



INDOOR/OUTDOOR AIR QUALITY NITROGEN-DIOXIDE RELATIONSHIP AT SELECTED LOCATIONS IN AGBOR; NIGER-DELTA REGION.

J.I. Uche^{a*} and E.E. Ukpebor^b

^aDepartment of Chemistry, College of Education, Agbor.

^bDepartment of Chemistry, University of Benin, Benin-City.

E-Mail: ucheifeanyij@yahoo.com

Abstract

Much of the health impact from air pollution worldwide seems to occur among the poorest and most vulnerable populations. The main aim of this study was to determine the levels of nitrogen dioxide concentrations both outside and inside homes and estimate the indoor/outdoor ratios at selected areas in Agbor. Measurement was carried out at ten locations indoor and outdoor by use of Palmes diffusion tubes. Indoor and Outdoor NO₂ levels were determined every two weeks for the period October, 2008- September, 2009. The maximum mean monthly indoor and outdoor NO₂ values of 50.02µg/m³ and 47.32µg/m³ respectively were recorded at site AG/TL. The annual mean outdoor NO₂ concentration ranged from 8.75 – 37.86µg/m³ while the annual mean indoor NO₂ levels ranged from 7.28 – 39.93µg/m³. The mean indoor/outdoor ratio was 0.91 (s = 0.11). Statistical analysis show that there exist a positive correlation between indoor and outdoor NO₂ (r = 0.87; p ≤ 0.05). When compared with statutory limits nationally (FMENV) and internationally (WHO), the values of NO₂ obtained in this study both indoor and outdoor lies below the permissible limits.

Keywords: Nitrogen dioxide, Correlation, indoor, outdoor, Agbor, Palmes diffusion tube.

Introduction

As a result of a variety of human activities, a large number of different toxic pollutants are emitted into the atmosphere. Exhaust emissions from automobiles contribute directly to adversely high ambient air pollutant concentration which is emitted in close proximity of the breathing zone of people. Although Nitrogen-dioxide (NO₂) occurs naturally in the environment, the largest contribution to ambient concentration comes from the burning/use of fossil fuel^(1,2). The principal removal mechanism of NO₂ from the air is its oxidation by OH to form nitric acid (HNO₃), which then collects on aerosols or dissolves in precipitation and is subsequently scavenged by rainfall. Other pathways for direct NO₂ removal occur through canopy scavenging of NO₂ and direct, dry deposition of NO₂, HNO₃ and particulate nitrates to the land surface and the ocean.

Wide ranges of air pollutants are generated outdoors and are either known or suspected of causing adverse effects to human health and the environment⁽³⁾. Principally, pollutants found in urban areas are from short range sources including pollution from vehicle exhausts, combustion, standby generators, construction, demolition and nuisance sources such as cooking smells from kitchen extracts.

People spend most of their time indoors yet the majority of pollutant concentration data are based on measurement conducted outdoors. Outdoor

pollutant concentration may not be reliable indicators of indoor and personal pollutant sources⁽⁴⁾. Although the toxicology of NO₂ exposure suggest the potential for respiratory symptoms and loss of lung function^(5, 6) evidence from three decades of epidemiological studies linking NO₂ exposure to adverse health effects has been inconsistent. Some inconsistency may be explained by differences in measures of exposures [acute^(7, 8) vs Chronic^(7, 8, 9, 10)]. Long term exposure to high concentration of NO₂ is associated with 20% increase in the risk of respiratory illnesses in children^(5, 11). The main aim of this study was to determine the levels of nitrogen dioxide concentrations both outside and inside homes and estimate the indoor/outdoor ratios at selected areas in Agbor.

Methodology

Sampling Location

This study was conducted in Agbor(headquarters of Ika South Local Government Area, Delta-State) ,an emerging commercial transitory town for commuters travelling to the North, East and western parts of Nigeria.

Sampling Sites

Ten sampling points outdoor evenly and well spread with different traffic densities, human and commercial activities were selected on the basis of security of equipment, accessibility and available receptors to obtain maximum spatial variation on



NO₂ levels. The outdoor sampling points and their codes are presented in Fig 1 below. For the Indoor NO₂ monitoring, ten homes (one from each site) located approximately 6m away from the outdoor sampling points were selected for the indoor monitoring purpose. Indoor sampling was done at the centre of the living room. However, the selection of homes used in this study was purely voluntary.

