in Challawa Industrial Area of Kano Metropolis, Nigeria. Journal of Experimental Research March 2019, Vol 7 No 1 ^{*}Bulus H¹, Ezeh EC², Shekarri TB¹, Pascalina Li¹ Email: editorinchief.erjournal@gmail.com editorialsecretary.erjournal@gmail.com Nigerian Institute of Leather and Science Technology, PM B 1034 Samaru, Zaria, Kaduna State, Nigeria. Aug., 2018 Received: ²Department of Industrial Chemistry, Enugu State University of Science Accepted for Publication: Jan., 2019 and Technology, Agbani, Enugu State, Nigeria. *Author for Correspondence:bulus1973@gmail.com

Abstract

This paper reports the evaluation of heavy metal deposits and distribution in Challawa industrial area of Kano metropolis, using atomic absorption spectroscopy (AAS). The results showed that lead (Pb), Cobalt (Co), Zinc (Zn) and Iron (Fe), were present in an increasing order. The comparatively high iron content in both incinerated tannery wastes, and contaminated soils has high implication on plant growth and therefore be of great concern to soil conservationist for agricultural purposes.

Key Words: Incinerated waste, contaminated soils, tannery, and metals.

INTRODUCTION

could be hazardous to plants and animals and as (Khan et al. 1999). It has been estimated that such is a critical reason why soils are analyzed. during tanning at least approximately 300 kg The sources of such contamination could be chemicals are used per ton of hides or skins conjectured; however, the extent of the (Verheijin et al. 1996). Tannery effluent and solid contamination can only be ascertained by wastes constitutes the hazardous pollutants from modern instrumental analytical techniques the industry because of heavy metals, toxic (Montero et al. 2007). For example, during chemicals, chloride and much more. Several leather tanning and finishing processes analytical techniques such as inductively (Geremew and Tekalign, 2017). Heavy metals coupled plasma-atomic emission spectroscopysuch as Zinc, Iron, Lead and Cobalt may be ICP-AES, inductively coupled-mass involved, and therefore the extent of their spectroscopy-ICP-MS (Christoph et al. 2001; contamination to the environment need to be Montero et al. 2007; Achi et al. 2012). However, ascertained (Leke et al. 2011; Mohammad and in recent time atomic absorption spectroscopy Nwaedozie, 2011; Igwe and Abia, 2007), and (AAS), has proved to be the most versatile mixture of many compounds may complicate the instrument for heavy metal determination due to wastewater treatment such as: acids, alkalis, chromium salts, tannins (natural or synthetic), techniques (Gennaro et al. 2007). solvents, sulphides, dyes, surfactants, auxiliaries (Mariliz et al. 2014). Not more than 20 % of the impact of other heavy metal deposit and chemicals are absorbed by leather; the remainder flows out with the effluent (Muthukkauppan and Parthiban, 2018). Excess of lead in plant is known to inhibit photosynthesis and toxicity to soft tissues, such as liver, kidney and brain (Ogieva, 2003). Similarly, cobalt if found in

excess in animals could inhibit cellular respiration. The tanning industry is known to Heavy metal presence in an environment generate the highest toxic wastes per unit output its high sensitivity, low cost and simple

> The aim of this paper is to report the distribution in Challawa industrial areas in Kano municipality, by determining these metal pollutants other than chromium and copper in incinerated tannery wastes and contaminated soil samples.

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Evaluation of Heavy Metal Deposits and distribution

MATERIALS AND METHODS

Description of the Study Area.

Kumbotso Local Government area lies between latitudes 11°56'S to 12°N and longitude 8°24'W to 8°4'E it falls within the Kano State settlement zone bordering the south and west by Madobi local government area in the north-west: Rimingado in the north by Gwale and east by Tarauni local government areas respectively.

Sample Collection

Incinerated tannery wastes, and contaminated soil samples were collected randomly on weekly basis and were stored in plastic bags (Ayodele and Gaya, 1998) labeled A₁ $-Z_1$ and $A_2 - Z_n$ respectively. The soil samples were collected 500 meters away from the dump site at a depth of 15 cm from the surface. Total of blank for Zinc, Iron Lead and Cobalt 52 samples in each case were collected and concentrations using Atomic Absorption analyzed for Zinc (Zn), Iron (Fe), Lead (Pb) and Spectrophotometer (Buckley and Cranston, Cobalt (Co) using standard methods (Buckley 1993; Deepali and Gangwar, 2010). and Cranston, 1993; Balasubramanian et al. 1997).

