

Edta, Water-Extractable Arsenic and Bioavailability of Arsenic in Soil/Tailings: the Effects of Zeolite and Beringite Amendments

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Abstract

The effects of zeolite and beringite (0, 30 and 60 kg ha⁻¹ w/w) on EDTA-extractable Arsenic (As), water soluble As, and bioavailability of arsenic (As) in non-mined agricultural soil and tailings of a mined site(capped and uncapped) were studied in a pot experiment. Un-amended non-mined agricultural soil was included as a control. EDTA-extractable and water extractable forms of As were monitored for 0-12-weeks after amendments and bioavailability of As was evaluated by growing lettuce (*Lactuca sativa*) on amended tailings/soil for 4-12 weeks after incubation. For EDTA-extractable As at week 12, beringite amendment (60 kg ha⁻¹ (w/w) was 29 %, 8.6 % and 26 % more efficient over zeolite for soil/capped and uncapped tailings respectively. For water-extractable As at week 12, beringite amendment 60 kg ha⁻¹ (w/w) again recorded 84 %, 54.5 % and 72 % less extractable As over zeolite in soil/capped and uncapped tailings respectively. Analysis for bioavailable-As in edible leafy aerial parts of lettuce, indicated 52.5 %, 82.0 % and 39.0 % less extractable As for beringte over zeolite in capped and uncapped tailings (week 12, 60 kg ha⁻¹ (w/w)... Beringite (week 12, 60 kg ha⁻¹ w/w) reduced As of capped tailing (0.3 mg kg⁻¹) below the FAO & WHO recommended maximum limit of 1.0 mg kg⁻¹ for human consumption.

Keywords: Immobilization, solubility, mobility, remediated soil, Lactuca sativa



1. Introduction

Arsenic (As) is a frequently found contaminant in contaminated sites. This is due to its many industrial applications. Owing to its toxic nature, soil contamination by arsenic has always been a major environmental concern. Apart from its ability to bio-accumulate in edible plant parts and enter the food chain, arsenic contaminated soils could be transported by wind and run-off into community water sources. Exposure to arsenic has been linked to a variety of cancers, cardiovascular disease, diabetes, and anemia, as well as having reproductive, developmental, immunological, and neurological effects (US EPA, 2003). Methods used to remediate arsenic contaminated soils include phytoremediation, solidification/stabilization, fixation, vitrification, soil flushing, and electro kinetic remediation (Date, Terakado, Sasaki, Aota, Matsumoto, Shiku, Inoa, Watanabe, Matsue & Ohmura, 2012; Jiang, Tao & Liao, 2011). A common objective of these methods is to reduce the hazard potential of the waste by limiting its solubility and mobility. Chemical stabilization has received wide patronage for arsenic-contaminated soil in recent years, probably because of its cost effectiveness. The conventional soil remediation technologies, although suitable for relatively small and heavily contaminated areas, are financially and physically inefficient in extensive and moderately contaminated soils. The capping of contaminated tailings with lateritic soil as practiced by some mining companies in Ghana does not remove arsenic from the soil/tailings. Amendments that have been screened for their potential to immobilize heavy metals in soils are alkaline materials such as beringite and zeolite (Edwards, Rebedea, Lepp, & Lovell, 1999.; Oste, Lexmond, & Van Riemsdijk, 2002; Mench, Vansgronsveld, Beckx, & Ruttens, 2006).

The bioavailability of heavy metals in mediated soils can be assessed by growing plants such as lettuce that bio-accumulate substantial amounts of these metals in their edible aerial parts (Nadia & Zaghlou, 2007). The objective of this study therefore was to assess the effectiveness of zeolite and beringite on the forms and bioavailability of arsenic (As) in non-mined agricultural soil, capped and uncapped tailings of a mined site.

2. Materials and Methods

Soil and tailings used in this study were sampled from a mining site in Ghana. These were reclaimed tailings (capped and uncapped), and a non-mined agricultural soil (control). Top soils/tailing samples (0-15 cm depth) were collected. Forty soil cores each were taken for non-mined, capped tailings and uncapped tailings, bulked into three composite samples and transported to the laboratory. The samples were prepared by air drying, crushing, discarding of foreign material and sieving through a 2-mm nylon mesh. The properties of soil/tailings are presented in Table 1.