NO₂ Monitoring

NO₂ measurement was carried out by using Palmes diffusion tubes. A Palmes diffusion tube/sampler is a device capable of taking up a sample of a gas or vapor from the atmosphere through diffusion or permeation into the interior of the tube where it is subsequently trapped by means of adsorption on

the adsorbent. The sampler is basically an acrylic tube of 8.20cm long with cross-sectional area of 0.82cm² machined at each end to accommodate closely fitting polyethylene cap. The sensitivity of this particular tube length and the sensitivity and selectivity of triethanolamine for NO₂ have been reported^(1, 12, 13). At the sampling locations, the tubes were exposed at a height of about 1.5 – 2.0m above the ground level. Nitrogen dioxide monitoring was done every two weeks for a period of twelve months (October 2008 – September 2009). A UV/visible spectrophotometer (Spectronic, 21 D) was used to determine the absorbance of both blanks and air samples at 540nm. The atmospheric concentration of NO₂ during the measuring period was calculated as described by Palmes et.al⁽¹⁴⁾.

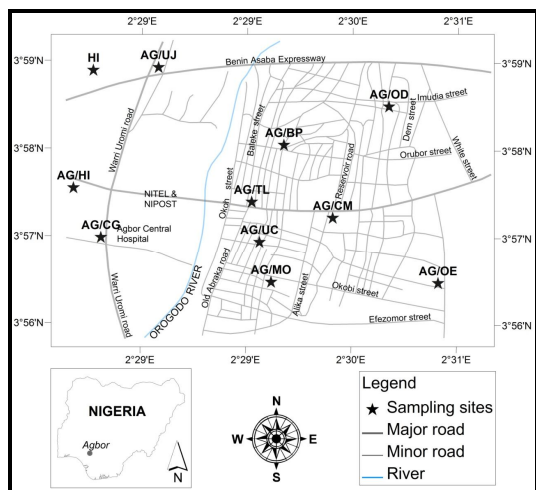


Fig 1 Base map of Agbor (Boji-Boji) town situated in Delta state, Nigeria showing street pattern and sampling sites

Results and Discussion

The result of the outdoor and indoor NO₂ levels for the period is shown in Table 1 below. From the results, the maximum monthly indoor (ID) and outdoor (OD) values of 50.02µg/m³ and 47.32µg/m³ respectively was obtained in the dry season months of December'08 and January'09 at site AG/TL. The high indoor and outdoor NO₂ values recorded for this site correlates with the high traffic density and the use of generators for commercial activities observed in this area. Other locations with high ID/OD values are; AG/UC 44.18µg/m³ (ID), 42.18µg/m³ (OD) and AG/CM 40.87µg/m³ (OD). Generally the ID and OD values recorded for sites AG/OD and AG/OE were low (6.11µg/m³, 6.92µg/m³ and 7.00µg/m³). This may be due to the fact that these sites are purely

residential and located at the outskirts of the city with very low traffic.

The annual mean ID/OD NO₂ concentration for the different locations does not follow any clearly defined pattern (Table 2), in some locations the ID values were higher than the OD values (AG/TL). This could be due to the fact that some of the homes studied had very poor ventilation system and also likely due to the fact that greater photochemical reactions of NO₂ can occur in the atmosphere. Generally, the mean indoor/outdoor ratio was 0.91 (s = 0.11). This high indoor/outdoor ratio is a sign of a high level of penetration of outdoor air indoor.

Regression Equation=4.097+0.898x
 SpearmanRank Correlation=0.867

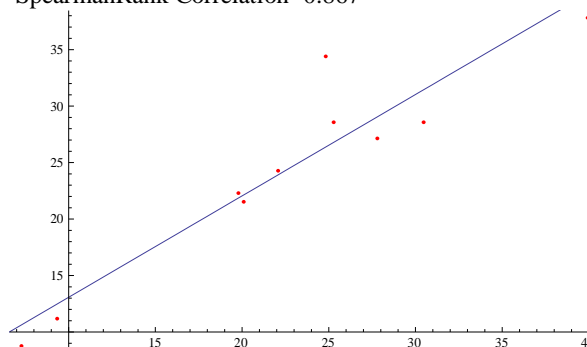


Fig 2. Graph of indoor vs outdoor NO₂ Concentration

Fig 2 shows the correlation between ID and OD NO₂. There was high correlation between ID and OD air (r = 0.87, p < 0.05); this could be due to factors such as generation rate of pollutant indoors, outdoor pollutant concentration, air exchange rate and pollutant penetration efficiency from outdoor to indoor environment^(15, 16).