Determination of Metals in Soil Samples

About 0.25 g of sample was digested with 10 cm³ hydrofluoric acid and 1.0 cm³ aqua regia, which is hydrochloric acid and nitric acid (3:1) in a flask. Thereafter, 5.0 cm³ perchloric acid was added and again heated to dryness on a hot plate; distilled water was added, filtered through Whatman, no 42-filter paper and made up to 100 cm³. Digested soil samples were analyzed in triplicates including the blank for Zinc, Iron, Lead and Cobalt concentrations using Atomic Absorption Spectrophotometer Alpha 4 Model (Deepali and Gangwar, 2010).

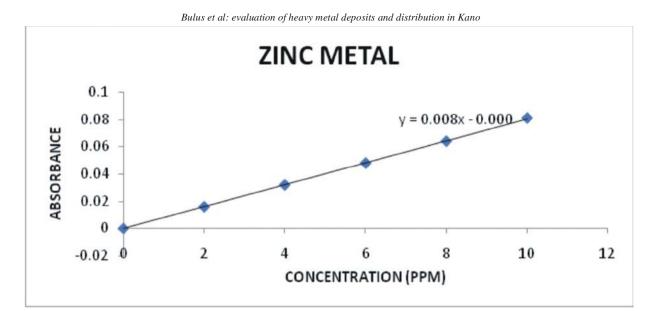
Determination of Metals in Incinerated Tannery Wastes

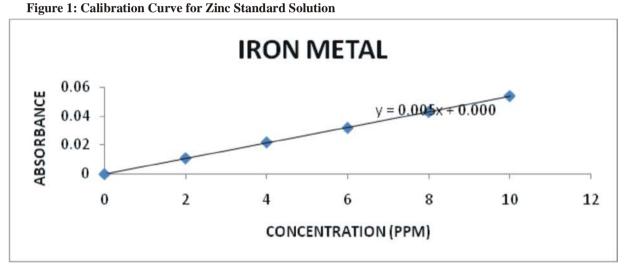
Burnt tanneries wastes from the sampling site were collected, crushed, mixed and sieved crucible and ashed at 400 °C (Gary, 2004) to a constant weight. The ash was quantitatively transferred from the platinum crucible using

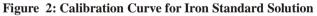
spatula into a 250 cm³ conical flask. A mixture of concentrated oxidizing acids of nitric (5 cm^3) , sulphuric (3.5 cm³) and perchloric (11.5 cm³) were transferred into conical flask containing the ash (John, 2003) antibumping granules and funnel was introduced into the conical flask. The mixture was heated on a hot plate in a fume cupboard. The initial colour of the solution varied from dirty green to orange (SLTC, 1996; Balasubramanian et al. 1997). The digestion was completed after $1\frac{1}{2}$ hours and the heating stopped and cooled at room temperature. 100 cm³ of distilled water was added into the solution and further boiled for 10minutes, cooled and filtered through a Whatman filter paper no (1) 90 mm into a 250 cm³ volumetric flask and was made up to the mark with water. The resultant solutions were analyzed in triplicates including the reagent

Preparation of Standard Solution

The following were prepared independently for their standard stock solutions as follows: Zinc nitrate (1.1375 g); Iron nitrate (1.8085 g); Lead (II) nitrate (0.3998 g); Cobalt nitrate (1.0082 g), transferred into a beaker containing 2.5 cm³ of 10 % nitric acid and made up to 250 cm³ with distilled water respectively. 2.5 cm³ of 250 mg/dm³ (standard stock solution) were prepared and made up to 25 cm³ mark with distilled water from the resultant solution 2.0. 4.0, 6.0, 8.0, 10.0. were pipetted and made up to 25 cm³ mark with distilled water each with new concentrations of 2.0, 4.0, 6.0, 8.0 and 10.0 mg/dm³ (Chemiasoft, 2011) for Zinc: Iron: Lead and Cobalt salts respectively (John, 2003). The absorbance of these serial dilutions was recorded. Average reading of both standards and samples were corrected from the blank readings. Calibration curves were plotted for their using 0.4 mm mesh for homogeneity. One gram standards. The concentrations of each element of each sample was weighed into platinum under investigation in mg/dm³ or mg/kg were determined from the calibration curve of its standard as presented in Figures 1.0-4.0.







