

## Investigating Heavy Metal Pollution in Soils at Tarkwa, Ghana

### Abstract

The study was done at Tarkwa in the Western Region of Ghana which has a long history of mining activity. Soil samples were collected from 15 communities in the Tarkwa mining area and the concentrations in parts per million (ppm) of Mercury (Hg), Arsenic (As), Cadmium (Cd), and Lead (Pb) were determined in each sample using Instrumental Neutron Activation Analysis (INAA). Heavy metal concentrations of soils in the Tarkwa mining area were generally lower than expected of a typical mining area. The results of the study showed that except for isolated cases at Old Town, Akyempim, and Mile 10½ respectively, arsenic, cadmium, and mercury, levels in soil were far below the limits set by the International Atomic Energy Agency- Soil-7. Lead was within limits at all the sampling stations. It may, therefore, be concluded that contamination of soils by heavy metals is of no great significance in the study area. Periodic analysis of water sources in the area should be done to ascertain the concentration levels of heavy metals so that the local people as well as the miners could be advised accordingly.

**Key Words:** contamination, geochemistry, geology, infiltration, topography, concentration

### Introduction

Globally, the overall impacts of mining activities are severe (Hilson, 2001) especially in areas where small scale mining is practiced. This is because most small scale miners do not use approved and scientific methods for mining and processing. There have been earlier reports that

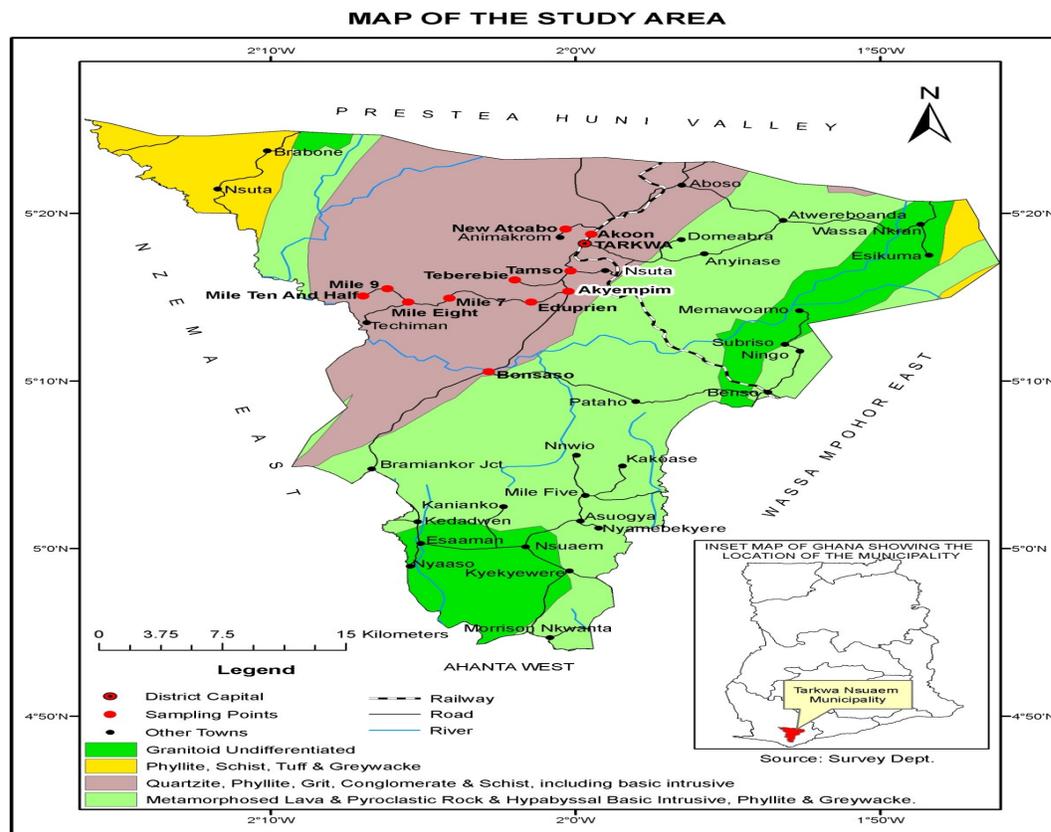
22 contamination of soils in mining areas depends on the geochemistry and the geology of an area  
23 over a period of time (Bell, 1998).