Table 1. Mean monthly Indoor/Outdoor NO₂ concentrations (µg/m³)

		Locations									
Period		AG/UJ	AG/HI	AG/CG	AG/TL	AG/UC	AG/BP	AG/OE	AG/CM	AG/MO	AG/OD
Oct '08	ID	27.00	25.10	16.31	32.41	26.26	19.10	5.62	23.11	23.63	11.42
	OD	24.00	23.11	18.21	29.70	24.70	19.86	7.00	34.10	27.11	15.22
Nov '08	ID	30.10	26.80	22.44	40.65	32.47	23.42	5.94	25.31	21.77	12.69
	OD	29.49	31.06	24.04	38.93	30.20	26.59	9.04	39.35	25.21	17.44
Dec '08	ID	30.42	27.02	25.00	50.02	44.18	22.76	8.03	28.93	23.04	12.16
	OD	33.40	35.31	25.36	46.80	42.54	26.27	10.00	40.87	26.82	11.70
Jan '09	ID	30.18	27.10	21.37	49.34	34.71	24.00	8.68	28.76	22.97	13.00
	OD	33.17	35.06	24.36	47.32	33.11	25.87	10.03	38.39	25.77	15.69
Feb '09	ID	29.25	26.88	23.79	46.77	33.01	24.03	7.99	26.91	22.57	13.32
	OD	30.88	32.44	24.02	45.63	32.04	25.81	9.61	37.88	25.01	15.49
Mar '09	ID	29.88	26.75	24.00	48.00	32.68	25.04	7.85	26.88	22.01	13.44
	OD	31.66	32.95	24.77	45.60	31.33	26.00	9.20	37.10	24.63	15.21
Apr '09	ID	27.41	27.18	17.61	30.00	21.72	24.79	7.44	25.25	15.11	6.04
	OD	18.66	20.14	18.10	27.60	18.79	26.50	9.02	30.23	14.77	7.02
May '09	ID	28.76	25.33	18.06	32.74	23.91	23.68	6.92	24.17	16.14	6.73
	OD	22.01	22.66	19.42	30.20	22.60	25.20	8.11	31.47	18.62	7.11
Jun '09	ID	25.32	22.60	18.73	35.00	32.64	15.66	7.00	22.11	17.20	6.00
	OD	27.44	27.40	20.00	34.10	30.15	18.70	8.94	30.00	19.77	8.04
Jul '09	ID	24.91	22.13	18.19	35.96	30.18	20.04	7.68	23.06	17.14	5.18
	OD	25.23	28.10	20.46	34.77	28.26	24.20	8.11	32.11	21.00	6.11
Aug '09	ID	25.00	24.00	17.24	40.00	27.34	20.68	8.00	22.00	18.73	6.21
	OD	24.01	28.00	19.77	37.20	24.10	23.79	7.97	30.02	20.18	6.92
Sept '09	ID	25.72	22.11	18.39	38.23	26.95	21.25	6.23	21.65	17.00	6.03
	OD	26.20	27.01	20.20	36.41	24.66	23.00	8.00	30.77	19.02	8.00



Table 2 Indoor/Outdoor NO₂ Ratio

s/n	Code	Annual mean NO ₂ (µg/m ³)		
		Indoor	Outdoor	Ratio
1	AG/UJ	27.83±2.187	27.18±4.651	1.02
2	AG/HI	25.25±2.040	28.60±4.936	0.88
3	AG/CG	20.09±3.034	21.56±2.718	0.93
4	AG/TL	39.93±7.088	37.86±7.056	1.05
5	AG/UC	30.50±5.920	28.54±6.195	1.07
6	AG/BP	22.08±2.785	24.32±2.619	0.91
7	AG/OE	7.28±0.950	8.75±0.924	0.83
8	AG/CM	24.85±2.580	34.36±4.098	0.72
9	AG/MO	19.78±3.160	22.33±3.936	0.88
10	AG/OD	9.35±3.522	11.16±4.355	0.84

Intercomparison of the annual mean ID and OD values with regulatory limits show that the values obtained in this study are well below the World Health Organization ⁽¹⁷⁾ annual limit of 40µg/m³ and the Federal Ministry of Environment ⁽¹⁸⁾ daily limit of 75-113µg/m³. Mandel et al., ⁽¹⁹⁾ studied indoor NO₂ concentrations in some residences of Calcutta and also found annual mean NO₂ concentration well below the values prescribed by EC for European countries (45ppb).

