1	Original Research Article
2	Accumulation of Heavy Metals in Soil and Maize Plant (Zea mays) in the Vicinity of
3	Two Government Approved Dumpsites in Benin City, Nigeria
4	Abstract
5	Soil contamination by heavy metals is of great concern with respect to human health risks,
6	groundwater contamination, phytotoxicity to plants, adverse effects on microbial activity and
7	diversity, long-term effects on soil fertility and depreciation of land. Soil samples were
8	obtained with the aid of soil Augar within a depth of $0 - 20$ cm from the vicinity of the two
9	selected dumpsites in Benin City, Edo State Nigeria. The soil samples were assessed for some
10	physico-chemical properties using standard methods. Maize plants found growing in the
11	dumpsite and control areas were also sampled, partitioned into leaves, stems, and roots prior
12	to analysis in the laboratory for heavy metals determination using atomic absorption
13	spectrophotometer (AAS). The soil sample showed Zn, Cu, Cd, Pb and Cr levels ranging
14	from 42.66-243.81, 2.16-21.41, 0.35-2.59, 1.11-7.76 and 2.99-10.99 mg/kg respectively.
15	Pollution indices such as contamination factor, contamination degree and pollution load index
16	of the heavy metals analyzed were calculated and their values varied among the heavy metals
17	and between the dumpsites. Concentrations of the metals in the dumpsite soil and plant were
18	found to be higher compared to those of the control sites. Significant differences of heavy
19	metals accumulations were observed per plant parts, roots having the highest concentrations.
20	The translocation factor, biological concentration factor and biological accumulation
21	coefficient values of the plant species varied for all the metals. These results imply that the
22	dumpsites have associated human health and ecological risks.

23 Keywords: Dumpsites, Heavy Metals, translocation factor, Soil Maize (Zea mays).

24 Introduction

Dumpsites exist throughout developing countries. Most of these dumping sites are 25 uncontrolled and years old, having grown over time from small dumps to large, unmanaged 26 27 waste sites (Joan *et al.*, 2016). This constitutes serious health and environmental concerns because of the effects on the host soils, crops, animal and human health. Many cities in 28 29 Nigeria have developed without proper planning and it has led to the presence of open dumps 30 within built-up areas inhabited by millions of people (Amadi and Nwankwoala, 2013). 31 Consequently, such waste dumps become point source for soil pollution as they serve as host 32 for leachate from dumpsites. Composition of solid wastes in major cities in Nigeria comprises 33 domestic garbage, wood, agricultural waste, industrial waste, hospital waste, polythene bags, 34 plastics, broken glasses, abandoned automobiles, demolition waste, ash, dust, human and 35 animal waste (Awomeso *et al.*, 2010). The proper wastes disposal has been a serious problem 36 in Benin City and most cities in Nigeria. Solid and fluid wastes generation and their poor 37 disposal mechanism in the urban areas of most developing countries have become a threat to 38 the environment (Amadi et al., 2010). The contamination of soil with heavy metals is an 39 environmental concern because accumulated metals may have adverse effects on soil 40 ecology, agricultural production, animal and human health as well as groundwater quality 41 (Okiemen et al., 2011). While many heavy metals are essential elements at low levels of 42 concentration, they can exert toxic effects at concentrations higher than permitted in the 43 environment (Anegbe et al., 2014; Mmolawa et al., 2011). They may be volatilized to the 44 atmosphere, especially during dry seasons (Okuo and Okolo, 2011). In Benin City 45 metropolis, most of the dumpsites are used as fertile soils for the cultivation of some fruits, 46 food crops and vegetables due to the high cost of fertilizer. Some farmers collect the 47 decomposed parts of the dumpsites and apply to their farms as manure. These cultivated 48 plants take up these heavy metals either as mobile ion in the soil solution through their roots 49 or through their leaves thereby making it unfit for human consumption (Ganesh et al., 2010). 50 Recent studies have also reviewed that waste dumpsite can transfer significant levels of these 51 toxic and persistent metals into the soil environment. Eventually these metals are taken up by 52 plant part and transfer same into the food chain (Anhwange and Asemave, 2013). 53 Consequently, higher soil heavy metals concentration can result in higher levels of uptake by 54 plants. Although, the rate of metal uptake by crop plants could be influenced by factors such 55 as metal species, plants species, plant age, plant part soil composition, geographic and atmospheric conditions (Dulama et al., 2012). Transfer of heavy metal from soils to plants 56 57 has been proved as an efficient way for removal of these heavy metals through harvestable 58 plant parts such as roots, stems and leaves (Malik et al., 2010). However, the metal 59 availability and toxicity to plant can be determined by the soluble and exchangeable fraction 60 of metals in particular (OK et al., 2011). Intake of heavy metals via the crop-soil system has 61 been regarded as the predominant pathway of human exposure to toxic metals (Liu et al., 62 2007), and is normally chronic. This is because these metals are non-biodegradable and can 63 undergo global ecological circles (Opaluwa et al., 2012). Thus, it become necessary to assess 64 the uptake of heavy metals by maize plant (zea mays) in two government approved dumpsites in Benin City, Nigeria in order to determine their potential hazards to human beings and 65 66 animals. The objective of this study was to determine and compare the content of heavy metals in various parts of the maize plant, namely the leaves, stems and roots compared to the 67 68 levels in the soil around Oluku and Ikhueniro dumpsite zones.

