ROOF RUNOFF WATER AS SOURCE OF POLLUTION: A CASE STUDY OF SOME SELECTED ROOFS IN ORLU METROPOLIS, IMO STATE, NIGERIA

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ABSTRACT. Samples collected from different runoffs; AR, GMR, TR, ASR, and DR for aluminum roof, galvanized metal roof, thatch roof, asbestos roof and ambient rainfall respectively were analyzed for pH, conductivity, turbidity, TDS, TSS, NO3, PO43-, Cd, Cu, Fe, Zn and Pb. The result indicates that the mean concentration of the parameters analyzed ranged from 5.8± 0.39 – 7.10±0.70, 22.25±11.70 -79.99± 3.40µScm⁻¹, 1.47±0.43 - 46.53±1.60mg/l, 11.90±0.93 - 59.83±1.62NTU, 15.53±0.70 - 204.53±5.08mg/l, 0.93±0.06 - 2.55±0.13 mg/l, 1.33±0.22 - 7.30±0.57mg/l respectively for pH, conductivity, TDS, turbidity, TSS, PO43- and NO3, and the levels of the heavy metals (in mg/l); Cd, Cu, Fe, Zn and Pb ranged from 0.0023±0.001– 0.0521±0.004, 0.052±0.01–0.2483±0.02, 0.0348±0.01–1.1120±0.07, 0.0161±0.01- 0.8093;±0.02 and 0.0106±0.01– 0.0499±0.002 respectively. Ranking of the heavy metal in roof runoff is in the order; Fe>Zn>Cu>Cd>Pb. The result compared with WHO standard showed elevated level of the parameters analyzed with Cd and Pb exceeding the limit. Though the result of this study showed some variability which is an indication of the type of roofing material; air quality of the environment and industrial activity going on in the area. It could be deduced from the result that roof runoff may be a non point source of environmental pollution owning to the release of heavy metals and other pollutants into the environment, and increased concentration of some of the pollutants as reported by this study suggests that roof runoff water could impact negatively to the environment and if consumed without being treated may be injurious to human health.

1. INTRODUCTION

Harvested rainwater has been increasingly used for potable and non-potable purposes which are as a result of decrease in the availability and quality of traditional water resources [1]. Significant level of water quality reduction could be attributed to non point sources of pollution [2]. Roof runoff water has been identified as a major contributor to the pollution of streams, rivers and estuaries [3,4,5], resulting from increasing urbanization [6], since roof runoffs are discharged into the environment without being previously treated [7]. Roof surfaces in urban settlements contribute excess nutrients and toxic metals to receiving waters [8,9,10,11], and these surfaces could cover from 12% in residential areas to 21% in commercial areas [8,12]. This therefore increases the overall toxic burden of the environment [13]. And the level of pollutants that may emanate from roof runoff depends on the roof materials, age, orientation and slope of the roofs, atmospheric depostions, rain events and meteorological conditions [14,15,16].

In order to develop alternative technologies to control roof runoff pollution accurate data on the quality and properties of runoff pollution from different roof surfaces is required before it is mixed and transported in a sewer. This type of information seem to be lacking at present, though there has been some reports of roof runoff waters. High concentration of heavy metals like Cd, Cu, Pb and Zn has been reported in roof runoff [17,18,19,20]. Olaoye, and Olaniany reported high level of pH, TH, Al, Cu, NO3, SO42- and presence of coliform bacteria in samples from asbestos, concrete and corrugated plastic roof, they noted that the aluminium roof was free from pathogenic
contamination [21]. Also reports about roof runoff water have been made elsewhere [22,23]. The roof materials mainly used in Orlu Metropolis are made of a variety of substances ranging from plant materials, galvanized metal sheets, aluminum to asbestos and others which seen to have direct influence on the tendency for the release of toxins to the environment.

The need to evaluate pollution level emanating from rainwater samples in roof runoff in Orlu metropolis became necessary owning to paucity of reports on this issue about Orlu and its surroundings and the fact that there are significant number of people who still depend on harvested rainwater from roof runoff because of inadequate supply of portable water to every home in the town. In addition to the fact that Orlu (fig.1) is very close to Ohaji/Egbema/Oguta area of Imo State known for crude oil related activities and gas flaring.

2. MATERIAL AND METHODS

Study area
Orlu area of Imo State is located within the geographical coordinate 5°47' North and 7° 2' East, within the tropical rainforest region with two distinct seasons; wet and dry seasons. In the recent time, the population of Orlu is fast increasing owning to urbanization and industrialization with a lot of commercial activities. Orlu is in the same zone with Ohaji/Egbema/Oguta area of Imo State where there has been crude oil exploration, drilling and gas flaring for more than forty years now.