Table 1. Properties of soil/tailings at 0 - 15 cm depth

Properties	None M	ined Capped Tailings	Uncapped Tailings
Organic carbon (%)	2.13	0.63	1.02
Total N (%)	0.19	0.06	0.10
C:N ratio	11.21	10.50	10.20
pH (1:2.5 soil:water)	5.10	6.79	7.22
P - Bray I (mg kg ⁻¹)	6.58	16.57	1.53
Ca ⁺² (cmol _c kg ⁻¹)	3.47	11.75	15.39
$\mathrm{Mg}^{+2} \left(\mathrm{cmol}_{\mathrm{c}} \mathrm{kg}^{-1} \right)$	1.60	8.54	9.61
K^+ (cmol _c kg ⁻¹)	0.21	0.42	1.40
Na ⁺ (cmol _c kg ⁻¹)	0.13	0.42	1.67
CEC (cmol _c kg ⁻¹)	5.41	21.13	28.07
Exhange Acidity			
$(\text{cmol}_{\text{c}} \text{ kg}^{-1})$	0.35	0.10	0.05
Effective CEC			
(cmocl _c kg ⁻¹)	5.76	21.23	28.12
Bulk density (g cm ⁻³)	1.26	1.33	1.39
Total Porosity (%)	52.5	49.8	47.5
Sand (%)	44	49	35
Silt (%)	46	41	61
Clay (%)	10	10	4

2.1 Soil Amendment and Incubation Experiment

Five hundred grams (500 g) air dry soil/tailings from each of the three sites were placed in 0.5 L plastic containers. The soil/tailings were amended with zeolite and beringite at the rates of 0, 30 and 60 kg ha⁻¹ (w/w). The amendments were thoroughly incorporated into soil/tailings and moistened to field capacity. Each amendment was replicated there times. The completely randomized design was used. The samples were incubated at room temperature of 25 °C in the laboratory and moistened weekly to maintain soil moisture at 60 % field capacity. The soil/tailings were sampled at 0, 4, 8 and 12 weekly intervals, mixing thoroughly before sampling. Five random sub-samples were taken with a spatula from each container and bulked for analysis.



2.2 Extraction of (EDTA) – As

Ten (10) grams of air-dried soil/tailings were weighed and pulverized in a clean, nitric acid washed mortar and pestle, for the Ammonium tetra sodium-ethylene-diamine-tetraacetic acid (Na₄-NH₄-EDTA) extraction (ADAS, 1986; Sabiene, Brazuaskiene & Rimmer, 2004). The soil was wrapped in a paper towel before being pulverized with the mortar and pestle to minimize the nitric acid washing necessary for the mortar and pestle. The soil was placed into a 100 ml conical flask and 50 ml of 0.05 M (Na₄-NH₄-EDTA) (pH 7) was added. The flask with the soil: Na₄-NH₄-EDTA mixture was shaken at room temperature on a mechanical shaker at 125 rpm for 1 hr. (ADAS, 1986; Sabiene, Brazuaskiene & Rimmer, 2004). The slurry was then passed through a Whatman's no. 42 filter paper into a sterile 100 ml conical flask. The filtrate was brought to exactly 100 ml with the extraction solution. The filtrate was analyzed using the Atomic Absorption Spectrophotometer (AAS). The concentrations of As was measured at wavelengths of 193.7 nm.

2.3 Extraction of Water Soluble As

Water soluble As was determined by weighing 5.0 g of soil/tailings samples into centrifuge tubes and 50 ml of distilled water added. The tubes were capped and shaken for 30 min. After shaking, the content of the tube was filtered with Whatman no.42 filter paper. Arsenic levels in the extracts were determined by AAS at a wavelength of 193.7 nm.