69 Materials and Methods

70 Study Area

71 Ikhueniro dumpsite is the largest and the major open dumpsite site in Benin metropolis. It has

been in operation as a disposal facility; permitted to receive commercial and municipal solid

vaste. However, the absence of waste management and sorting systems lead to the dumping

of industrial waste into the dumpsite (Ighodaro *et al.*, 2015). Ikhueniro and Oluku dumpsites

- 75 comprise household materials, hospitals disposables, metals scraps, polyethylene bags and
- 76 papers, plants materials and debris among other substances. They also consist of scavengers
- 77 that are involved in sorting some of these materials for re-us





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81 Collection of Soil Samples

Soil and plant samples were collected from two government approved dumpsites and their control sites within Benin City, namely Oluku dumpsites (A), Oluku control site (B), Ikhueniro dumpsites (C) and Ikhueniro control site (D). In this research, Soil samples at 0- 20 cm depth from rhizosphere of the maize plant were taken from each site from where plant sample was rooted. At each site, three different points were chosen using cluster random 87 sampling technique to collect the sample. The soil samples were obtained by the use of soil 88 auger, and then blended (mixed) to obtain a representative sample. The samples were air 89 dried and ground to pass through a 2 mm sieve and used for both physico-chemical analysis 90 (Anegbe and Okuo, 2013). Prior to the analysis of plant material, leaf, stem and roots of 91 plants were separated and carefully washed with tap and deionized water in order to remove soil or dust deposits. Then the plant samples were oven-dried at 70°C to constant weight, 92 93 pulverized, passed through 2 mm steel sieve and weighed in order to determine the heavy metals concentrations by atomic absorption spectrometry (Tsvetomil et al., 2013). The 94 95 control soil and plant samples were taken at about 1500 m away from each of the dumpsites. 96

97 Analysis of the Soil and Plant Samples

98 The pH and the CEC were determined as described by Anegbe and Okuo (2013). The hydrometer method described by Ugbune and Okuo (2011) were used in evaluating the 99 100 particle size. The method described by Anegbe et al. (2017) were used to determine the organic carbon content, while the total heavy metals determination was carried out using the 101 102 Tessier's method described by Okuo *et al.* (2016). According to the method, 5 ml of aqua regia (BDH, England) and 1 ml of perchloric acid (BDH, England) were added to 1 g of soil 103 sample in a 150 ml digestion tube and digested on a heating digester until white fumes of 104 perchloric acid appeared. The tube was cooled and the sides rinsed with distilled water and 105 106 then filtered through a Whatman 1 filter paper into a 100 ml volumetric flask. The volume 107 was made up with distilled water All glasswares used were soaked and washed with chromic 108 acid and rinsed with distilled water. Bulk scientific standard solution was used to calibrate the 109 Atomic Absorption Spectrometer (Pg A500 model, USA). Procedural blank samples were 110 subjected to similar extraction method using the same amount of reagents.

