Particle Counts and Size Distributions in Local Garri Processing Environment

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Abstract

Previous studies have linked morbidity and mortality of individuals to airborne particles resulting from anthropogenic sources. Fine particles (particles $\leq 2.5 \mu m$) are known to have harmful effects on human health and environment compared to the coarse particles (2.5 μ m < d_p <10 μ m). Sadly, developing countries experiencing rapid growth are characterized with inadequate environmental action and policy. Garri processing has been linked to contribute to ambient particulate matter pollution. The present study investigates the number and sizes of particulate emitted as a result of this anthropogenic activity. Continuous measurement of number concentration and size distributions of particles of diameters 0.3 µm, 0.5 µm, 1.0 µm, 2.0 µm and 5.0 µm were conducted, using a GT -321 particle counter for day and night cases November 2012 at four major local garri processing locations in the ancient city of Oyo, Nigeria. On the average, the total particle number concentration for the selected four locations were $2.3 \times 10^6 - 8.9 \times 10^6$; $0.085 \times 10^6 - 4.487 \times 10^6$; $0.073 \times 10^6 - 0.912 \times 10^6$; $0.022 \times 10^6 - 0.654 \times 10^6$; and $0.002 \times 10^6 - 0.155 \times 10^6$ for particle size 0.3μ m, 0.5μ m, 1.0μ m, 2.0μ m and 5.0 μ m respectively for day time readings and $1.021 \times 10^6 - 8.793 \times 10^6$; $0.071 \times 10^6 - 3.323 \times 10^6$; 0.051×10^6 -1.903×10^{6} ; $0.012 \times 10^{6} - 0.278 \times 10^{6}$; and $0.003 \times 10^{6} - 0.015 \times 10^{6}$ for particle size $0.3 \mu m$, $0.5 \mu m$, $1.0 \mu m$, 2.0 µm and 5.0 µm respectively for night time measurements. Daily cycle of particle numbers were obvious with high concentrations recorded during day time but lower level at night time which is consistent with the trend of the garri processing activity. The findings suggested that higher fraction of fine particles $(0.3 - 2 \mu m)$ are produced and retained in the ambient air from this activity.

Keywords: Garri, number concentration, size distribution, particulate matter

1.0 Introduction

Antropogenic contribution of particulate matter and other gaseous pollutants increase the ambient loading of aerosols in air. Epidemiological studies have linked morbidity and mortality of individuals to exposure of particulate matter especially the fine particles, $PM_{2.5}$ and the coarse particles, PM_{10} (Brauer *et al.*, 2001; Penttinen *et al.*, 2001; WHO, 2003; Franklin *et al.*, 2007; Bell *et al.*, 2009; Ostro *et al.*, 2009). Aside the health impacts, they affect visibility (Andreae and Merlet, 2001; Mahmoud, 2010), soil surfaces when deposited (Chow *et al.*, 2002), and affect the earth's climate (MacCracken, 2008; Alleman *et al.*, 2010). Studies have tend to investigate

ambient concentrations of pollutants resulting from some anthropogenic sources (Sonibare and Jimoda, 2009; Jimoda *et al.*, 2011). However, the issue of air quality impact from some local manufacturing processes is yet t receive tangible attention from the policy makers and air quality researchers in the developing countries.

Cassava is one of the most important staple food crops grown in tropical Africa (IITA, 1990). It plays a major role in efforts to alleviate the African food crisis because of its efficient production of food energy, year-round availability, tolerance to extreme stress conditions, and suitability to present farming and food needs in Africa (Oyegbami *et al.*, 2010). Sanni and Olubamiwa (2004) and IITA (1990) estimated that over 70% of cassava yield in Nigeria is processed into 7- 10 million tonnes of garri (cassava flour) annually (FAO, 2004). People engage in the local processing of the product in many towns of southwestern Nigeria (Adebayo, 2009; Adejumo and Raji, 2010; Makanjuola *et al.*, 2012). Oyo town is known for its high garri processing capability. It provides a high percentage of garri consumed in the adjourning towns and cities.

The processing of cassava to garri after harvesting in Oyo town is intensive during the dry season, which may contribute significantly to the ambient air pollution. Open frying is the most common method used by garri processors. Such frying often takes place in large extended pots supported by muds. Fire woods which are in plentiful supply are used majorly to supply the required heat for the processing. This study focuses on emissions from open burning of agricultural biomass such as cassava peels and firewood involved in garri processing. This is for the assessment of the number concentration of particulate matter of different sizes resulting from garri processing.

