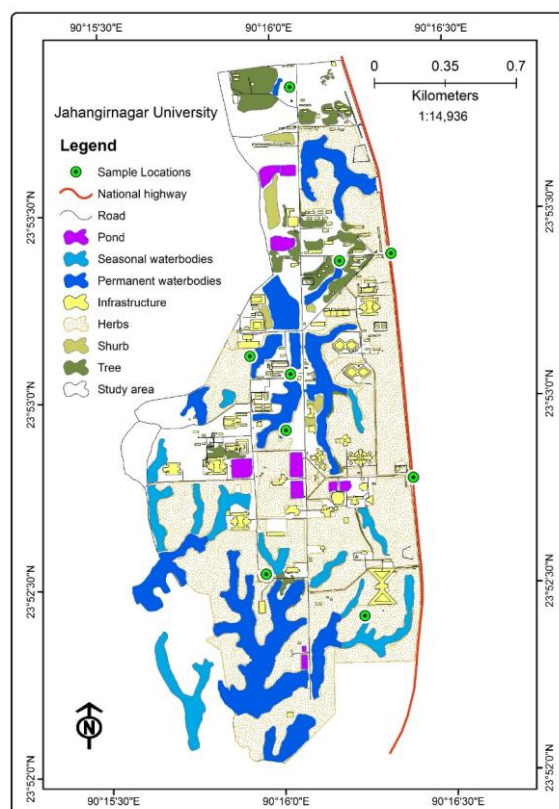
Greenhouse gases (GHGs) have been recognized to be responsible for global warming and climate change (IPCC, 2007b). Emissions of GHGs such as carbon dioxide (CO<sub>2</sub>), nitrous oxide (N<sub>2</sub>O) and methane (CH<sub>4</sub>) are ever increasing and other gases such as oxides of nitrogen (NO<sub>x</sub>), carbon monoxide (CO), and sulphur dioxide (SO<sub>2</sub>) often accompany the release of these GHGs (Sookun *et al.*, 2014). Carbon dioxide (CO<sub>2</sub>) is the primary greenhouse gas emitted through human activities. Carbon dioxide (CO<sub>2</sub>) emissions are produced in our daily lives through burning fossil fuels to meet essential needs such as electricity, heating, and transportation. As a major component of greenhouse gas emissions, increased CO<sub>2</sub> emissions due to anthropogenic activities have been suggested to contribute to global warming. Carbon dioxide (CO<sub>2</sub>) has significantly increased over recent centuries with atmospheric concentrations rising by 31% since 1750, with fossil fuel burning accounting for approximately three quarters of the increased emissions (IPCC, 2001). Methane, a greenhouse gas, has 23 times the global warming potential of carbon dioxide over a 100-year time horizon (IPCC, 2001). Methane (CH<sub>4</sub>) is a radiatively active trace gases whose abundance in the atmosphere has more than doubled during the past several hundred years and continues to rise (Cicerone and Oremland, 1988; Dlugokencky *et al.*, 1998; Shipham *et al.*, 1998). This increase is correlated with increasing human population (Rasmussen and Khalil, 1984). CH<sub>4</sub> contributes to global warming as a result of its ability to trap heat, 21 times more effectively than carbon dioxide over a 100-year period (IPCC, 2001). Also, the oxidizing capacity of the atmosphere is diminished by increasing CH<sub>4</sub> through reaction with hydroxyl radicals. The short lifetime of CH<sub>4</sub> in the atmosphere, about 9 years compared with 120 years for carbon dioxide, coupled with its potency makes it an ideal candidate for emissions reduction (WMO, 1998; IPCC, 2001).

Jahangirnagar University is the only residential university in Bangladesh close to the capital city

Dhaka. So far, no efforts have been made to monitor ambient concentration of CO<sub>2</sub> and CH<sub>4</sub> in Jahangirnagar University and its surroundings. The present study focuses on direct measurements and to develop a general profile of temporal variations in the ambient concentration of CO<sub>2</sub> and CH<sub>4</sub> in this area.