Sampling and preservation of sample
Water samples were collected from four rain events in triplicate from each of the different roof types including sample from the ambient environment, referred to as direct rain water, in the locations referenced with Garmin GPSmap76. The study was conducted in Orlu, Imo State, within the months of June and August, 2013. The samples were collected from five different sources in triplicate in four different rain events at five different locations which include Umuaka, Banana junction, Umuna, Ogboko junction and Umuago Urualla. The samples from aluminum roof, galvanized metal roof, thatch roof, asbestos roof and direct rainfall from ambient environment were labeled AR, GMR TR, ASR and DR respectively. The parameters analyzed were pH, electrical conductivity, total dissolved solid (TDS), turbidity, total suspended solid (TSS), phosphate (PO$_4^{3-}$), nitrate (NO$_3^-$) and heavy metals (Cd, Cu, Fe, Zn, and Pb). Polyethylene bowls and bottles for sampling were previously washed with detergent and 0.1MHNO$_3$ and then rinsed with deionized water [22,23]. Contaminations of the samples were significantly reduced as samples were collected over elevated platform. Samples meant for heavy metal analysis were preserved according to [24,25].

![Fig. 1 Showing map of the study area](image-url)
3. ANALYSIS OF SAMPLE

pH and electrical conductivity (E.C) were measured with Hanna pH meter Hi 98107 and Hanna conductivity meter Hi98303. Turbidity, TDS and TSS were determined according to standard procedure [24,26,,27,28]. Nitrate was determined by cadmium reduction method using Hanna H1 83200 multiparameter bench photometer at a wavelength of 525nm. 10ml of the sample was poured into two sample cells, one was used as blank to zero the instrument. Nitrate reagent was added to the second sample cell and inserted into the cell compartment and timed for 4 minutes 30 seconds. After this time the read button was displayed and result displayed in mg/l of Nitrate. Also phosphate was determined by amino acid method using Hanna H1 83200 multiparameter bench photometer at 525nm. 10ml of the sample was poured into two sample cell bottles. One was used as blank to zero the instrument and 10 drops of H19371 A-O molybdate reagent and a packet of Hi9371B phosphate HR reagent B was added to the second sample cell bottle. It was gently shaken to dissolve and inserted into the cell compartment and timed for 5 minutes. At the end of this time the read button was pressed and result displayed in mg/l of phosphate. Heavy metals (Cd, Cu, Fe, Zn and Pb) were analysed in the sample using atomic absorption spectrometer AAS, Agilent FS 240 AAS (USA). The metals were quantified by appropriate calibration curves prepared in the same acid matrix with standard solution of the metals [29].

Quality Control
Glasswares and polythene bottles used were soaked in 0.01MHNO₃ overnight, washed, rinsed with deionized water and allowed to dry. The reagents that were used in this work were of analytical grade standard. The reagents HCl, HNO₃ and metal standard were supplied by Chemscience , Overri.

Statistical Analysis
All the results of this work were analyzed with Microsoft Excel 2007 version which was presented as mean of standard deviation, minimum and maximum values.

4. RESULTS AND DISCUSSION

Table 1 is the result of physiochemical analysis showing mean values, minimum values, maximum values and the standard deviations while fig.2 to fig.5 is the chart representation of the results. From table 1 and fig.2, it is observed that the mean pH values ranged from 5.8± 0.39 ~7.10± 0.70 and the values are in the order DR< AR<GMR<TR<ASR. This indicates that all the samples were acidic, indicating presence of acidic oxides in the atmosphere. The acidic nature of ambient rainwater may cause the chemical compounds (such as cadmium, copper, lead, zinc, and chromium) from roofing materials to leach into the harvested rainwater [30,31]. The mean pH value of the harvested runoff samples are within the WHO drinking water standard range of 6.5 to 8.5 which could be used for domestic purposes. Similar studies have reported an increase in pH value from ambient rainwater to harvested roof runoff water [7,32,33].

The mean conductivity as shown in table 1 and fig.2 ranged from 22.25 ± 11.70 -79.99± 3.40μScm⁻¹, the mean values of the conductivity are in the order TR>ASR>AR>DR>GMR. The result is typical of roof runoff water [33,34]. The sample from thatch roof runoff showed significantly high conductivity when compared with other roof runoff samples. And these values agree with reports made in similar studies [13,35].

The values for TDS (table 1 and fig.4) vary from (1.47±0.43) mg/l for DR (46.53±1.60)mg/l for TR. The results showed elevated level of TDS when compared with other samples this followed by ASR. This could be attributed to the tendency of this roof to retain more precipitation [2], which could facilitate vigorous decomposition of the roof surfaces and as well being released with roof runoff. TDS correlates with E.C, the higher the E.C the greater the TDS [36]. The result is in the order TR>ASR>AR>GMR>DR which are within the WHO limit of 1000mg/l.