2.0 Materials and Methods

2.1 Sampling Area

Oyo is a town in Southwestern Nigeria. It is located in the region of Oyo State with latitude and longitude values of 7° 47' 26" N and 3° 56' 15" E respectively. Its population ranges between 250000 and 500000. It has an equatorial climate notably with dry and wet season, average daily temperature ranging between $25^{\circ}C$ (77°F) and $35^{\circ}C$ (95°F). The measurement and samplings were done at various stations namely Irepo (G1), Sabo (G2), Ajonibadi (G3), Ajegunle (G4) and G_c (a residential area) ischosen as the control location as shown in Figure 1. The sampling stations (G1, G2, G3 and G4) are characterized by numerous cassava processing sheds, where cassava is being peeled, ground and fried. At the sampling sites, the particle concentrations were measured at about 1 m above the ground overlooking the locations where continuous burning takes place. There are no tall buildings or structures nearby to stop free flow of ambient air for quality readings of particle concentration.

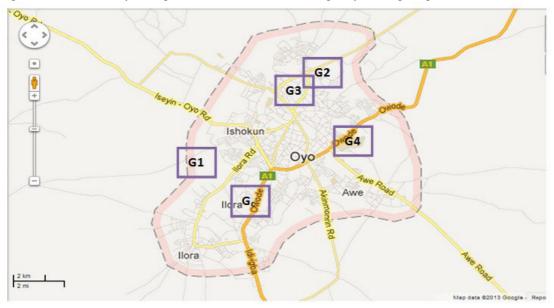


Figure 1: The study locations in Oyo, Southwestern Nigeria (Google Maps, 2013)

2.2 Particulate Matter Number and Size Characterization

Particulate matter (PM) was measured with GT-321 Particle Counter, an equipment from Met One Instruments. It is handheld, battery operated and completely portable unit measuring five number ranges of Total Suspended Particles: $PM_{0.3}$, $PM_{0.5}$, PM_1 , PM_2 , and PM_5 . It has a sampling period of 2 minutes and a flow rate of 2.83 l/min. To measure, it is placed at 1 m above the ground level, switched on in the environment of interest and the measured concentration read directly on the screen after particle capturing. Measured concentrations were compared with the International Standard Organisation standard (ISO 14644) for airborne particulates for cleanrooms and clean zones (Table 1).

| ISO Classification Number (N) | Maximum Concentrations Limits (particles/m ³) | | | | | | |
|-------------------------------|-----------------------------------------------------------|--------|--------|----------|---------|--------|--|
| | 0.1 µm | 0.2 μm | 0.3 µm | 0.5 µm | 1.0 µm | 5.0 µm | |
| ISO Class 1 | 10 | 2 | | | | | |
| ISO Class 2 | 100 | 24 | 10 | 4 | | | |
| ISO Class 3 | 1000 | 237 | 102 | 35 | 8 | | |
| ISO Class 4 | 10000 | 2370 | 1020 | 352 | 83 | | |
| ISO Class 5 | 100000 | 23700 | 10200 | 3520 | 832 | 29 | |
| ISO Class 6 | 1000000 | 237000 | 102000 | 35200 | 8320 | 293 | |
| ISO Class 7 | | | | 352000 | 83200 | 2930 | |
| ISO Class 8 | | | | 3520000 | 832000 | 29300 | |
| ISO Class 9 | | | | 35200000 | 8320000 | 293000 | |

Source: ISO (2004)

3.0 Results and Discussion

3.1 Particulate Matter size distribution and number concentration

On the average, the total particle number concentration for the four selected garri processing locations G1, G2, G3 and G4 range between 2.3 x 10^6 and 8.9×10^6 ; $0.085 \times 10^6 - 4.487 \times 10^6$; $0.073 \times 10^6 - 0.912 \times 10^6$; $0.022 \times 10^6 - 0.654 \times 10^6$; and $0.002 \times 10^6 - 0.155 \times 10^6$ for particle size 0.3μ m, 0.5μ m, 1.0μ m, 2.0μ m and 5.0μ m respectively for day time readings and $1.021 \times 10^6 - 8.793 \times 10^6$; $0.071 \times 10^6 - 3.323 \times 10^6$; $0.051 \times 10^6 - 1.903 \times 10^6$; $0.012 \times 10^6 - 0.278 \times 10^6$; and $0.003 \times 10^6 - 0.015 \times 10^6$ for particle size 0.3μ m, 0.5μ m, 1.0μ m, 2.0μ m and 5.0μ m respectively for night time readings. In all the locations, it was noticed that the particulate matter size of 0.3μ m (PM_{0.3}) has the concentrations far above other particulates (PM_{0.5}, PM₁, PM₂ and PM₅) measured. Figures 2 - 5 show the percentage contribution of each particulate matter size to the overall particulate matter number concentration in each processing location. For the control location,

