1	Original Research Article
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3	<b>BIOREMEDIATION OF HEAVY METALS IN THE SOIL BY</b>
4	PSEUDOMONAS AERUGINOSA AND TRICHODERMA HARZIANUM
5	USING SOLANUM LYCOPERSICUM AS TEST PLANT
6	

## 7 ABSTRACT

8 This study determined the heavy metal concentrations of contaminated stream water and assessed the heavy metal 9 contents of pre- and post-cropped sterilized soil. It also determined the heavy metal uptake of the S. 10 lycopersicum plant. This was with a view to assessing the potential of Pseudomonas aeruginosa and Trichoderma 11 harzianum for transforming heavy metals in heavy metal contaminated stream water. Experimental pots containing 12 3000 g of sterilized soil was used for this experiment whereby 60 sample pots were used with various treatments in 13 this study. Solanum lycopersicum seeds were raised in the nursery for a period of 3 weeks and treatments applied 14 just before transplanting into the experimental pots. The plants were left for a week so as to be established properly 15 and overcome transplanting shock before watering with the contaminated stream water. Heavy metal analysis using 16 Atomic Absorption Spectroscopy (AAS) method was carried out on the contaminated stream water to determine the 17 amount of heavy metal in the stream water before the commencement of the experiment. The contaminated stream 18 water was applied to the pots in measured quantities; 0, 5 and 0%. Pre and post soil heavy metal analysis were 19 carried out on the soil samples. At harvest, plant tissues were analysed for heavy metals using AAS method. The 20 results showed that heavy metals were present in high concentration in the stream water sample. The values of the 21 heavy metals in the stream water sample used for watering were Iron - 138.15 mg/L, Zinc - 68.4 mg/L, Lead - 7.89 22 mg/L and Copper – 8.98 mg/L. Heavy metal analysis of the soil and all the treatments revealed that treatments with 23 P. aeruginosa inoculation had the lowest level of Iron, Copper, Zinc and Lead followed by treatments inoculated 24 with T. harzianum. The study concluded that the use of contaminated stream water for irrigation could be a potential 25 source of heavy metals in tomato. However, inoculation of microorganisms for the treatment of the heavy metal 26 contaminated sites was effective Phytoremediation, for increased health, growth and yield of tomato fruits.

## 27 **KEYWORDS:** Phytoremediation, Pseudomonas aeruginosa, Trichoderma harzianum, Solanum lycopersicum

## 28 INTRODUCTION

29 Heavy metals represent a great environmental concern, because of their widespread use and distribution, and 30 particularly their toxicity to human beings and the biosphere. However, they also include some elements that are 31 essential for living organisms at low concentrations (Alloway, 1990). These elements are usually transition metals. 32 They have high densities (>5 g cm-3) when compared with other materials (Baird and Cann, 2005). Human 33 activities such as industrial production, mining, agriculture and transportation lead to release of high amount of 34 heavy metals into the biosphere. The primary sources of metal pollution are the burning of fossil fuels, smelting of 35 metal like ores, municipal wastes, fertilizers, pesticides and sewage (Nriagu, 1979, 1996; Pendias and Pendias, 1989; 36 Rai, 2009). Heavy metal contamination may occur due to factors which could include irrigation with contaminated 37 water, addition of fertilizers and metal based pesticides, industrial emissions, and transportation (Radwan and 38 Salama, 2006; Tuzen and Soylak, 2007; Duran et al., 2007). Heavy metal pollution does not only affect the 39 production and quality of crops, it also influences the quality of the atmosphere and water bodies. This threatens the health and life of animals as well as human beings by the way of food chain and most phenomenal is that, this kind 40 of pollution is covert, long term and non-reversible (Zhang, 1999). Heavy metals are also one of the major 41 42 contaminating agents in our food supply (Zaidi et al., 2005; Khair, 2009). Bioremediation is a process that uses