111 Results and Discussion

112 The physico-chemical properties of the soil samples at various sites are shown in Table 1.

113 Soil pH plays a major function in the sorption of heavy metals as it directly controls the

- solubility and hydrolysis of metal hydroxides, carbonates and phosphates (Tokalioglu et al.,
- 115 2006). The pH of the studied areas ranged from neutral to moderately alkaline. Most mineral

- and nutrients are more soluble or available in acidic soils than in neutral or slightly alkaline
- soils. Soils tend to become acidic as a result of rain water leaching away basic ions $(Ca^{2+},$

118 Mg^{2+} , K⁺ and Na⁺) (Bickelhaupt, 2015).

119 Table 1: Physico-chemical Properties and Total Heavy Metal Concentration of the Soil

120 Samples.

Parameters	Units	Dumpsite soil (Oluku)	Control soil (Oluku)	Dumpsite soil (Ikhueniro)	Control soil (Ikhueniro)
pH		8.10±0.01	7.14±0.02	7.40±0.00	7.05±0.02
CEC	meq/100g	12.82±1.22	5.22±0.15	11.21±0.30	5.20±0.22
TOC	%	1.90±0.26	1.56±0.14	1.74±0.15	1.23±0.30
OM	%	3.28±0.26	2.69 ±0.14	3.00±0.15	2.12±0.30
Clay	%	10.71±0.12	16.71±0.22	28.24±0.40	31.72±0.30
Silt	%	2.09±0.11	2.09±0.10	2.56±0.15	2.49±0.17
Sand	%	87.20 ±0.35	81.20±0.50	69.20±0.31	65.79±0.25
Zn	mg/kg	156.78±2.35	58.93±2.07	243.81±5.00	42.66±2.00
Cu	mg/kg	20.17±1.44	2.30±0.24	21.41±1.71	2.16±0.22
Cd	mg/kg	1.40±0.19	0.35±0.08	2.59±0.42	0.73±0.10
Pb	mg/kg	6.36±0.25	1.32±0.56	7.76±0.55	1.11±0.05
Cr	mg/kg	7.92±1.12	4.68±0.50	10.99±0.64	2.99±0.23

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122 The CEC parameter particularly measures the ability of soils to allow for easy exchange of 123 cations between its surface and solution. The soil samples from the two dumpsites and their 124 controls shows a low CEC which indicates that they are more likely to develop deficiencies in potassium (K^+), magnesium (Mg^{2+}) and other cations, while high CEC soils are less 125 126 susceptible to leaching of these cations (CUCE, 2007). The low values of the CEC were 127 attributed to high sandy nature of the soil samples. Textural analysis showed the 128 preponderance of sand fraction, followed by clay then silt, thus classifying the parent soil as 129 loamy sand. Sandy soils are known to have a poor retention capacity for both water and metals (Wuana et al., 2010). Low organic matter (2.12 – 3.28 %) was observed in all the soil 130 131 samples. The low organic matter content of the dumpsite and the control soil samples is an

indication that these soils will have low adsorption strength and an increased metal mobilityand bioavailability (Udeigwe, 2010).

134 The soil samples showed the presence of Zn, Cu, Cd, Pb and Cr in all the sites analysed This 135 could be attributed to the availability of wastes containing those metals at the dumpsites 136 which are eventually leached into the underlying soils. The concentrations of heavy metals 137 are higher in each of the dumpsites than their respective control sites. Liu et al. (2007), 138 observed that heavy metal concentration in soils is usually high near the sources, and decline with both distance and depth due to physical dilution and increasing limits in mobility. In 139 140 overall terms, the results of the present study suggested that the five metals decline in the 141 following order: Zn>Cu>Cr>Pb>Cd for each of the two sites. The results of heavy metals 142 obtained from the analysis also indicate that the concentrations of the heavy metals were 143 found to be higher in Ikhueniro dumpsite when compared with Oluku dumpsite. The higher 144 concentration could be as a result of waste carrying more concentrations of these metals in 145 Ikhueniro dumpsite compared to that of Oluku dumpsite.

146 Assessment of Metal Contamination

147 Contamination Factor (CF)

- 148 The level of contamination of soil by metal is expressed in terms of a contamination factor
- 149 (CF) calculated as:

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$$CF = \frac{Cm Sample}{Cm Background}$$
 (1)

151 Cm Sample = metal concentration in Sample

152 Cm Background = metal concentration in background or control Sample. (Lin *et al.*, 2009)

Where the contamination factor CF < 1 refers to low contamination; $1 \le CF < 3$ means moderate contamination; $3 \le CF \le 6$ indicates considerable contamination and CF > 6indicates very high contamination.