## Material and methods

2.1 Study area: Jahangirnagar University situated in Savar Upazila, 32 kilometers away from the capital city Dhaka (Fig. 1). Total area is 697.56 acres (2.8 km<sup>2</sup>) and about 17,000 people staying here. This area encompasses with approximately 25% seasonal and permanent water bodies and about 20% vegetation cover. In the Northern site of this university a landfill site and a biogas plant is situated.



**Fig. 1.** Map of the Study Area.

### Data collection

Concentration of CO<sub>2</sub> and CH<sub>4</sub> was measured by Landfill Gas analyzer LFG20 from Jahangirnagar University area in August of 2013 and 2014 consequently. Data were measured around 11.00 am

to 3.00 pm (GMT+6.00). Landfill Gas analyzer LFG20 was settled at 3.0 m height from ground level during operation (Sahay and Ghosh, 2013). Data was collected from 9 selective points (Table 1).

**Table 1.** Name of the sampling sites with GPS locations.

Site ID	Name of the site	Latitude	Longitude
S1	Bishmile bio gas plant	23.89722	90.2675
S2	Prantic gate	23.88972	90.27222
S3	Prantic mosque	23.88944	90.26972
S4	Al-Beruni hall	23.88528	90.26528
S5	Fazilatunnesa Hall	23.88444	90.26722
S6	New Register building	23.88194	90.26694
S7	Gymnasium	23.87556	90.26583
S8	Dairy gate	23.87972	90.27306
S9	MH Hall	23.87361	90.27056

*Data preparation*

GPS location of the monitoring sites was measured using Garmin GPS. Collected percentage data of CO<sub>2</sub> and CH<sub>4</sub>, then converted them into ppm using the following equations.

$$X \text{ (ppm)} = 10000 \times X \text{ (\%)}$$

*GIS based methods*

ArcGIS-10.2 is used for the analysis of temporal variation of GHGs emission in the study area. Geo-databases were developed and Geo-referenced with GCS\_WGS\_1984. The features were projected using WGS\_1984\_UTM\_Zone\_45N (Universal Transverse Mercator).

*Spatial Analysis*

IDW (Inverse Distance Weighted) is an interpolation technique in which interpolated values are made based on measurement values at nearby locations weighted only by distance from the interpolation location. This technique does not assume any type of spatial relationship, except the basic assumption that nearby points ought to be more closely related to one another than more distant points (Davis, 1986; Maguire *et al.*, 1991). The only complication is the resulting contour is a function of sampling density.

The data points define the interpolation, the resulting contours are fully representative of the data. This is not always true of kriging and other contour models (Gerlach *et al.*, 2001).

Spokas *et al.* (2003) stated the formula for IDW which used for spatial distribution of CO<sub>2</sub> and CH<sub>4</sub> calculation by the way

$$Z_{est}^j = \frac{\sum(\frac{z_i}{(h_{ij} + S)^2})}{\sum(\frac{1}{(h_{ij} + S)^2})}$$

Where,  $Z_{est}^j$  is estimated value for location j,  $z_i$  is measured sample value at point i,  $h_{ij}$  is distance between  $Z_{est}^j$  and  $z_i$ ; and  $S$  is smoothing factor (0 was used).

**Results and discussion**

Results include the general findings of day time emissions of CO<sub>2</sub> and CH<sub>4</sub> in August 2013 and 2014 in Jahangirnagar University. Spatial analysis is carried out for the distribution which represent the temporal variation of the emissions of CO<sub>2</sub> and CH<sub>4</sub> in the study area.

*Statistical Analysis*

In 2013, maximum value of CO<sub>2</sub> was 500 ppm, which was found in several sites (values and site names are given in Table-1 and Table-2). These sites are characterized by solid waste dumping site and high way road sites. The minimum and average value of CO<sub>2</sub> in 2013 were 300 ppm and 388.889 ppm respectively (Table-2). In 2014, maximum value of CO<sub>2</sub> was 600 ppm which was found around the high way road site area. The minimum and average value of CO<sub>2</sub> in 2014 were 300 ppm and 422.222 ppm respectively (Table-2).