2.4 Bioavailability Experiment and Plant Analysis

Lettuce (*Lactuca sativa*) was chosen as the plant indicator of the bioavailability of As for its ability to accumulate heavy metals in the edible leafy aerial parts (Nadia & Zaghloul, 2007). The field practices for the cultivation of lettuce as described by (Grubben, 2004) were followed. The lettuce seeds were planted in a seedbed that was shaded with oil palm fronds. The seedlings were pricked out three weeks after germination and planted at a spacing of 4 cm × 4 cm. The seedlings were natured for another three weeks and subsequently transplanted into the containers/pots containing the mediated non-mined soil/tailings which were incubated for 3 weeks. The soil/tailings samples were then moistened to 60% of field capacity and then moistened weekly to maintain moisture at 60% field capacity. Fertilizer was incorporated into the soil/tailings samples at the rates of 50 kg N ha⁻¹, 45 kg P ha⁻¹ and 65 kg K ha⁻¹ at the time of transplanting. Three-weeks after transplanting, 50 kg N ha⁻¹ was applied as recommended by (Grubben, 2004). The aerial parts of the lettuce plants were harvested and prepared for the analysis for Arsenic. One (1.0) g of prepared plant material was digested in freshly prepared mixture of HNO₃-HCl (1:3 v/v) on a digester at 110 °C for three hours to determine the concentration of As.

2.5 Data Analysis

The GenStat statistical package (GenStat, 2016) was used for the analysis of data. The analysis of variance (ANOVA) was used to determine the effects of the amendments on water-soluble and EDTA-extractable As in the non-mined soil, capped and uncapped tailings. The least significant difference (LSD) was used for the separation of means (95 % confidence level). The percentage change was used to explain the differences in the effects of the



amendments. The percentage change in treatment effect on the value of the variable (arsenic) was estimated from the relationship:

%
$$\Delta Varible = 100 \left(\frac{V_{\times Rate} - V_0}{V_0} \right)$$

The percentage change in value of variable at given rate of application of amendment is represented by % Δ Variable, the value of the variable at a specific rate of application is given by V_X Rate and the value of the variable at 0 kg ha-1 rate of application is represented by V_0 .

3. Results and Discussion

3.1 Effects of Amendments on EDTA-extractable Arsenic in Non-Mined Soil, Capped and Uncapped Tailings

The results of EDTA-extractable arsenic (As), in non-mined soil, capped and uncapped tailings following beringite and zeolite amendment are presented in tables 2, 3 and 4. At 4 weeks after incubation (WAI), there were significant (P = 0.05) differences in EDTA-As between beringite amendments (30 and 60 kg ha⁻¹) and the control in non – mined, capped and uncapped tailings. The concentration of EDTA-As was significantly lower when beringite was applied at 60 kg ha⁻¹(15.46 mg kg⁻¹) than at 30 kg ha⁻¹ (17.02 mg kg⁻¹) and decreased by 21.8 % and 13.9 %, respectively compared to the non-amended non-mined soil at 4 WAI. The concentration of EDTA-As, when beringite was applied to capped tailings at 60 kg ha⁻¹(135.3 mg kg⁻¹) and 30 kg ha⁻¹ (160.7 mg kg⁻¹) decreased by 24.5% and 10.3 %, respectively compared to the non-amended soil at 4 WAI (Table 3). A similar trend was observed for beringite-amended uncapped tailings at 4 WAI (Table 4). Generally, the concentrations of EDTA-As beringite applied at the rates of 30 and 60 kg ha⁻¹ were significantly lower than (P = 0.05) that of non-amended non-mined soil, capped and uncapped tailings from 8 WAI to 12 WAI (Tables 2, 3 and 4). These results indicated that the exchangeable and strongly bonded fractions of arsenic in non-mined soil, capped and uncapped tailings that were extracted with 0.05 M EDTA buffered at pH 7 decreased with the application of beringite at 4, 8 and 12 WAI (Sabiene, Brazuaskiene & Rimmer, 2004; Xu, Xie & Xue, 2011; Garau, Castaldi, Santona, Deiana, & Melis, 2007). The extracting solution, Ammonium-EDTA (0.05 M, pH 7) extracted significantly lower concentrations of As compared to un-amended non-mined soil capped and uncapped tailings due to the formation of As complexes that are strongly bonded in the inner sphere of beringite (Pacwa-Plocini, Plaza, Piotrowska-Seget & Cameotra, 2011; Rascio & Navari-Izzo, 2011). The complexation of As to form calcium arsenate occurs in the inner sphere of berringite that is a constituent of beringite. Furthermore, the adsorption of As due to the presence of Ca2+ and Mg2+ ions in beringite contributes to a decrease in extractable As in contaminated soil (Bhargava, Carmona, Bhargava & Srivastava., 2012; Trgo, Peric & Vukojevic-Medvdovic, 2006; Shazia, Alia, Iftikhar, & Muhammad, 2015). The presence of bivalent cations, in berringite that is a constituent of beringite, aids in the adsorption of As which is not easily desorbed (Pubmed, Elena, Elizabeth, & Rosanna, 2011; Valeska, Elena, Luz, Michel, Alexander, & Rosanna.,