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Metals	Oluku Dumpsite	Oluku Classes	Ikhueniro Dumpsite	Ikhueniro Classes
Zn	2.66	Moderate contamination	5.72	Considerable contamination
Cu	8.77	Very high contamination	9.91	Very high contamination
Cd	4.00	Considerable contamination	3.55	Considerable contamination
Pb	4.82	Considerable contamination	6.99	Very high contamination
Cr	1.69	Moderate contamination	3.68	Considerable contamination

161 Table 2: Contamination Factor of Each Metal in the Two Dumpsites

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163 From the results of the contamination factors, the soil samples may be classified as 164 moderately contaminated with respect to Zn and Cr, considerably contaminated with respect 165 to Cd and Pb, and very highly contaminated with respect to Cu in the vicinity of Oluku 166 dumpsite. The soil in the vicinity of Ikhueniro dumpsite may be classified as considerably contaminated with respect to Zn, Cd and Cr, and very highly contaminated with respect to Cu 167 168 and Pb. It was also observed that the contamination factor of each metal in the vicinity of Ikhueniro dumpsite is greater than the contamination factor of the same metal in Oluku 169 170 dumpsite except for cadmium which proved otherwise. This is because contamination factor 171 is directly proportional to the concentration of each metal in the sediment, and all the metals 172 in the vicinity of Ikhueniro dumpsite except cadmium have higher sediment concentrations 173 than their counterpart in Oluku dumpsite.

174 Degree of Contamination

The sum of contamination factors for all elements examined represents the contamination
degree (Cdeg) of the environment. Using the sum of contamination factors obtained in table 2
for all elements in each dumpsite.

The contamination degree (Cdeg) values were 21.94 and 29.84 for Oluku and Ikhueniro dumpsites respectively. Therefore, the soils in the vicinity of the two dumpsites where the plant samples were gotten can be classified as having considerable degree of contamination according to Dasaram *et al.* (2011). However, it is worthy of note that the degree of contamination of soil in the vicinity of Ikhueniro dumpsite is greater than that of Oluku dumpsite; this might be attributed to the higher concentration, and hence higher contamination factor of each metal at the vicinity of Ikhueniro dumpsite than that of Oluku dumpsite.

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187 The Pollution Load Index (PLI)

188 Generally, pollution load index (PLI) as reported by Harikumar *et al.* (2009), is as follows:

189 PLI =
$$\sqrt[n]{Cf1 x Cf2 x Cf3 x Cf4 Cfn}$$
 (2)

190 Where, CF = contamination factor, n = number of metals

191 The PLI value of > 1 is polluted, whereas < 1 indicates no pollution (Harikumar *et al.*, 2009).

The PLI values as calculated for both sites were 3.77 and 5.53 for Oluku and Ikhueniro dumpsites respectively. This showed a strong sign of pollution deterioration by the five measured metals in both sites since their PLI values are greater than 1. The PLI value was higher in the vicinity of Ikhueniro dumpsite than that of Oluku dumpsite. Hence, the vicinity of Ikhueniro dumpsite may cause more pollution to the environment than Oluku dumpsite.

197 Uptake of Heavy Metals by Plants in the Vicinity of the Open Dumpsites in Benin City

Although certain trace elements are essential in plant physiology, plants growing in a polluted
environment can bioaccumulate trace elements at very high concentration in their edible parts
and can present public health concerns (Quierolo, *et al.*, 2000).