43 naturally occurring micro-organisms to transform harmful substances to nontoxic compounds, these processes which 44 take advantage of microbial degradation of organic and inorganic substances can be defined as the use of micro-45 organisms to remove environmental pollutants of soils, water and sediments (Pala et al., 2006). Bioremediation 46 involves the use of organisms for the treatment of polluted soils. These organisms which could be micro-organisms 47 or green plants eliminate, attenuate or transform the harmful substances via biological processes to a less harmful 48 substance (Mrayyana and Battikhi, 2005). Micro-organism breaks down organic molecules to carbondioxide, 49 fattyacid and water in order to obtain energy and nutrients. Bioremediation occurs naturally (even though it could be 50 enhanced by a number of processes), thus, it is widely accepted by the general public as a safe way of treating 51 polluted soils. Trichoderma harzianum has potential in stimulating phytoremediation directly and indirectly and 52 therefore, inoculation of plants with this fungus could be a feasible approach to enhance the transformation of 53 hydrocarbons in polluted soil. T. harzianum also have the ability to solubilize metal ions and produce siderophores 54 to chelate iron, making metal ions required for plant growth more available to the plant (Harman et al., 2004). The 55 fungus is thought to colonize roots of annual plants for their entire lifetime by penetrating the outer layers of the 56 roots (Harman et al., 2004). This makes the plants release more root exudates to the surrounding soil, thus, 57 stimulating microbial degradation of pollutants. Trichoderma harzianum has been shown to induce the production of 58 larger and deeper root systems, and plants inoculated with Trichoderma harzianum also produce greater plant 59 biomass. Such plants are more resistant to abiotic stress and take up nutrients more effectively (Harman et al., 60 2004). Edwards et al., (2006) noted that various bacteria such as Pseudomonas aeruginosa produce surfactants that 61 aid in the biodegradation. A recent study has found a P. aeruginosa strain that actually supports plant growth. This 62 characteristic, along with the fact that *P. aeruginosa* can transform polycyclic aromatic hydrocarbons, suggests the

- 63 future uses of *P. aeruginosa* for environmental detoxification of synthetic chemicals and pesticides and for industrial
- 64 purposes (Botzenhardt and Doring, 1993).
- 65

#### 66 MATERIALS AND METHOD

#### 67 Collection of Contaminated water, seeds and microorganisms

- Heavy metals contaminated stream water was obtained from a flowing stream. It is situated at 7°30' Northern
   latitude and 4°28' Eastern longitude. The sampling point was located at the back of the Ife Iron and Steel Nigeria
- Limited along Ife-Ibadan expressway. Surface water samples was collected at downstream into clean plastic kegs.
- 71 The water samples were collected during the month of April, 2015. Seeds of *Solanum lycopersicum* cultivar (ROMA
- 72 VF) were obtained from Institute of Agricultural Research and Training, Moor Plantation, Ibadan.

#### 73 Culturing of Organisms

- 74 A culture of *Pseudomonas aeruginosa* was obtained from the Department of Microbiology, Obafemi Awolowo
- 75 University (OAU), Ile-Ife. A culture of *Trichoderma harzianum* was also obtained from the Mycology unit of the
- 76 Department of Crop Production and Protection, OAU, Ile-Ife. A single colony of *P. aeruginosa* was subcultured by
- vising nutrient agar in Petri dishes and kept in the incubator for 48 hours at 37°C to a medium after which it was
- harvested by flooding with sterile distilled water. The bacterium inoculum was prepared by streaking a single colony
- 79 of *P. aeruginosa* earlier isolated on plated nutrient agar plate and incubated at  $37^{\circ}$ C for 48 hours. Cells of *P*.
- 80 *aeruginosa* were harvested from agar plates by flooding with sterile distilled water and standardized using a 81 colorimeter to  $10^8$  CFU/ml. Spores of *Trichoderma harzianum* was subcultured by using potato dextrose agar in
- Petri dishes and kept in the incubator for 7 days at 37°C to a medium after which it was harvested by flooding with
- sterile distilled water. The fungal spore solution was prepared by picking spores of *T. harzianum* earlier isolated on
- serve distinct water. The fungar spore solution was prepared by picking spores of *T. harzunum* carnet isolated of potato dextrose agar plate and incubated at  $37^{\circ}$ C for 7 days. Spores of *T. harzunum* were harvested from agar plates
- by flooding with sterile distilled water and standardized using a colorimeter to  $10^7$  spores/ml.