2012; Rajindiran, Dotaniya, Vassanda Coumar, Panwar, & Saha, 2015). The adsorption and complexation of As through the application of beringite could therefore be the major factors contributing to the concentration of As-EDTA compared to the control. A decrease of 34 % in extractable arsenic in contaminated soil has similarly been obtained and attributed to the presence of bivalent cations (Ca²⁺ and Mg²⁺) in beringite that aid in the adsorption of arsenic (Ruttens & Boulet, 2011; Saha, 2014).

The rate of application of zeolite resulted in an insignificant (P = 0.05) concentration of EDTA-extractable arsenic compared to un-amended non-mined soil when applied at 4 WAI (Table 5). Also, at 8 and 12 WAI, the concentrations of EDTA-As were similar (P = 0.05) in zeolite-amended non-mined soil at rates 30 and 60 kg ha⁻¹ with values in the ranges of 19.55 mg kg ⁻¹ to 19.62 mg kg ⁻¹ and 19.41 mg kg ⁻¹ to 19.48 mg kg ⁻¹, respectively but significantly lower (P = 0.05) than non-amended soil (Table 3).

Table 2. Effects of rate of application and type of amendment on EDTA- extractable arsenic of non-mined soil as affected by incubation period

Amendments	Rate of application (kg ha ⁻)		Weeks after incubation			
	(kg nu)		4	8	12	
			EDTA	EDTA –extractable As (mg kg ⁻)		
Zeolite	0	20.01		19.95	20.04	
	30	19.73		19.62	19.55	
	60	19.65		19.48	19.41	
Beringite	0	19.76		19.79	20.20	
· ·	30	17.02		16.35	16.18	
				14.72	14.34	

Table 3. Effects of rate of application and type of amendment on EDTAextractable arsenic of capped tailings as affected by incubation period

Amendments Rate of application			
(kg ha ⁻)	We	eks after incub	ation
	4	8	12
	EDTA -	extractable As	(mg kg ⁻)
	22 111	chiractacie i is	(mg mg)



Zeolite	0	178.6	177.9	177.5	
	30	178.0	176.9	176.4	
	60	177.1	175.1	174.6	
Beringite	0	179.2	178.8	179.2	
	30	160.7	156.6	153.6	
	60	135.3	129.8	123.5	
LSD (0.05):	.18.6				

Table 4. Effects of rate of application and type of amendment on EDTAextractable arsenic of uncapped tailings as affected by incubation period

Amendments		Rate of application (kg ha ⁻)	Weeks after incubation		
		4	8	12	
			EDTA –extract	table As (mg kg ⁻)	
Zeolite	0	789.0	789.7	790.3	
	30	787.4	785.8	784.7	
	60	786.1	784.5	783.5	
Beringite	0	789.8	789.3	790.4	
	30	744.8	743.1	738.7	
	60	725.9	722.5	716.2	
LSD (0.05):	2.90				