Table 3: Uptake of Zn, Cu, Cd, Pb and Cr (mg/kg) by Different Parts of the Maize Plants Collected from Oluku Dumpsite and its Control

Metals	Roots	Stems	Leaves	Shoot
Zn	62.30(17.64)	25.14(7.36)	7.37(3.11)	32.51(10.47)
Cu	9.50(0.78)	3.21(0.36)	1.98(<0.05)	5.19(0.36)
Cd	0.62(0.10)	0.21(<0.05)	0.26(0.09)	0.47(0.09)

Pb	2.05(0.32)	0.62(<0.05)	0.35(<0.05)	0.97(0.00)
Cr	3.08(0.90)	2.15(0.72)	1.07(0.54)	3.22(1.26)

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Table 4: Uptake of Zn, Cu, Cd, Pb and Cr (mg/kg) by Different Parts of the Maize Plants Collected from Ikhueniro Dumpsite and its Control. 205

Metals	Roots	Stems	Leaves	Shoot
Zn	89.55(13.83)	40.94(5.79)	22.87(3.88)	63.81(9.67)
Cu	10.30(0.44)	4.98(0.26)	1.08(<0.05)	6.06(0.26)
Cd	0.96(0.19)	0.54(0.09)	0.60(0.15)	1.14(0.24)
Pb	2.62(0.21)	1.05(0.10)	0.26(<0.05)	1.31(0.10)
Cr	5.07(0.72)	2.02(0.29)	1.77(<0.05)	3.79(0.29)

206 **Key**: Shoot = Stem + Leaf. Note: the values in bracket are for control.

207 Among all plant parts, roots accumulated the highest metal contents in all the sites (Table 4). 208 Highest uptake of metals in roots compared to other parts was also observed in similar studies 209 (Sekara et al., 2005). This could be due to roots direct exposure to the contaminated soil. It 210 was also found that plant parts from the dumpsites have higher metal concentrations than control sites. This might be attributed to higher concentrations of metals in the dumpsite than 211 the control site (Amusan et al., 2005; Li et al., 2007). The results also indicated that the levels 212 213 of metals in plants are dependent upon their concentrations in their habitual soil environment 214 (Ayari et al., 2010; Malik et al., 2010).

Transferability of Metals. 215

Biological Accumulation Coefficient (BAC) =
$$\frac{[Metals] \text{ shoot}}{[Metals] \text{ soil}}$$
 (3) (Li *et al.*, 2007).
Biological Concentration Factor (BCF) = $\frac{[Metals] \text{ root}}{[Metals] \text{ soil}}$ (4) (Yoon *et al.*, 2006).

218 Translocation Factor (TF) =
$$\frac{[Metals] \text{ shoot}}{[Metals] \text{ root}}$$
(5) (Cui *et al.*, 2007).

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Metals	Biological Accumulation Coefficient (BAC)	Biological Concentration Factor (BCF)	Translocation Factor (TF)
Zn	0.21(0.18)	0.40(0.30)	0.52(0.59)
Cu	0.26(0.16)	0.47(0.34)	0.55(0.46)
Cd	0.34(0.26)	0.44(0.29)	0.76(0.90)
Pb	0.15(0.00)	0.32(0.24)	0.47(0.00)
Cr	0.41(0.27)	0.39(0.19)	1.05(1.40)

Note: the values in bracket are for control

220 Table 5: BAC, BCF and TF for Zn, Cu, Cd, Pb and Cr in the Maize Plant at Oluku

223 For Oluku dumpsite, the biological accumulation coefficient (BAC) decreased in the 224 following order Cr>Cd>Cu>Zn>Pb, the biological concentration factor (BCF) decreased as 225 follows Cu>Cd> Zn>Cr>Pb while the translocation factor (TF) followed the order of 226 Cr>Cd>Cu>Zn>Pb. For Oluku control site, the BAC decreased in the following order 227 Cr>Cd>Zn>Cu>Pb, the BCF decreased as follows Cu > Zn >Cd >Pb>Cr while the TF 228 followed the order of Cr>Cd>Zn>Cu>Pb. Pb was not detected in the various plant parts but 229 occurred in low concentrations in the roots of the maize plant. The TF of the maize plant is 230 greater than 1 with respect to chromium shows the special ability of the maize plant to absorb 231 chromium from soils and transport and store it in its above-ground part (Wei et al., 2002). 232 BAC was categorised as: < 1 excluder, 1-10 accumulator and > 10 hyperaccumulator (Ma 233 et al., 2001). Hence, using the results obtained for BAC in table above, it could be 234 suggested that the maize plant is an excluder with respect to all the heavy metals analysed in 235 Oluku dumpsite and its control site because all the BAC values were less than 1. TF > 1 236 signifies that the plant effectively translocates heavy metals from roots to the shoots (Baker 237 and Brooks, 1989). Hence it could be observed from Tables 5 that the maize plant effectively 238 translocate chromium from roots to the shoots since the TF of the maize with respect to 239 chromium is greater than 1 both in Oluku dumpsite and its control site.