Conclusion

In this study, ID/OD NO₂ relationship was investigated in ten locations. Indoor NO₂ values correlate positively with the Outdoor values. All sampling sites from where air samples were investigated gave NO₂ levels below the limits set nationally (FMENV) and internationally (WHO). The potentials of the data generated in this study can form the basis for environmental awareness and implementation of appropriate ambient air quality.

REFERENCES

1. E.E. Ukpebor, J.E. Ukpebor and K. Obahiagbon (2005); Measurement of Nitrogen dioxide in Ewohimi, Edo-state, Nigeria using diffusion tubes. *Global Journal of Pure and Applied Sciences*. 11(3):379-382.

2. R. A. Etiuma, A. U. Etiuma and I. E. Uwah (2007). "Monitoring of NO₂ concentration in Calabar South Local Government Area of Cross-River State, Nigeria." *J. Chem. Soc. Nigeria*. 32 (2): 211 – 217.

3. DETR (2000). The air quality strategy for England, Scotland, Wales and Northern Ireland. Department of the Environment, Transport and the Regions, the Scottish Executive. The National Assembly for Wales and the Department of Environment for Northern Ireland. The Stationary Office, United Kingdom, ISBN 0101254821.

4. L.Wallace, J. Quakenboss, and C. Rhodes (1997): AWMA/EPA Symposium on the Measurement of Toxic and related air pollutants. Research Triangle Park, NC, 860 – 871.

5. J. M. Samet and M. J Utell (1990). The risk of nitrogen dioxide: what have we learnt from epidemiological and clinical studies? *Toxicol Ind Health* .6:247-262.

6. M. Utell and M.W Frampton, (1997) Oxides of nitrogen in Roth R.A(ed) Comprehensive toxicology: Oxides of nitrogen. Cambridge, UK: Cambridge University Press. Pp303-312.

7. R. Basu and J.M .Samet (1999). A review of the epidemiological evidence on health effects of nitrogen dioxide exposure from gas stoves. *J. Environ Med*. 1:173-187.

8. M. Nitschke, B.J. Smith; L.S. Polotto, D. L. Pisaniello; M. J. Abramson and R.E. Ruffin (1999), Respiratory health effects of nitrogen dioxide exposure and current guidelines. *Int.J Environ Health Res*. 9:39-53.

9. B. Brunekreef (2001), NO₂: the gas that won't go away. *Clin Exp Allergy*.31:1170-1172.

10. J. M. Samet and M. L. Bell (2004), Commentary: nitrogen dioxide and asthma redux. *Int J Epidemiol*. 33:215-216.

11. R. A. Etiuma, I. E. Uwah and A. U. Etiuma (2006); "Level of Nitrogen Dioxide (NO₂) in Calabar city, Nigeria and Health Implications." *International Journ. Chem.*, 16 (4): 229 – 233.

12. G.W. Campbell, J. R. Stedman and K. Stevenson (1994). A Survey of Nitrogen dioxide Concentrations in the United Kingdom Using Diffusion tubes, July – December 1991. *Atmospheric Environment* 28: 477-486.

13. D. H. F. Atkins and D.S. Lee (1995). Spatial and Temporal Variation of Rural Nitrogen dioxide Concentrations across



- the United Kingdom. *Atmospheric Environment* 29 (2): 223-229.
14. E. D. Palmes, A. F. Gunmison, J. Dimattio and C. Tomzyk (1976). Personal Sampler for Nitrogen dioxide. *Am. Ind. Hyg. Assoc. J.* 37: 570-577.
15. R. Kamens, C. T. Lee, R. Weiner, D. Leith (1991); A study to characterize indoor particles in three non smoking homes. *Atmospheric Environment.* 25:939-948.
16. T. L Thatcher and D. W. Layton (1995); Deposition, re-suspension and penetration of particles within a residence. *Atmos Environ.* 29 (13):1487-1497.
17. WHO (2000), World Health Organisation: Guidelines for Air Quality, Geneva.
18. FMENV (1991); Guidelines and Standards for Environmental Pollution control in Nigeria, Federal Ministry of Environment, Abuja, Nigeria.
19. R. Mandal, S. Thakur, B.K. Sen, I. Bandyopadhyay and S. Sen (1997); Indoor nitrogen dioxide concentrations in some Calcutta residences. *Indian J. Environment Protection.* 17 (2):99-102.