#### 86 **Preparation of Sterilized Soil for Field work**

- 87 Top soil and river sand were mixed together and sieved before it was sterilized using an autoclave by heating for 5
- 88 hours at 131°C and left to cool for four (4) days.
- 89 Planting of seeds and contamination of experimental pots

90 Seedlings of S. lycopersicum were raised on nursery beds for a period of three weeks. Sixty pots, each containing

- 91 three kilograms of soil from sterilized soil was used for this study. *Pseudomonas aeruginosa* inoculum solution (30
- 92 ml) was poured into a hole that was made in the middle of a set of 15 experimental pots containing sterized soil
- before *S. lycopersicum* seedlings are transplanted to it. *Trichoderma harzianum* spore solution (30 ml) was also poured into a hole that was made in the middle of another set of 15 experimental pots before *S. lycopersicum*
- poured into a hole that was made in the middle of another set of 15 experimental pots before *S. lycopersicum* seedlings are transplanted to them. The third set of 15 pots received dual inoculation of *Trichoderma harzianum*
- spore solution (15 ml) and *P. aeruginosa* innoculum before *S. lycopersicum* seedlings were transplanted into it; with
- 97 the final set of 15 pots acting as control at various levels. Thereafter, pot preparation was arranged in a completely
- 98 randomized design in the screenhouse.
- 99 Seedlings were left for a week to establish and overcome transplanting shock before wetting with the contaminated 100 stream water at various concentrations of 0%, 5% and 10% v/v. Contaminated stream water was quantified using the 101 formula: percentage soil contamination = (Volume of polluted stream water applied / Volume of soil) x 100. Each
- treatment of the experiment was replicated three times. Twenty four pots were watered with the contaminated stream
- 103 water once during the experiment and another 24 pots watered daily with the contaminated stream water. The
- remaining 12 pots which served as the control experiment were watered daily with distilled water. Pots containing *S*.
- 105 *lycopersicum* was watered regularly to ensure adequate moisture. Heavy metal analysis on the contaminated stream
- 106 water was carried out using AAS (Atomic Absorption Spectrophotometer) for Iron, Copper, Lead, and Zinc pre
- experiment. Plant samples were also subjected to heavy metal analysis using AAS (Atomic Absorption
   Spectrophotometer) for Iron, Copper, Lead, and Zinc post experiment. Pre and post soil tests were carried out to
- determine soil nutrients. Soil samples were also subjected to heavy metal analysis using AAS (Atomic Absorption
- 110 Spectrophotometer) for Iron, Copper, Lead, and Zinc pre and post soil tests. Data obtained was subjected to
- 111 statistical analysis using descriptive and inferential methods.

# 112 **Experiment (Treatment Layout)**

- 113 Sterilized soils were polluted with contaminated stream water at a calculated percentage using the formula;
- 114 Percentage soil contamination = (Volume of Contaminated stream water/Volume of soil) x 100.
- 115 The layout of the experiment is as follows;
- 116 Treatment 1- sterilized soil + *S. lycopersicum*
- 117 Treatment1d- sterilized soil + *S. lycopersicum* (2)
- 118 Treatment 2- sterilized soil + *Trichoderma harzianum* + *S. lycopersicum*
- 119Treatment 2d- sterilized soil + Trichoderma harzianum + S. lycopersicum (2)
- 120 Treatment 3- sterilized soil + *Pseudomonas aeruginosa* + *S. lycopersicum*
- 121 Treatment 3d- sterilized soil + *Pseudomonas aeruginosa* + *S. lycopersicum* (2)
- 122Treatment 4- sterilized soil + T. harzianum + P. aeruginosa + S. lycopersicum
- 123 Treatment 4d- sterilized soil + *T. harzianum* + *P. aeruginosa* + *S. lycopersicum* (2)
- 124 Note: (2) and d means daily wetting of pots with contaminated water
- Each of the layouts contaminated at 0, 5, and 10% (v/w) contaminated stream water concentration was replicated thrice. The experimental pots were watered regularly to ensure adequate moisture for proper growth of the test plant.