 $PM_{0.3}$ contributed the highest percentages ranging from between 71% and 87%. It is followed by $PM_{0.5}$ with contribution of between 8% and 22%. The least contributor is the PM_5 with less than 1% contribution in all the study locations. Figure 6 indicated that G4 is the highest contributor of TSP into the air followed by G3, G2 and then G1. This is consistent with the trend and number of active processing point in each location. G4 has the highest number of frying point. It has about 198 points with an estimated firewood usage of 148.5 tonnes/day. G3 has 50 frying points with an estimated firewood usage of 35.75 tonnes/day, G2 with 47 points with about 35.25 tonnes of firewood used per day and G1 with 21 frying points with about 15.75 tonnes of firewood used per day.

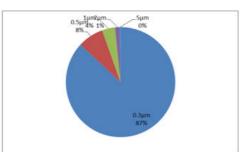
3.2 Comparison with Standard

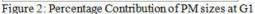
The measured particulate number concentration were compared with the ISO 14644 standards (Table 1) for particles sizes 0.3 μ m, 0.5 μ m, 1.0 μ m, and 5.0 μ m. PM₂ does not have a ISO 14644 standard. Averaged measured concentration for particulate size 0.3 μ m which is the most harmful of all the sizes measured were above the highest ISO class 6 of maximum concentration limit of 102000 particles/m³. However, particle sizes 0.5 μ m, 1.0 μ m, and 5.0 μ m measured fall within the ISO class 8 which has the maximum concentration limit of 3520000 particles/m³, 832000 particles/m³, and 29300 particles/m³ respectively (Tables 2 and 3). For the control location Gc, PM_{0.3} also exceeds the ISO 14644 limit of 102000 particles/m³. However, PM_{0.5} and PM₁, fall in class 7, and PM 5 falls in class 7 and 8 for night and day time readings respectively.

| Sampling Location | Mean Concentration (particles/m ³) | | | | |
|-------------------|------------------------------------------------|---------|--------|-------------------|--------|
| | PM 0.3 | PM 0.5 | PM 1.0 | PM _{2.0} | PM 5.0 |
| G1 | 4343034 | 233923 | 176066 | 65053 | 10605 |
| G2 | 5819408 | 608306 | 159190 | 106951 | 7268 |
| G3 | 5921870 | 609490 | 118105 | 91514 | 8290 |
| G4 | 7271768 | 2192791 | 440351 | 165325 | 6620 |
| Gc | 1973544 | 213655 | 71331 | 34022 | 3503 |

Table 2: Measured Day time Particulate Matter Number Mean Concentration measured during study

| Sampling Location | Mean Concentration (particles/m ³) | | | | | |
|-------------------|------------------------------------------------|-------------------|--------|-------------------|--------|--|
| | PM 0.3 | PM _{0.5} | PM 1.0 | PM _{2.0} | PM 5.0 | |
| G1 | 4355117 | 1413703 | 498491 | 69065 | 7130 | |
| G2 | 5517710 | 1052906 | 150855 | 76278 | 7714 | |
| G3 | 6161774 | 1049619 | 418660 | 92725 | 8823 | |
| G4 | 4355117 | 1413703 | 498491 | 69065 | 7130 | |
| Gc | 1936261 | 212435 | 52357 | 21830 | 1752 | |





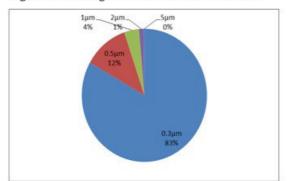


Figure 4: Percentage Contribution of PM sizes at G3

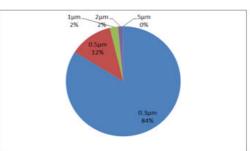


Figure 3: Percentage Contribution of PM sizes at G2

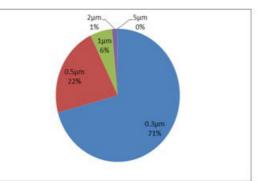


Figure 5: Percentage Contribution of PM sizes at G4

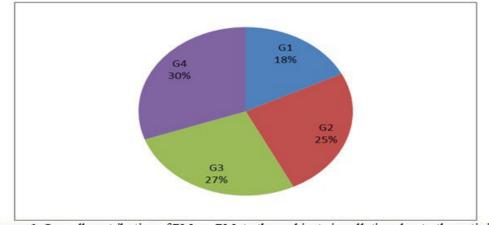


Figure 6: Overall contribution of PM0.3-PM5 to the ambient air pollution due to the activity