Comparing with the measured values of CO<sub>2</sub> in 2013 and 2014 shows that in some locations the concentration of CO<sub>2</sub> remains same and in a few locations CO<sub>2</sub> concentration increased (Fig. 2). Average concentration of CO<sub>2</sub> is increased in 2014 comparing with the concentration of 2013 (Fig. 3). In

2013, maximum value of CH<sub>4</sub> was 3500 ppm which was found in Bishmile, situated in the Northern site of study area. These sites are characterized by solid waste dumping site and biogas plant area. 2300 ppm and 1700 ppm concentration of CH<sub>4</sub> were found in some small scale waste dumping sites (Table 2). The

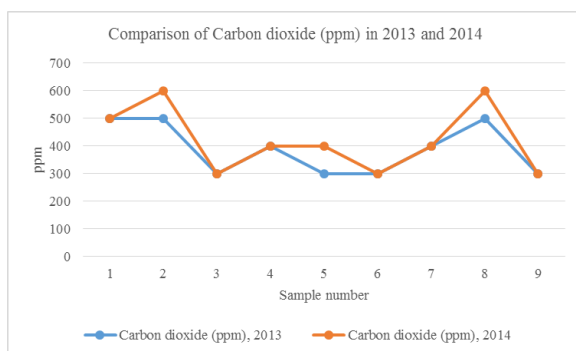
minimum and average value of CH<sub>4</sub> in 2013 were 100 ppm and 1077.778 ppm respectively (Table 3). In 2014, maximum value of CH<sub>4</sub> was 4000 ppm which was found in the similar region. The minimum and average value of CH<sub>4</sub> in 2014 were 100 ppm and 1144.444 ppm respectively (Table 3).

**Table 2.** Daytime measured concentration of CO<sub>2</sub> and CH<sub>4</sub> in the study area (2013 and 2014).

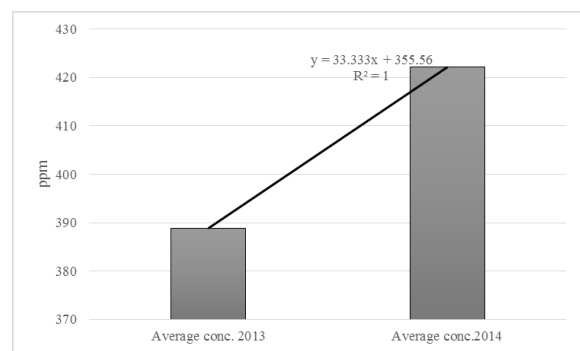
Site ID	Name of the site	2013				2014				Source Characteristics
		CO <sub>2</sub> (%)	CH <sub>4</sub> (%)	CO <sub>2</sub> (ppm)	CH <sub>4</sub> (ppm)	CO <sub>2</sub> (%)	CH <sub>4</sub> (%)	CO <sub>2</sub> (ppm)	CH <sub>4</sub> (ppm)	
S1	Bishmile bio gas plant	0.05	0.35	500	3500	0.05	0.4	500	4000	Solid Waste Dumping site
S2	Prantic gate	0.05	0.01	500	100	0.06	0.01	600	100	Bus Stand
S3	Prantic mosque	0.03	0.02	300	200	0.03	0.02	300	200	Lake
S4	Al-Beruni hall	0.04	0.23	400	2300	0.04	0.2	400	2000	Lake and small waste dumping site
S5	Fazilatunnesa Hall	0.03	0.08	300	800	0.04	0.09	400	900	Lake
S6	New Register building	0.03	0.07	300	700	0.03	0.08	300	800	Lake
S7	Gymnasium	0.04	0.02	400	200	0.04	0.01	400	100	Irrigated Land
S8	Dairy gate	0.05	0.02	500	200	0.06	0.02	600	200	Bus Stand
S9	MH Hall	0.03	0.17	300	1700	0.03	0.2	300	2000	Irrigated Land and small waste dumping site