24 Most major towns in Tarkwa-Nauaem Municipality in the Western Region of Ghana rely solely  
25 on groundwater and so the number of boreholes and hand-dug wells are increasing rapidly to  
26 match the demand for portable water (Kortatsi, 2004). Surface water taken from the river Bonsa  
27 at Bonsaso is treated and distributed to Tarkwa town. Borehole yield varies from 0.4-18 m<sup>3</sup>h<sup>-1</sup>  
28 with an average of 2.4 m<sup>3</sup>h<sup>-1</sup>. The depth varies between 18 m to 75 m with an average of 35.4  
29 (Kortatsi, 2004). The recharge of groundwater in the area occurs mainly by direct seepage or  
30 infiltration. In some places groundwater is in hydraulic contact with rivers and recharge from  
31 them can also take place (Kortatsi, 2004). A study by Ragnar and Bjorn (2005) showed that  
32 metal levels in ground water exceeded WHO guidelines for drinking water in many areas in the  
33 Western Region including Tarkwa where this study was done.

34 This study seeks to ascertain the level of heavy metal pollution of soils at Tarkwa which is a  
35 known mining town in Ghana and has a long history of small scale mining. The objective of the  
36 study was to investigate heavy metal pollution in soils in some selected communities of the  
37 Tarkwa-Nauaem Municipality in the Western Region of Ghana.

### 38 **The study area**

39 Tarkwa-Nauaem Municipality lies within latitudes 4°N and 5° 40'N, and 1° 45' W and 2° 10'  
40 W. The total land area of the district is 92,354 km<sup>2</sup> (Kusimi, 2007) (Fig 1) below.



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42 The geology of the Tarkwa mining area consists of a Basement complex, further divided into

43 different sub provinces including the metamorphosed and folded rocks of the Birimian and

44 Tarkwaian system. In several places, these systems are intruded by sills and dykes of igneous

45 rocks ranging from felsite and quartz porphyry to metadolerite, gabbro and norite (Kortatsi,

46 2004). The geomorphology of the area consists of a series of ridges and valleys parallel to each

47 other and to the strike of the rocks. The strikes of the rocks are generally in north-south direction

48 (Kortatsi, 2004). Both the Tarkwaian and Birimian systems are folded along axes that trend

49 northeast. The general type of topography reflects the underlying geology (Kortatsi, 2004).

50 There are two types of soil in the study area: the Forest Oxysols in the south, and the Forest

51 Ochrosol-oxysol integrates in the north (Kortatsi, 2004). The forest oxysols are porous, well

52 drained, and generally loamy-brown to orange. Due to heavy rains in the south, a high degree of

53 leaching and reduction of nutrients have occurred, making the soil acidic (Kortatsi, 2004). The  
54 soil consists mostly of silty-sands with minor patches of laterite, mainly on hilly areas (Kuma  
55 and Younger, 2001). The soil type distribution and their textures include: Banket (silty-sand  
56 Laterite), Tarkwa Phyllite (Laterite), Huni (silty-sand), Kawere (silt sand), and Weathered dyke  
57 (silt) (Kuma & Younger, 2001).

### 58 **Research Methodology**

59 Soil samples were collected from 15 communities in the Tarkwa mining area and the  
60 concentrations in parts per million (ppm) of Mercury (Hg), Arsenic (As), Cadmium (Cd), and  
61 Lead (Pb) were determined in each sample using Instrumental Neutron Activation Analysis  
62 (INAA). Analytical treatment of data was made in Excel, and Arcview.

### 63 *Sample collection and preparation*

64 Soil samples were collected from fifteen communities within the Tarkwa mining area. The soil  
65 samples were collected from farmlands at a depth of 0-20 cm using hand auger and plastic  
66 trowel. The samples were stored in dry polythene bags and sent to the Ghana Research Reactor-1  
67 (GHARR-1) laboratory at Ghana Atomic Energy Commission (GAEC) for analysis. The samples  
68 were collected into clean plastic bottles (obtained after thorough rinsing) and carried to the  
69 GHARR-1 laboratory for analysis.

70 The soil samples were air dried for five days. They were then ground and homogenized after  
71 which they were sieved through a micro mesh of size 180  $\mu\text{m}$  to obtain fine grains. 100 mg of  
72 soil sample was weighed in six folds for each test material on a clean polythene film, folded with  
73 forceps and heat-sealed with a air dryer. Each sample was in turn put into a rabbit capsule and  
74 smoothly heat-sealed with a soldering rod. Standard Reference Materials IAEA-Soil 7 and  
75 estuarine Sediment were equally weighed as the test samples.

76 The samples and the controls were irradiated at the Ghana Atomic Research Reactor-1  
77 (GHARR-1) at the Ghana Atomic Energy Commission (GAEC), operating at 15KW at a thermal  
78 flux of  $5 \times 10^{11} \text{ n s}^{-1} \text{ cm}^{-2}$ . The samples were transferred into irradiation sites via pneumatic  
79 transfer system operating at a pressure of 0.60 mpa. The irradiation was done according to the  
80 half lives of the elements of interest. The soil samples were irradiated for one hour and delayed  
81 for 24 hrs, with 10 mins counting time.