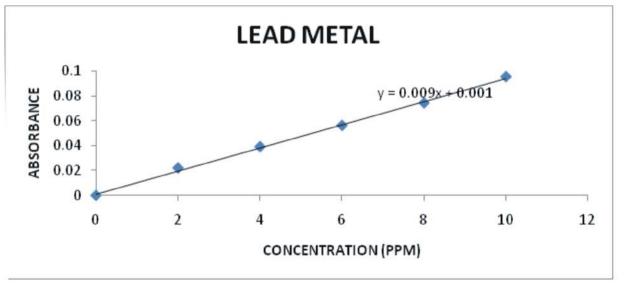


Figure 3: Calibration Curve for Lead Standard Solution

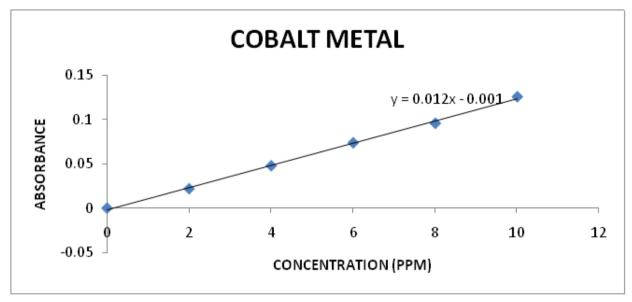


Figure 4: Calibration Curve for Cobalt Standard Solution

RESULTS AND DISCUSSION

in Figures 5 and 6, highlighting the metal carrots from farmland contaminated with tannery Concentrations of Zn, Fe, Pb and Co in Tannery waste could increase health risk. Incinerated Wastes and Contaminated Tannery Soil Sample (mg/kg).

Zinc

tannery wastes and farm soils were found to be The incinerated waste was above the standard 565.07 ± 225.71 and 251.20 ± 157.28 mg/kg limit for waste disposal but lower in the case of respectively. These values were below the set farm soils. These values were below 4,837-6,311 standard for sludge and soil contaminated with mg/kg range obtained from affected soils in tannery waste (Corning, 1979). Sources of Zinc Mexico as reported by Alvarez-Bernal et al. are: Animal skins 0.20% on dry basis (Adegboye (2006) Sources of iron in tannery waste are the and Agina, 1983) tanning salts, galvanized tanning agents because of their complexing materials or containers and pigments (Daintith, power with other substances such as (mineral 2004). The low concentration of Zinc could lead tannage), dyes other sources may include, natural to the formation of chlorophyll in plants because geochemical processes and machineries used ligand exchange processes in Zn^{2+} are usually which may include fleshing, setting out, shaving, rapid and does not disturb the genetic heredity staking and buffing (machines) in their and biosynthesis of enzymes (Sindhu, 2002) and production (Thainkaivelan et al. 2000; Lorede et activates of some enzymes and hormones to al.2003). Iron in high concentration could lead to enhance reproduction growth and development poor growth and development in plants (Ogieva, (Ogiera, 2003). Although Zinc is important in the 2003). Although iron is needed in haemoglobin body it becomes toxic above 15 mg/day and several enzymes, its bioaccumulation in the concentration due to bioaccumulation of the body in excess of 200 mg/day is considered toxic fumes from the burnt tannery waste the body for human (Butler, 1979; Bowen, 1979; through inhalation could lead to nausea and John,1986; Milacic et al. 1998; Sindhu, 2002). disorder of the immune system (John, 1986; Consequently, iron may be harmful to the body Ohnersorge and Wilhelm, 1991; Milacic et al.

1998; Sindhu, 2002) and consequent The results from this work are presented accumulation of the ingested fruits such as

Iron

The concentration for iron was found to be 1823.33±364.23 and 362.73±130.08 mg/kg The concentrations for Zinc in the for incinerated waste and farm soil respectively. cells when it catalyses the production of

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season when the leather wastes are burnt in soft tissues such as the liver, kidney and brain resulting in fumes causing poor visibility (Sezgin (Ademoroti, 1991). et al., 2004) with other related health problems. This is because iron bioaccumulates and does not **Cobalt** degrade naturally even when organic iron complexes breakdown by hydrolysis, fresh incinerated waste and farm soils are (Corning, 1984; Shen et al. 1986).