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Dumpsite and its Control.					
lls Biol Coef	ogical Accumulation ficient (BAC)	Biological Concentration Factor (BCF)	Translocation Factor (TF)		
0.26	(0.23)	0.37(0.32)	0.71(0.70)		
0.28	(0.12)	0.48(0.20)	0.59(0.59)		
0.44	(0.33)	0.37(0.26)	1.19(1.26)		
0.17	(0.09)	0.34(0.19)	0.50(0.48)		
0.34	(0.10)	0.46(0.24)	0.75(0.40)		
	Dumpsite and Is Biol Coef 0.266 0.286 0.446 0.176 0.346	Dumpsite and its Control. Is Biological Accumulation Coefficient (BAC) 0.26(0.23) 0.28(0.12) 0.44(0.33) 0.17(0.09) 0.34(0.10) 0.34(0.10)	Biological Accumulation Coefficient (BAC) Biological Concentration Factor (BCF) 0.26(0.23) 0.37(0.32) 0.28(0.12) 0.48(0.20) 0.44(0.33) 0.37(0.26) 0.17(0.09) 0.34(0.19) 0.34(0.10) 0.46(0.24)		

245 **Table 6: BAC, BCF and TF for Zn, Cu, Cd, Pb and Cr in the Maize Plant at Ikhueniro**

247 **Note**: the values in bracket are for control.

248 For Ikhueniro dumpsite, the biological accumulation coefficient (BAC) decreased in the 249 following order Cd> Cr>Cu>Zn>Pb, the biological concentration factor (BCF) decreased as 250 follows Cu> Cr>Cd= Zn> Pb while the translocation factor (TF) followed the order of 251 Cd>Cr> Zn>Cu> Pb. For Ikhueniro control site, the BAC decreased in the following order 252 Cd>Zn>Cu> Cr>Pb, the BCF decreased as follows Zn >Cd > Cr >Cu>Pb while the TF 253 followed the order of Cd>Zn>Cu>Pb >Cr. Using the results obtained for BAC in table 6, it 254 could be suggested that the maize plant is an excluder with respect to all the heavy metals 255 analysed in Ikhueniro dumpsite and its control site (Ma et al., 2001). Also considering the TF 256 values in table 6, it could be observed that the maize plant effectively translocate cadmium 257 from roots to the shoots since the TF of the maize with respect to cadmium is greater than 1 258 (Baker and Brooks, 1989). High accumulation of heavy metals in roots and low translocation 259 in shoots may indicate appropriateness of a plant species for phytostabilisation (Archer and 260 Caiwell, 2004; Malik et al., 2010). Phyto-stabilization process depends on roots' ability to 261 limit the heavy metals' mobility and bioavailability in the soils and these occurs through 262 sorption, precipitation, complexation or metal valance reduction (Ghosh and Singh, 2005). 263 High root to shoot translocation of metals indicate that the plants have vital characteristics to 264 be used in phytoextraction of the metals (Malik et al., 2010).

265 Conclusion

The two dumpsites studied in Benin City had revealed that indiscriminate disposal of wastes such as municipal wastes, industrial wastes, agricultural wastes, etc are major sources of soil contamination and pollution by heavy metals. A knowledge of the total concentration of these heavy metals through soil analysis (as indicator) could be considered as a starting point for evaluating the degree of pollution as investigated in this study. All of the heavy metals

271 studied were found to accumulate mainly in the roots of the maize plant. It can be concluded 272 that Ikhueniro dumpsite area is more polluted than Oluku dumpsite area and that the 273 indiscriminate disposal of wastes on these land had contributed to the increment in 274 concentrations of these metals in the land. All of the heavy metals studied were found to 275 accumulate mainly in the roots of the maize plant. Linearity dependence was found between 276 the total heavy metal content in the soil and in the plant for all the elements studied. This may 277 suggest that, plant absorption is controlled by the content of heavy metals in the soil solution 278 and also by the content that is bioavailable in the soil.

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