3.2 Effects of Amendments on Water-soluble Arsenic in Non-mined Soil, Capped and Uncapped Tailings

Tables 5, 6 and 7 show the results of the effects of the application zeolite and beringite on water soluble arsenic in non-mined soil, capped and uncapped tailings. The levels of water soluble arsenic (As-water) after the application of either 30 kg ha⁻¹ or 60 kg ha⁻¹ beringite to non-mined soil, capped and uncapped tailings resulted in significant (P = 0.05) successive decreases in As-water at 4, 8 and 12 WAI compared to un-amended soil (Tables 5, 6 and 7). The decrease in extractable arsenic in contaminated soil/tailings occurred possibly due to the presence of bivalent cations (Ca²⁺ and Mg²⁺) in beringite that aid in the adsorption of arsenic (Shazia, Alia, Iftikhar, & Muhammad, 2015). This could have resulted in the significantly lower concentration of As-water of beringite-amended soil/tailings compared to un-amended samples. The reduction in extractable As-EDTA (Tables 2, 3 and 4) as a result of the complexation of arsenic to form calcium arsenate in the inner sphere of ettringite that is a constituent of the beringite structure and the sorption of As in beringite (Rascio &



Navari-Izzo, 2011) could also be assigned to the significantly lower (P = 0.05) concentration of water soluble As in beringite-amended samples compared to un-amended samples in non-mined, capped and uncapped tailings (Tables 5, 6 and 7). There was no significant difference in the level of As-water between zeolite applied at either 30 kg ha⁻¹ or 60 kg ha⁻¹ and un-amended uncapped tailings at 4 WAI (P = 0.05). At 8 and 12 WAI, the application of 60 kg ha⁻¹ zeolite gave the same effects as un-amended uncapped tailings (P = 0.05) (Table 7). The insignificantly different concentrations of water soluble As compared to un-amended samples at the 60 kg ha⁻¹ rate of application (Table 7) indicates that zeolite is not effective in remediating/immobilizing arsenic in uncapped tailings.

The high pH of zeolite (7.0 - 9.2) and the resultant neutral to slightly alkaline pH of zeolite-amended uncapped tailings (7.3 - 7.5) could lead to a decrease in the sorption of As. The concentration of water soluble As was not significantly different between un-amended samples and non-mined soil/capped tailings amended with either 30 kg ha⁻¹ or 60 kg ha⁻¹ zeolite throughout the incubation periods at 4, 8 and 12 WAI (P = 0.05) (Tables 5 and 6). The concentration of As-EDTA adsorbed onto the soil/tailings was not significantly different with the application of either 30 kg ha⁻¹ or 60 kg ha⁻¹ zeolite compared to un-amended capped tailings at the end of the incubation experiment (Table 4). This implies that there were slight and insignificant (P = 0.05) changes in the adsorption of extractable and available arsenic within the soil matrix. Hence the insignificantly (P = 0.05) different levels of water soluble arsenic extracted from zeolite and un-amended capped tailings.

Table 5.Effects of rate of application and type of amendment on water soluble arsenic of non-mined soil as affected by incubation period

Amendments		Rate of application (kg ha)		Weeks after incubation		
			_	_4	8	12_
				EDT	A –extractable A	s (mg kg ⁻)
Zeolite	0	0.229	0.299)	0.228	
	30	0.238	0.236	;	0.237	
	60	0.234	0.233		0.233	
Beringite	0	0.229	0.229		0.228	
	30	0.095	0.088		0.089	
	60	0.045	0.036		0.035	
LSD (0.05):	0.134					



Table 6. Effects of rate of application and type of amendment on water soluble arsenic of capped tailings as affected by incubation period

Ameno	lments	Rate of applicat (kg ha ⁻)	ion Wee	ks after incubation
		4	8	12
		ED	OTA –extractat	ole As (mg kg ⁻)
	0	0.446	0.445	0.446
Zeolite	30	0.438	0.446	0.450
	60	0.438	0.441	0.442
	0	0.446	0.447	0.446
Beringite	30	0.316	0.304	0.299
	60	0.289	0.222	0.201
LSD (0.05):	.0.027			

Table 7. Effects of rate of application and type of amendment on water soluble arsenic of uncapped tailings as affected by incubation period

Amendment	s	Rate of application (kg ha ⁻)	Weeks after incubation			
			4	8	12	
			EDTA –extractable As (mg kg			
Zeolite	0		3.11	3.11	3.11	
	30		3.15	3.22	3.24	
	60		3.12	3.13	3.14	
Beringite	0		3.11	3.11	3.11	
	30		2.20	1.18	1.60	
	60		1.99	1.14	0.87	
LSD (0.05):	.0.11					