**Table 3.** Descriptive statistics of GHGs (CO<sub>2</sub> and CH<sub>4</sub>) emission in the study area.

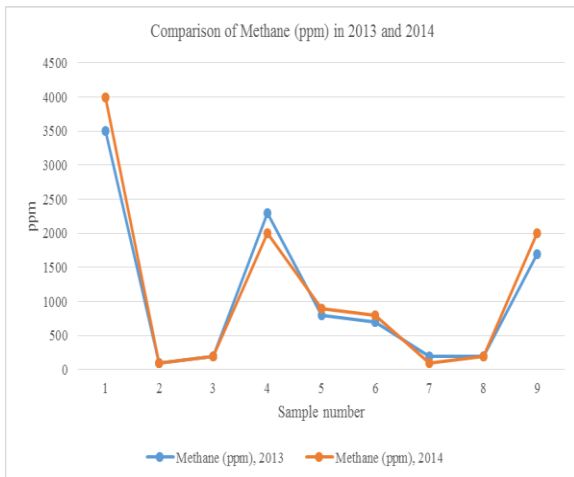
GHGs	2013					2014				
	Min	Max	Range	Mean	Std. Deviation	Min	Max	Range	Mean	Std. Deviation
CO <sub>2</sub> (ppm)	300	500	300-500	388.889	92.796	300	600	300-600	422.222	120.185