The mean turbidity value ranged from 11.90±0.93 - 59.83±1.62NTU as shown in table 1 and fig.4. The least value was obtained from DR while the most turbid is that from TR and this is in the
order TR>AR>ASR>GMR>DR. The elevated value for thatch roof is not unexpected owning to its organic nature as noted for TDS.

The results as shown (tab.1 and fig.4) indicates that the TSS values varies from (15.53±0.70) mg/l in DR to (204.53±5.08)mg/l in TR. The level of TSS observed is in the order, TR>ASR>GMR>AR>DR, the level of TSS may depend on the dry days, roughness of the surface and rain intensity [37]: Elevated level of TSS can serve as carriers of toxics, which readily cling to suspended particles [32].

The mean phosphate level varies from (0.93±0.06) mg/l in DR to (2.55±0.13) mg/l in TR, and values are in the order DR<ASR<AR<ASR<TR. The phosphate level in roof runoff samples was greater than direct rain water from ambient air (table 1 and fig4). Again the concentration of phosphate were greater in TR runoff followed by ASR, this may be due to the ability of these roofs to retain contaminants [6]. In thatch roof, the fertilizer used during plant establishment could be the probable source of phosphate [38].

All the samples including that from ambient air contain elevated level of nitrate (table1 and fig.4). The concentration ranged from (1.03±0.22)mg/l in DR from ambient rain to (7.30±0.57) mg/l in TR runoff water sample and the values are in the order

| Parameters | Unit | Roof type | WHO |
|------------|------|-----------|-----|
| pH         |      | AR        | GMR | TR  | ASR | DR  |
| MIN        | 6.60±0.64 | 6.65±0.57 | 6.88±0.36 | 7.10±0.70 | 5.80±0.39 | 6.5-8.5 |
| MAX        | 7.20  | 7.30      | 7.20 | 7.90 | 6.30 | 6.90 |
| E.C µS/cm  | 50.43±2.18 | 22.25±1.10 | 79.99±3.40 | 65.68±7.18 | 33.05±3.35 | 100 |
| MIN        | 48.40 | 18.40     | 76.50 | 58.20 | 58.70 | 50 |
| MAX        | 53.50 | 25.80     | 84.30 | 75.40 | 36.50 | 50 |
| Turbidity NTU | 18.11±1.30 | 13.19±0.85 | 59.83±1.62 | 15.01±1.20 | 11.90±0.93 | 50 |
| MIN        | 16.75 | 12.76     | 57.85 | 13.96 | 10.51 | 5 |
| MAX        | 20.34 | 13.63     | 62.31 | 16.25 | 13.63 | 50 |
| TDS mg/l   | 10.34±5.62 | 7.88±0.58 | 46.53±1.60 | 13.19±0.76 | 1.47±0.43 | 1000 |
| MIN        | 9.20  | 7.30      | 44.80 | 12.30 | 1.14 | 1000 |
| MAX        | 11.75 | 8.52      | 48.20 | 14.20 | 2.06 | 1000 |
| TSS mg/l   | 48.51±3.39 | 92.93±6.50 | 204.53±5.08 | 189.2±6.48 | 15.53±0.70 | 500 |
| MIN        | 46.30 | 84.50     | 199.80 | 184.20 | 14.60 | 100 |
| MAX        | 51.40 | 98.40     | 210.30 | 193.60 | 16.30 | 100 |
| Nitrate mg/l | 3.85±1.41 | 3.68±0.84 | 7.30±0.57 | 4.35±0.95 | 1.33±0.22 | 50 |
| MIN        | 2.20  | 2.80      | 6.50 | 3.40 | 1.10 | 50 |
| MAX        | 5.30  | 4.80      | 7.80 | 5.40 | 1.60 | 50 |
| Phosphate mg/l | 1.75±0.13 | 1.38±0.17 | 2.55±0.13 | 2.23±0.25 | 0.93±0.06 | 0.03 |
| MIN        | 1.60  | 1.20      | 2.40 | 1.90 | 0.89 | 0.03 |
| MAX        | 1.90  | 1.60      | 2.70 | 2.50 | 1.02 | 0.03 |
| Cd mg/l    | 0.019±0.007 | 0.0349±0.005 | ND | 0.052±0.004 | 0.002±0.001 | 0.003 |
| MIN        | 0.0131 | 0.0302    | ND | 0.0486 | 0.0012 | 0.003 |
| MAX        | 0.0281 | 0.0412    | ND | 0.0561 | 0.0034 | 0.003 |
| Cu mg/l    | 0.2483±0.02 | 0.1878±0.02 | 0.0520±0.01 | 0.397±0.02 | 0.012±0.00 | 0.003 |
| MIN        | 0.2301 | 0.1280    | 0.0430 | 0.3740 | 0.012 | 0.003 |
| MAX        | 0.2710 | 0.2340    | 0.0610 | 0.4230 | 0.014 | 0.003 |
| Fe mg/l    | 0.507±0.23 | 1.112±0.07 | 0.0348±0.01 | 0.636±0.04 | 0.382±0.04 | 2.00 |
| MIN        | 0.4812 | 1.0210    | 0.0230 | 0.5800 | 0.3310 | 2.00 |
| MAX        | 0.5312 | 1.2030    | 0.0450 | 0.6840 | 0.4230 | 2.00 |
| Zn mg/l    | 0.577±0.03 | 0.8093±0.02 | ND | 0.552±0.32 | 0.016±0.01 | 3.00 |
| MIN        | 0.5420 | 0.7850    | ND | 0.0713 | 0.0101 | 3.00 |
| MAX        | 0.6210 | 0.8210    | ND | 0.7320 | 0.0221 | 3.00 |
| Pb mg/l    | 0.015±0.004 | 0.0224±0.01 | ND | 0.049±0.002 | 0.010±0.006 | 0.01 |
| MIN        | 0.0121 | 0.0080    | ND | 0.0484 | 0.0023 | 0.01 |
| MAX        | 0.0210 | 0.0412    | ND | 0.0522 | 0.0142 | 0.01 |
TR>ASR>AR>GMR>DR. Also higher level of nitrate was observed in TR runoff compared to others followed by ASR runoff. The mean value for all the samples exceeded the WHO permissible limit of 50 mg/l. This is an indication of atmospheric pollution by nitrate which could be deposited on the surface of the roof. The higher values observed in TR and ASR runoff water though within WHO permissible limit of 45mg/l could be attributed to higher retention capacity of the roofing materials and degradation of plant material in the case of thatch roof in a study of road and roof runoff waters as a source of pollution in a big urban agglomeration (reported Higher concentration of nitrate in the range of 0.92-62.000 mg/l were reported in road and roof runoff in Didansk, Polaul [5]. Similarly, Olaoye and Olaniyan reported nitrate range of 31.90-39.00 mg/l in their study of the quality of rainwater from different roof material in Ogbomoso north L.G.A., Oyo State f Nigeria [21].