3.3 Concentration of Arsenic in Lactuca sativa grown on Mediated Soil/Tailings

Edible plants grown on polluted soil can accumulate potentially toxic elements at high concentrations that pose a threat to human health (Vosta, Graminas &Samara, 1996; Sharma, Bangar, Rajesh Jain. & Sharma., 2004; Wauchope, 1983). Ultimately, lettuce was cropped on mediated soil/tailings so as to assess the uptake and toxicity levels of As. Figures 1, 2 and 3 show the results of the concentration of Arsenic in lettuce (*Lactuca sativa*) harvested from non-mined soil, capped and uncapped tailings after mediation with beringite and zeolite. Arsenic (As) accumulated in the aerial part of the lettuce plant has been used as an index of bioavailability for the purpose of this study. Generally, the concentration of As in zeolite and beringite-amended soil/tailings were significantly lower than that of un-amended samples (P = 0.05) (Figures 1, 2 and 3).

3.4 Bioavailability of Arsenic in Lettuce Grown on Soil/Tailings Amended with Zeolite and Beringite

The concentration of arsenic accumulated in lettuce ranged from 0.27 - 4.13 mg kg⁻¹. This concentration of arsenic is within the limits given by (FSA, 2007). Plants that are not hyper-accumulating species accumulate arsenic in a normal range of 0.1 - 5 mg As kg⁻¹ dry weight in their leaves (FSA, 2007).

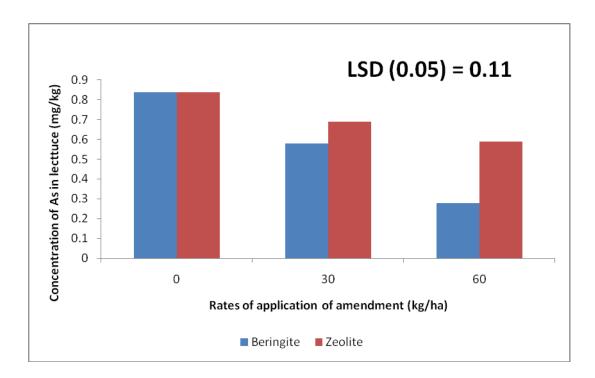


Figure 1. Bioavailability of As in Lactuca sativa L. grown on amended non-mined soil



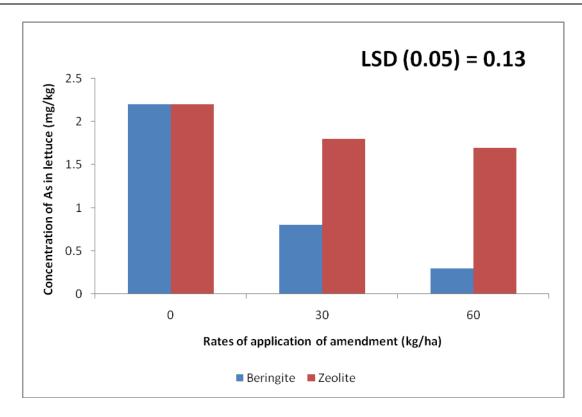


Figure 2. Bioavailability of As in Lactuca sativa L. grown on amended capped tailings

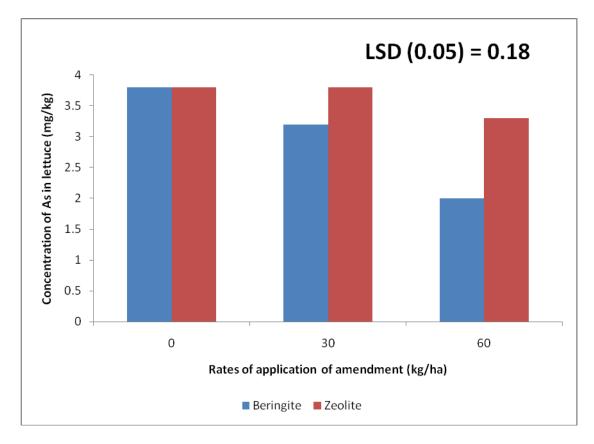


Figure 3. Bioavailability of As in Lactuca sativa L. grown on amended uncapped tailings