CH <sub>4</sub> (ppm)	100	3500	100-3500	1077.778	1185.093	100	4000	100-4000	1144.444	1311.593



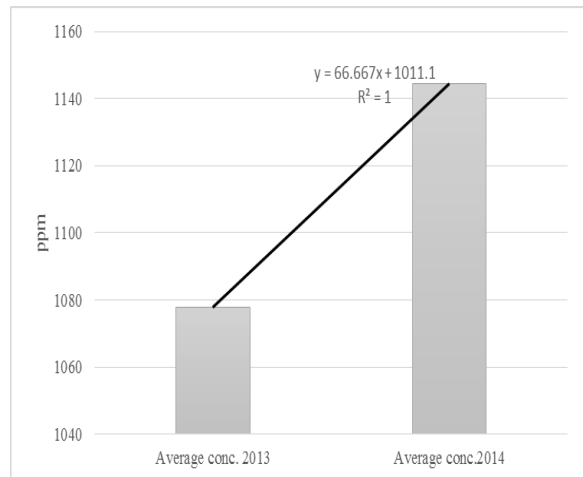
**Fig. 2.** Comparison of Carbon dioxide (ppm) in 2013 and 2014 with respective to their sampling number.



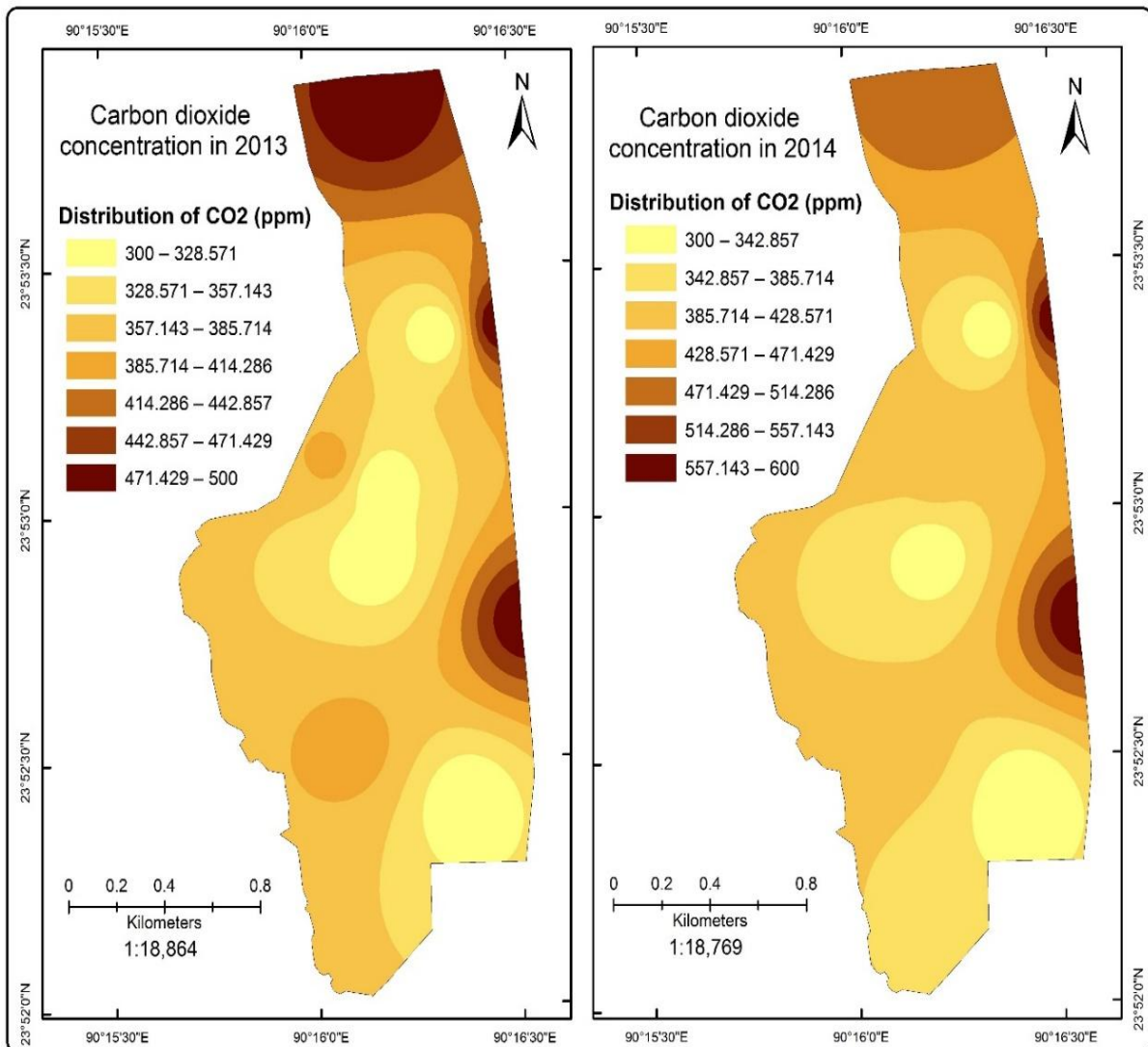
**Fig. 3.** Rising trend of Average Carbon dioxide concentration comparison with year (2013 and 2014).