Lead

incinerated waste and contaminated soil samples cellular respiration and enzymes of the citric acid were found to be 77.88 ± 25.23 and 29.19 ± 14.61 cycle, exposure to cobalt carboxyl may change mg/kg respectively. These values were found to reflexes and electric activity of the brain. The be below the set standards (Corning, 1979). The toxicity of cobalt is independent of its chemical presence of lead could be attributed to the use of form as its toxic effects are contributed by pigment such as lead chromes and basic lead oxides, metals and are not masked by alloy carbonate (white lead) the later has been (Coates and Watson, 1971; Steve, 1972; Bowen, curtailed for toxicity consideration. Other 1979; Taylor and Hawkins, 1987). Cobalt could sources could be from domestic effluent surface be found in the composition of some tannery water run offs lubricating oil, grease (Hodel and pigment, skins of the animals about 0.04 % on the Chang, 2004). Lead toxic effect on plant is the dry weight of the skins (Adegboye and Agina, inhibition of photosynthesis due to direct 1983). Other natural sources may include; soil, interference with the synthesis of carbohydrate dust, volcanic eruption, fresh fires, burning coal which results in flaccidity of stomata guard cells and oil, cars, trucks and airplane exhaust and and impaired transpiration and CO_2 exchange. from industrial processes that use the metal or its Furthermore, lead gets into man through his diet compound (Steve, 1972).

hydroxyl radicals (Hueber, 1991) during the dry by inhalation and accumulates to cause toxicity

The concentration of cobalt in the unreactive inorganic complexes are formed 116.23 ± 58.11 and 43.07 ± 22.39 mg/kg respectively. The concentration of cobalt in the incinerated waste was above the set standard limit (Corning, 1979) which could pose some The data for lead concentrations in both related health problems such as inhibition of

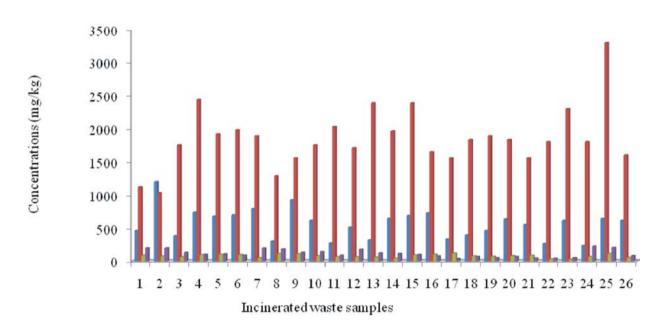
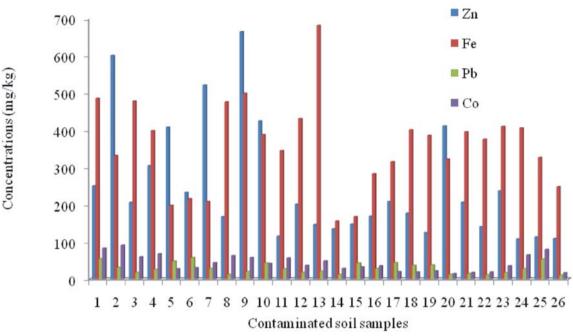


Figure 5: Concentration of Zn, Fe, Pb and Co in Tannery Incinerated Wastes (mg/kg)



Bulus et al: evaluation of heavy metal deposits and distribution in Kano

Figure 6: Concentrations of Zn, Fe, Pb and Co in Contaminated Tannery Soil Sample (mg/kg).

CONCLUSION

This study has shown that heavy metals in the tannery incinerated waste ranges from 77.88±25.23 (Pb), 116.23±58.11 (Co), 565.07±225.71 (Zn), 1823.33±364.23 (Fe) in mg/kg and for the contaminated soil sample it ranges from 29.19±14.61 (Pb), 43.07±22.39 (Co), 251.20±157.28 (Zn), 362.73±130.08 (Fe) in mg/kg respectively. The high concentration of iron in both parameters could be attributed to the use of iron salt for tanning in the industries with high percentages offered. Furthermore, some of the pigment dopes stinks after a long period of time when not properly preserved with biocides consequently ends in the dump site contributing to the concentration of these metals, to curtail this scenario a periodic monitoring of the incinerated wastes and contaminated soils should be embarked upon as a pollution index. The use of combination tannage with polyphenolic materials be encouraged and the composition of the pigments screened to check the bioaccumulation of heavy metals in the studied sites.

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