The concentration of the metal (table 1 and fig.5) detected ranged from (0.0023±0.00)mg/l for Cd in DR to (1.1120±0.07)mg/l for Fe in GMR, the heavy metals are in the order Fe>Zn>Cu>Cd>Pb. The level of Fe in the roof runoff samples are in the order GMR>ASR>AR>DR>TR. Cd, Zn and Pb were not detected in TR runoff sample. Elevated level of Cd, Cu, Fe and Pb were observed in ASR roof runoff sample than others, which were above 0.003mg/l and 0.01mg/l respectively for Cd and Pb WHO limit [39]. Cu and Fe were detected in all the samples and the concentrations are within the WHO limit. Higher values of the metals studied has been reported in a similar study [6,7]. It has been reported that the major source of Zn. Cu and Pb in urban runoff water are roof corrosion [17]. Also these metal could
leached into the environment due to vehicular traffic during tire and brake wearing, gas emissions and corrosion of metallic pieces [16]. Elevated values of Fe and Zn in aluminum and galvanized metal roof runoff samples may be due to erosion of zinc roofing material, galvanized gutters and bulk atmospheric deposition [10, 16, 40, 41]. Higher value of Pb and Cd could be due to tendency of Pb to strongly adhere to particles [42], Pb and Cd are associative [43]. Cd, Zn and Pb which were not detected in TR runoff sample may be due to the tendency of this roof type to retain heavy metals, owing to stabilization of the media through repeated wetting, drying cycles and formation of chelates with organic materials [37, 44], since composited biosolids have been found to lower the solubility of Zn and increases its retention [45].

5. CONCLUSION

Though the result of this study showed some variability which is an indication of the type of roofing material; air quality of the environment and industrial activity going on in the area. The study showed that roofs could be a source of pollution since the surfaces are available for deposition of atmospheric particles and the leach-ability of the roof surfaces could have contributed to the level contamination observed in the roof runoff samples. Also it could be deduced from the study that roof runoff may be a non point source of environmental pollution owning to the release of heavy metals and other pollutants into the environment. The increased concentration of some of the pollutants as reported by this study suggests that roof runoff water could impact negatively to the environment and if consumed without being treated may be injurious to human health.

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