**Fig.4.** Comparison of Methane (ppm) in 2013 and 2014 with respective to their sampling number.



**Fig. 5.** Rising trend of Average Methane concentration comparison with year (2013 and 2014).



**Fig. 6.** Spatial distribution of Carbon dioxide concentration in 2013 and 2014.

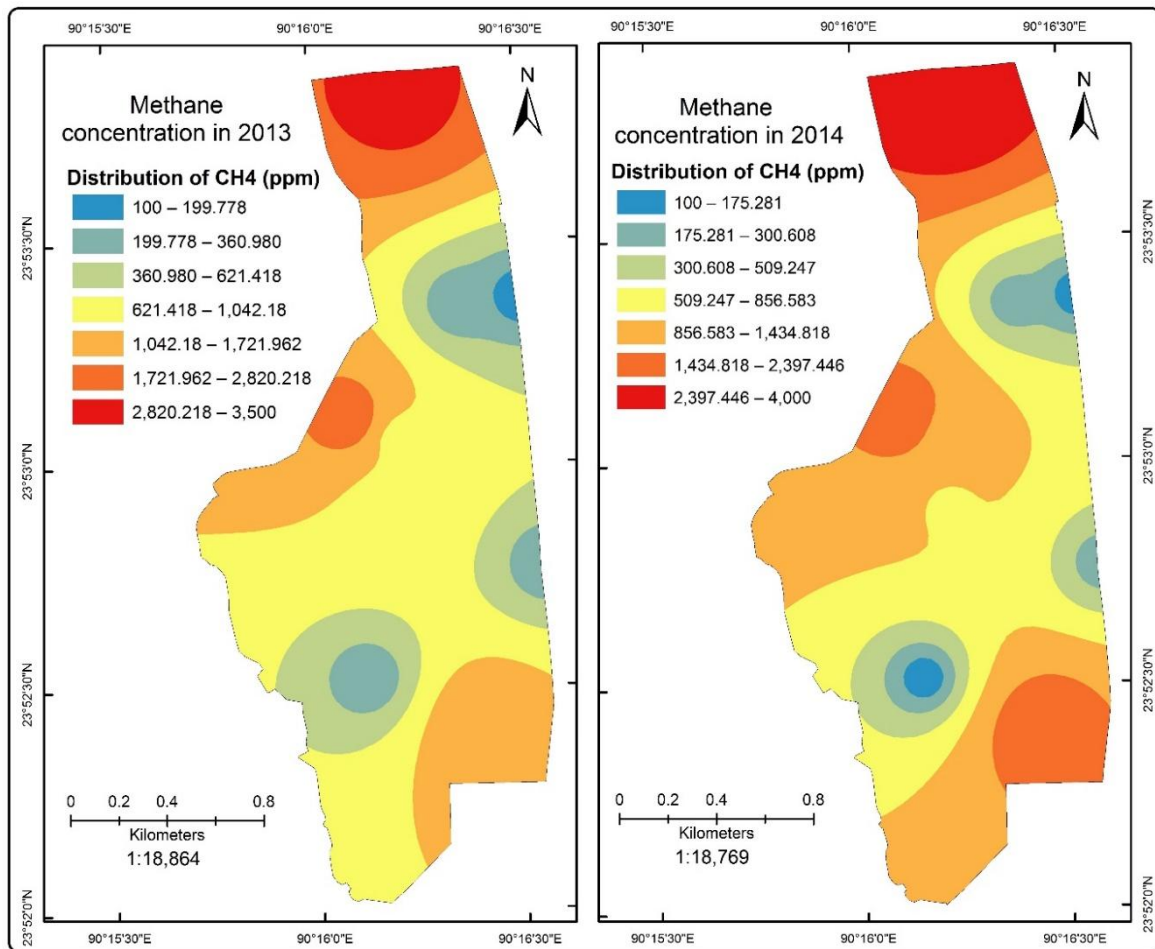


Fig. 7. Spatial distribution of Methane concentration in 2013 and 2014.

Comparing with the measured values of CH<sub>4</sub> in 2013 and 2014 shows that in few locations the concentration of CH<sub>4</sub> remains same and in some locations CH<sub>4</sub> concentration increased (Fig. 4). The average concentration of CH<sub>4</sub> is increased in 2014 compared with the concentration of 2013. Fig. 5 represents the rising trend of CH<sub>4</sub> with respect to the time in the study area.

*Geo-spatial Analysis*

Spatial Analysis is the process of examining the locations, attributes, and relationships of features in spatial data through overlay and other analytical techniques in order to address a question or gain useful knowledge (Wade and Sommer, 2006). Spatial analysis of the CO<sub>2</sub> emission in the study area provides an interpretation of the distribution of CO<sub>2</sub>. From the observation of CO<sub>2</sub> distribution in the Bishmile landfill site area and surroundings of

Dhaka-Aricha highway areas are under risk for high CO<sub>2</sub> emission. From the temporal analysis it is found that the concentration of CO<sub>2</sub> in this area in a rising trend (Fig. 2).

The results of the geospatial calculations show that in 2013, 43.355% of total area covers 357.143 to 385.714 ppm concentration of CO<sub>2</sub> while 385.714 to 500 ppm concentration of CO<sub>2</sub> covering 33.825% of the total area. In 2014, 45.896% of the total area covers 385.714 to 428.571 ppm concentration of CO<sub>2</sub> while 28.075% of the total area covering 428.571 to 600 ppm concentration of CO<sub>2</sub> (Table-4). This confirms the qualitative use as reported by Borjesson *et al.*, (2000).

Gratani and Varone (2005) reported near-surface CO<sub>2</sub> concentration measured at different sites in Rome from 1995 to 2004 with intervals. The concentrations

varied from 414 to 505 ppm at city center and showed strong correlation with the traffic density. The problem of contribution from vehicular exhaust affecting the measurements was reduced in our study as the monitoring sites were selected away from such streets. Javoy (2003), near surface CO<sub>2</sub> concentration measured in Paris, France reached as high as 950 ppm as monitoring was done close to streets with

heavy traffic loads, while outside the city, it averaged 415 ppm. Though Jahangirnagar University is a residential area, but the CO<sub>2</sub> emission value is close to the urban area with high traffic density similar value of Rome and Paris city. About 9000 vehicles are passing everyday along with Dhaka Aricha highway. High traffic density is the main offender for the elevated CO<sub>2</sub> emission.