Except for the non-mined soil, there were no significant (P = 0.05) differences in the concentration of arsenic in lettuce grown in either 30 kg ha⁻¹or 60 kg ha⁻¹zeolite amended soil/tailings (Figures 1, 2 and 3). However, the application of zeolite at 60 kg ha⁻¹ resulted in a significantly (P = 0.05) lower concentration of arsenic in lettuce compared to un-amended soil/tailings. This trend was observed due to minimal effects of zeolite on As-EDTA and water soluble Arsenic (Tables 5-7) in non-mined soil, capped and uncapped tailings. Non-mined soil mediated with zeolite resulted in a lower content of As in lettuce aerial biomass. The concentration of arsenic in un-amended non-mined soil (Figure 1) was lower than the UK standard of 1.0 mg As kg⁻¹ dry mass (FAO & WHO, 2001). On the other hand, capped and uncapped tailings amended with zeolite resulted in the concentrations of arsenic in lettuce plants (Figures 2 and 3) that were above the UK acceptable limits for food (FAO & WHO, 2001) The cause of arsenic contamination may not necessarily be geologic in origin but anthropogenic. This is because the concentration of arsenic in lettuce grown on non-mined soil was within acceptable limits despite the underlying arseno-pyritic rocks. The beneficiation of gold in the study area is from arseno-pyrite rock ores which result in the release of high amounts of As during gold ore processing into the feed of tailings effluent, capped and uncapped tailings. The effects of mining activities in arsenic contamination were manifested through the concentrations of arsenic in lettuce that were above the UK acceptable limits for food. The application of beringite to soil/tailings resulted in the least content of arsenic in lettuce plants. The concentrations of arsenic in lettuce plants grown in un-amended non-mined soil, capped and uncapped tailings were 0.83, 2.13 and 3.87 mg kg⁻¹, respectively (Figures 1, 2 and 3). These were in contrast to the lower arsenic values of 0.27 – 0.47 mg kg^{-1} , $0.37 - 0.83 \text{ mg kg}^{-1}$ and $2.1 - 3.23 \text{ mg kg}^{-1}$ in lettuce cultivated on beringite-mediated non-mined soil, capped and uncapped tailings respectively. The significantly lower concentration of arsenic in lettuce could be attributed to the significantly immobilized arsenic by beringite and lower concentrations of EDTA-extractable (Tables 2-4) and water soluble arsenic (Tables 5-7) in beringite-amended soil/tailings. The concentration of arsenic in lettuce grown in beringite-mediated capped tailings was therefore lower than the United Kingdom (UK) limit of 1.0 mg As kg⁻¹ in food (FAO & WHO, 2001). However, the concentrations of arsenic in lettuce cultivated on un-amended capped tailings (2.13 mg kg⁻¹) (Figure 2) and amended uncapped tailings $(2.1 - 3.23 \text{ mg kg}^{-1})$ (Figure 3) was higher than the UK acceptable limit of 1.0 mg As kg⁻¹. Heavy metals in the soil can be absorbed by plants. The health of humans and other organisms can be adversely affected through the ingestion of both water and food that have been contaminated by the soil According to FAO & WHO (2001), arsenic toxicity usually occurs through contaminated food and drinking water. This implies that it would be safe to crop only on beringite-mediated capped tailings but not on either un-amended capped tailings or beringite-amended uncapped tailings.

4. Conclusion

Beringite was more effective in adsorbing and immobilizing arsenic experimental soil/tailings as compared to zeolite. The bioavailability test using lettuce, a leafy vegetable indicated that beringite applied at 60 kg ha⁻¹ reduced As below the recommended maximum limit of Food and Agriculture Organization (1.0 mg kg⁻¹). Although the study was done in a pot/laboratory,



field trials would be necessary to consolidate the potential of beringite in remediating arsenic contaminated soils, most especially at mine-contaminated sites.

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