82 After the irradiation, radioactivity measurement of the induced radionuclide was performed by a  
83 PC- based gamma ray spectrometry set-up which consists of an n-type HPGE detector, coupled  
84 to a computer based multi-channel analyzer (MCA) via electronic modules. The relative  
85 efficiency of the detector was 25% and its energy resolution was 1.8 keV at gamma ray energy of  
86 1,332 keV, belonging to  $^{60}\text{Co}$ . Through appropriate choice of cooling time, detector's dead time  
87 was controlled to be less than 10%. Identification of the gamma ray of product radionuclide was  
88 done using the energies and the quantitative analysis of the elements was achieved by the use of  
89 gamma ray spectrum analysis software, ORTEC MEASTRO-32. The quantitative analysis  
90 involved the conversion of the area under the photo peaks of the identified elements into  
91 concentrations.

92 For the determination of lead, 1.5 g of soil sample was digested in a microwave oven at a  
93 maximum power of 450 W for 10s, using a Teflon closed Parr bomb. The reagents used included  
94 a combination of 6.0 ml of 65%  $\text{HNO}_3$ , 3.0 ml of 35%  $\text{HCl}$ , and 0.25 ml of 30%  $\text{H}_2\text{O}_2$ . After  
95 that, samples were filtered carefully through Whatman 541, and quantitatively transferred into 50  
96 ml plastic flasks and undertaken through AAS measurements.

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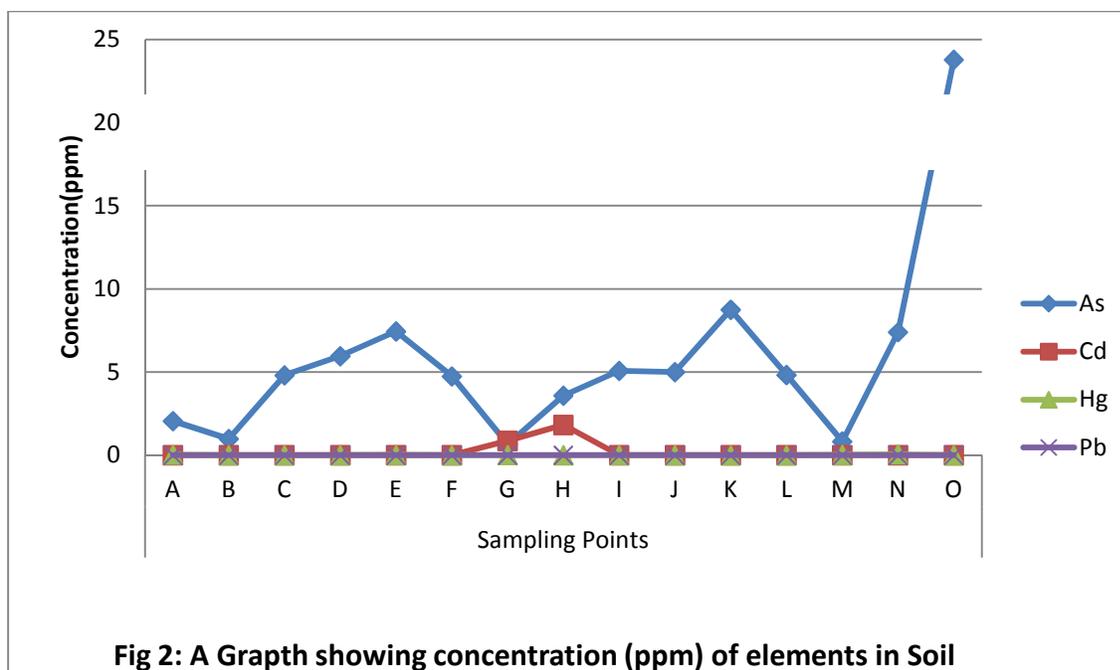
99 **Results and Discussion**

100 The concentrations of arsenic, cadmium, mercury, and lead in  $\mu\text{g/g}$  for soil and  $\text{mg/l}$  for water as  
101 obtained from the laboratory analysis were sampled from different points and represented with  
102 the letters A-O (Fig 1).

103 The sampling points for the analysis were:

104 A= Mile 8, B= Naboaso (Tarkwa), C= Post Office Sq.(Tarkwa), D= Tamso, E=Eduprien,  
105 F= Awunaben Basin, G= Akoon, H= Akyempim, I= Teberebie, J= Mile 9, K= New Atuabo, L=  
106 Mile 7, M= Bonsaso, N= Mile 10½, O= Old Town (Tarkwa).