**Table 4.** Results of IDW analysis of CO<sub>2</sub>.

Classes	CO <sub>2</sub> concentration (ppm) in 2013			CO <sub>2</sub> concentration (ppm) in 2014		
	Min Value	Max Value	Area Percentage	Min Value	Max Value	Area Percentage
1	300	328.571	6.605	300	342.857	5.537
2	328.571	357.143	16.215	342.857	385.714	20.492
3	357.143	385.714	43.355	385.714	428.571	45.896
4	385.714	414.286	12.767	428.571	471.429	16.582
5	414.286	442.857	10.531	471.429	514.286	8.212
6	442.857	471.429	5.934	514.286	557.143	1.847
7	471.429	500	4.593	557.143	600	1.434

**Table 5.** Results of IDW analysis of CH<sub>4</sub>.

Classes	CH <sub>4</sub> concentration (ppm) in 2013			CH <sub>4</sub> concentration (ppm) in 2014		
	Min Value	Max Value	Area Percentage	Min Value	Max Value	Area Percentage
1	100	199.778	0.714	100	175.281	0.875
2	199.778	360.98	4.797	175.281	300.608	3.336
3	360.98	621.418	10.009	300.608	509.247	6.244
4	621.418	1042.18	38.604	509.247	856.583	25.602
5	1042.18	1721.962	32.272	856.583	1434.818	39.168
6	1721.962	2820.218	10.436	1434.818	2397.446	17.476
7	2820.218	3500	3.168	2397.446	4000	7.299

The results of the geospatial calculations show that in 2013, 38.604% of total area covers 621.418 to 1042.18ppm concentration of CH<sub>4</sub> while 1042.18 to 3500 ppm concentration of CH<sub>4</sub> covering 45.876% of the total area. In 2014, 39.168 % of the total area covers 856.583 to 1434.818 ppm concentration of CH<sub>4</sub> while 24.775 % of the total area covering 1434.818 to 4000 ppm concentration of CH<sub>4</sub> (Table-4).

Padhy and Varshney (2000) measured the ambient methane levels at different sites in the urban area of Delhi from November 1994 to June 1995 at an interval of 15 days. The CH<sub>4</sub> concentration varied between 173 and 949.2 ppm with an average of 412.1 ppm. Major sources were poorly managed solid waste

dumps, and choking sewers. Nguyen *et al.* (2010) analyzed the long-term patterns of urban CH<sub>4</sub> concentration in Seoul, Korea. Measurements were taken at an urban site, 3.8 m above ground level and 1 m from the busy road for 11-year period. The average CH<sub>4</sub> concentration measured during the study period was 224 ppm. The implementation of a nationwide air quality control policy was responsible for low values, where as in several other studies, high levels of methane in the urban environment, ranging from 200ppm (Altshuller 1968), 200ppm (Graedel *et al.*, 1986), and 330 ppm (Rowland *et al.*, 1990) have been reported. As mentioned earlier Jahanginagar University is a residential area but the CH<sub>4</sub> concentration are higher than the reported average

value. Scattered solid waste dumping site around the university area and biogas plant adjacent of University emit higher emission CH<sub>4</sub>. These reasons are similar with Padhy and Varshney (2000) work.

### Conclusion

In 2013 and 2014 average value of CO<sub>2</sub> were 388.889 and 422.222 respectively and average value of CH<sub>4</sub> in 2013 and 2014 were 1077.778 ppm 1144.444 ppm respectively. Geospatial analysis is showed to measure area 2013, 38.604% of total area covers 621.418 to 1042.18ppm concentration of CH<sub>4</sub> while 1042.18 to 3500 ppm concentration of CH<sub>4</sub> covering 45.876% of the total area. In 2014, 39.168 % of the total area covers 856.583 to 1434.818 ppm concentration of CH<sub>4</sub> while 24.775 % of the total area covering 1434.818 to 4000 ppm concentration of CH<sub>4</sub>. Spatial assessment is showed that high way road along the University area are main offender for higher CO<sub>2</sub> emission. The present study shows the impact of anthropogenic activities on the ambient concentration of GHGs emission of Jahangirnagar University being close the capital city of Bangladesh. This study is expected to have significant implications in rapidly urbanizing cities in delivering baseline information about GHGs emission.