3.3 Daily cycle for particulate matter

The daily cylces for the measured particulate matter number concentration were shown in figures 7 - 14 for the four locations considered in this study. The maximum concentrations were measured on a high activity day (day 4) which is a Monday. Higher concentrations were observed during the day time compared to the night-time measurements. The results signify that the concentration changes are consistent with the trend of activity as it is expected. Fine particulates has the highest concentration because they are lighter in weight and are not easily deposited compared to the higher diameter particulates. This conforms with the assumption of the previous studies (Al - Rajhi *et al*, 1996, Masmoudi *et al.*, 2002; Pryor, 2006; Piazzola *et al.*, 2010; Kozawa *et al.*, 2012; Fang *et al.*, 2012) that deposition is size dependent. The higher the diameter of the particle, the more the tendency of it being deposited. Premised on this fact, it is assumed that a higher percentage of particles emitted from the activity, most especially the smallest ($PM_{0.3}$) will be retained in the air for a very long period as reflected in the night time concentrations.

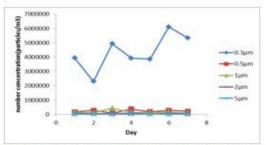


Figure 7: Day Time Number Concentration against Day for G1

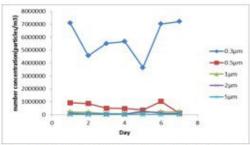


Figure 9: Day Time Number Concentration against Day for G2

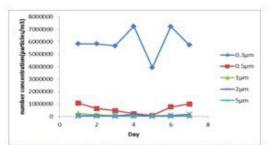


Figure 11: Day Time Number Concentration against Day for G3

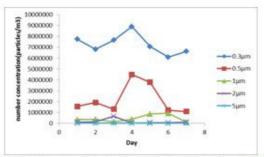


Figure 13: Day Time Number Concentration against Day for G4

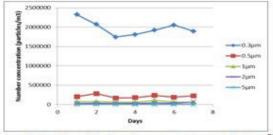


Figure 15: Day Time Number Concentration against Day for Gc

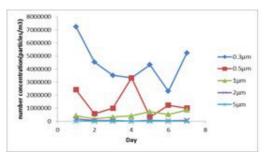


Figure 8: Night Time Number Concentration against Day for G1

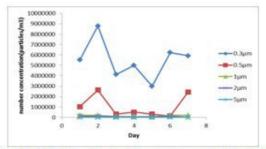


Figure 10: Night Time Number Concentration against Day for G.

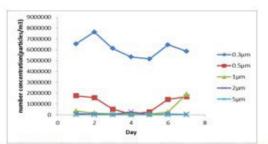


Figure 12: Night Time Number Concentration against Day for G3

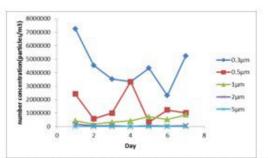


Figure 14: Night Time Number Concentration against Day for G4

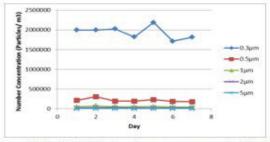


Figure 16: Night Time Number Concentration against Day for Gc

65

4.0 Conclusion

Particulate number concentration and their respective size distributions were measured over a period of seven days for five different particulate sizes in four different garri local processing locations. It can be said that data generated or measured at these sites should reasonably represent the ambient air quality in the domains. The finest fraction and the most harmful of the particulates measured ($0.3 \mu m$) in the locations have the highest concentrations and exceeded the maximum allowable limit of ISO 14644 standards. This suggests that most of the pollutants emitted from the activity are retained in the ambient air for long hours especially the fine particulate fractions. It is therefore imperative that the policy makers and all the authorities concerned need to find lasting solutions through the provision of kilns and industrial stoves to reduce/eliminate the use of firewood. Efforts should also be geared towards the industrial drying and conversion of cassava peels to animal feeds as against burning them. Future researches should tend to investigate the elemental composition of the particulate matter emitted from the activity and other gaseous pollutants that may arise especially methane.

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