107 This discussion was based on the premise that quality control bodies such as International  
108 Atomic Energy Agency (IAEA) have come out with specific guidelines regarding heavy metal  
109 (trace element) concentration in soils; and that at concentrations higher than the guideline value,  
110 there is the possibility of plants (including crop plants) taking up these metals and eventually  
111 transferring them to humans through the food chain. Fig 2 below shows the concentration of  
112 Arsenic, Cadmium, Mercury, and Lead in soil samples picked at 15 locations within the Tarkwa  
113 Mining Area.



**Fig 2: A Graph showing concentration (ppm) of elements in Soil**

114

115 ***Arsenic***

116 Arsenic concentration in soil ranges from 0.65  $\mu\text{g/g}$  to 23.78  $\mu\text{g/g}$  (Fig. 2).. The highest  
 117 concentration was obtained at Old Town (Tarkwa). The second highest concentration (8.75  $\mu\text{g/g}$ )  
 118 occurred at New Atuabo. New Atuabo and Old Town lie along the same geological plane and so  
 119 have the highest arsenic concentrations. Arsenic level in soil increased from Tamso (5.95  $\mu\text{g/g}$ )  
 120 to Eduprien (7.44  $\mu\text{g/g}$ ); it then decreased from Teberebie (5.07  $\mu\text{g/g}$ ) to Mile 9 (5.00  $\mu\text{g/g}$ ), and  
 121 further to 4.82  $\mu\text{g/g}$  at Mile 7. However, the concentration fluctuated again to 7.40  $\mu\text{g/g}$  at Mile  
 122 10½. The decrease trend may be due to geological factors whilst the sharp increase could be  
 123 blamed on some anthropogenic factors. The IAEA-Soil-7 recommended value for arsenic  
 124 concentration in soils is 13.50  $\mu\text{g/g}$ . This implies that all but only one station had soil arsenic  
 125 levels below the IAEA guideline value. The high arsenic concentration recorded at Old Town  
 126 was seen as an isolated case.

127 ***Mercury***

128 High mercury concentration in soil ( $0.05 \mu\text{g/g}$ ) occurred at Mile  $10\frac{1}{2}$ . This was followed by Mile  
129 8 which recorded  $0.03 \mu\text{g/g}$  (*Fig. 2*). The IAEA-Soil-7 recommended value for mercury level in  
130 soils is  $0.04 \mu\text{g/g}$ . This means that all the stations have mercury levels within limits except Mile  
131  $10\frac{1}{2}$  which had a concentration ( $0.05 \mu\text{g/g}$ ) slightly above the recommended value. A critical  
132 observation of the trend of mercury occurrence in soils in the study area shows that mining  
133 activities, especially small scale mining operations, play a significant role in mercury  
134 concentration in soils in the study area.

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#### 137 ***Cadmium***

138 Cadmium level in soils at all the stations was less than  $0.01 \mu\text{g/g}$  except at Akyempim and  
139 Akoon which recorded  $1.82 \mu\text{g/g}$  and  $0.86 \mu\text{g/g}$  respectively (*Fig.2*). The low concentrations  
140 could be due to geological factors. The IAEA-Soil-7 recommended value for cadmium  
141 concentration in soils is  $1.30 \mu\text{g/g}$ . Akyempim had a concentration of  $1.82 \mu\text{g/g}$  which is above  
142 the IAEA guideline value. A dose of 10 mg of Cadmium can produce symptoms of cadmium  
143 poisoning (ATSDR, 2007) and so plants must be monitored to avoid bioaccumulation in humans.

#### 144 ***Lead***

145 Lead concentration in all the samples was below  $0.01 \mu\text{g/g}$  (*Fig. 2*). The low concentration is  
146 probably due to the minute quantities in which lead minerals occur in the rocks of the area. The  
147 IAEA-Soil-7 recommended value for lead concentration in soils is  $60.00 \mu\text{g/g}$ . This means that  
148 the recorded concentrations are far below the IAEA guideline value. Hence, no soil  
149 contamination in terms of lead is occurring in the study area.

**150 Conclusions**

151 Heavy metal concentrations in the Tarkwa mining area, especially in soils, were generally lower  
152 than expected of a typical mining area. The results of the study showed that except for isolated  
153 cases at Old Town, Akyempim, and Mile 10½ respectively, arsenic, cadmium, and mercury,  
154 levels in soil were far below the limits set by the International Atomic Energy Agency- Soil-7.  
155 Lead was within limits in at all the sampling stations. It may, therefore, be concluded that  
156 contamination of soils by the named heavy metals is of no great significance in the study area.

**157 Recommendations**

158 There should be intensified education on precautionary measures to be taken by miners,  
159 especially small scale miners (e.g. wearing of overalls, hand gloves etc., avoiding spillage, etc.)  
160 when dealing with chemicals such as mercury. Further research should be done on  
161 bioaccumulation of heavy metals in plants.

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