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### References

**Altshuller AP.** 1968. Atmospheric analysis by gas chromatography. *Advances in Chromatography*, **5**, 229–262.

**Borjesson G, Danielsson A, Svensson BH.** 2000. Methane fluxes from a Swedish landfill determined by geostatistical treatment of static chamber measurements. *Environmental Science and Technology* **34 (18)**, 4044–4050.

**Cicerone RJ, Oremland RS.** 1988. Biogeochemical aspects of atmospheric methane. *Global Biogeochemical Cycles* **2 (4)**, 299–327.

**Davis JC.** 1986. *Statistics and Data Analysis in Geology*. Wiley, New York.

**Dlugokencky E, Masarie K, Tans P.** 1998. Continuing decline in the growth rate of the atmospheric methane burden. *Nature* **393**, 447.

**Gerlach TM, Douglas MP, McGee KA, Kessler R.** 2001. Soil efflux and total emission rates of magmatic CO<sub>2</sub> at the Horseshoe Lake tree kill, Mammoth Mountain, California, (1995-1999). *Chemical Geology* **177**, 101–116.

**Graedel TE, Hawkins DT, Claxton LD. (Eds.).** 1986. *Atmospheric chemical compound—Sources, occurrences and bioassay*. Orlando: Academic. 732.

**IPCC** 2001. Intergovernmental Panel on Climate Change, *Climate Change (2001): The Scientific Basis*. Cambridge University Press, Cambridge, UK.

**IPCC** 2007b. IPCC Forth Assessment report, *Climate Change: Synthesis Report*. IPCC.

**Maguire DJ, Goodchild MF, Rhind DW. (Eds.).** 1991. *Geographical Information Systems: Principles and Applications*. Longman Scientific and Technical, New York.

**Nguyen TH, Kim KH, Mab CJ, Cho SJ, Sohn JR.** 2010. A dramatic shift in CO and CH<sub>4</sub> levels at the urban location in Korea after the implementation of the Natural Gas Vehicle Supply (NGVS) program. *Environmental Research*, **110**, 396–409.

**Padhy PK, Varshney CK.** 2000. Ambient methane levels in Delhi. *Chemosphere - Global Change Science*, **2**, 185–190.



- Padhy PK, Varshney CK.** 2000. Ambient methane levels in Delhi. *Chemosphere - Global Change Science*, **2**, 185–190.
- Rasmussen RA, Khalil MAK.** 1984. Atmospheric methane in the recent and ancient atmospheres: concentrations, trends, and inter hemispheric gradient. *Journal of Geophysical Research* **89**, 11599–11605.
- Rowland FS, Harries NRP, Blake DR.** 1990. Methane in cities. *Nature*, **347**, 432–433.
- Sahay Samraj, Ghosh Chirashree.** 2012. Monitoring variation in greenhouse gases concentration in Urban Environment of Delhi. *Environ Monit Assess* **185**, 123–142 DOI 10.1007/s10661-012-2538-8.
- Shipham M, Bartlett K, Crill P., Harriss R, Blaha D.** 1998. Atmospheric methane measurements in central New England: an analysis of the long term trend and the seasonal and diurnal cycles. *Journal of Geophysical Research* **103 (D9)**, 10621–10630.
- Spokasa K, Graffa, Morceth M, Aranb C.** 2003. Implications of the spatial variability of landfill emission rates on geospatial analyses. *Waste Management* **23**, 599–607.
- Sookun Anand, Ravindra Boojhawon, Soonil DDV, Rughooputh,** 2014. Assessing greenhouse gas and related air pollutant emissions from road traffic counts: A case study for Mauritius. *Transportation Research Part D* **32**, 35–47.
- Wade Tasha and Sommer Shelly** 2001. A to Z GIS. -268p. ESRI (Environmental System Research Institute)-USA.
- WMO** 1998. Scientific Assessment of Ozone Depletion, 1998. World Meteorological Organization, Washington, DC. (Report No. 44).