# 129 **RESULTS**

128

# 130 Physicochemical Properties of Sterilized Soil Before Planting

131 The physicochemical properties of sterilized soil before planting was found to show that heavy metals (Iron, Zinc, 132 Copper and Lead) were present in the soil with iron (Fe) having the highest concentration (Table 4.1). Exchangeable 133 acidity ( $A^{3+}$ ,  $H^+$ ) was found to have a higher concentration in the sterilized soil than exchangeable bases ( $Na^+$ ,  $K^+$ ). 134 Organic carbon percentage was also found to be lower in concentration in sterilized soil compared than organic 135 matter percentage. The total nitrogen in the sterilized soil was found to be 0.19 g/kg while the electrical conductivity of the soil was 154.65 µs/cm. The pH of the soil was slightly acidic while the calcium content of the soil was higher 136 137 than that of the magnesium. The soil particle size was found to be 76% sand, 11% silt and 12% clay. The textural 138 class of the soil was loamy sand. 139 **Table 1: Physicochemical Properties of Sterilized Soil before Planting** 

Parameters Pa	Sterilised
Ph	<mark>6.5</mark>
T.N (g/kg)	<mark>0.19</mark>
E.C (μs/cm)	<mark>154.65</mark>
ECEC (mol/kg)	<mark>3.52</mark>
H <sup>+</sup> (cmol/kg)	<mark>0.09</mark>
K (cmol/kg)	<mark>0.81</mark>
Na (cmol/kg)	<mark>0.08</mark>
Ca (cmol/kg)	2.02
Mg (cmol/kg)	0.61
P (mg/Kg)	135.21
Fe (ppm)	22.75
Zn (ppm)	10.45
<mark>Pb (ppm)</mark>	<mark>1.89</mark>
<mark>Cu (ppm)</mark>	<mark>2.93</mark>
SAND (%)	<mark>76</mark>
SILT (%)	11
CLAY (%)	12
<mark>OC (%)</mark>	<mark>1.5</mark>
<mark>OM (%)</mark>	2.5

Textural class of the soil was Loamy sand.

- 141 The heavy metals analysis of the stream water showed that heavy metals (Iron, Zinc, Copper and Lead) were present
- in high concentration in the water. Iron (Fe) had the highest concentration of 138.15 mg/L followed by zinc (Zn)
- 143 which had a concentration of 68.4 mg/L. The order of concentration was Fe>Zn>Cu>Pb.
- 144

#### 145 **Physicochemical Properties of Contaminated Stream Water**

146 The physicochemical properties of the stream water showed that heavy metals (Iron, Zinc, Copper and Lead) were 147 present in high concentration in the water above the acceptable limits by World Health Organization (2004). Iron 148 (Fe) had the highest concentration of 138.15 mg/L followed by zinc (Zn) which had a concentration of 68.4 mg/L 149 (Table 4.2). The order of concentration was Fe>Zn>Cu>Pb. The turbidity of the water was found to be 18.9 NTU 150 which is within acceptable limit by the WHO (2004), but it had a high level of conductivity which is above the 151 acceptable limits by WHO (2004). The chloride and calcium concentration of the water were found to be within the 152 normal acceptable limits by WHO 2004, while the magnesium concentration of the water was found higher above 153 the acceptable limits by WHO (2004). The biological oxygen demand and chemical oxygen demand of the stream 154 water were found to be 351.8 mg/L and 628.4 mg/L respectively. The stream water was characterized with high 155 biological oxygen demand and high level of nitrate and phosphate. The pH of the water was 6.1 and was above the 156 acceptable limits by WHO (2004) which showed that the water was acidic.

157

## Table 2: Physicochemical Properties of Contaminated Stream Water

Parameters	Stream Water
Turbidity (NTU)	18.9
Acidity (mg/L)	29.3
BOD (mg/L)	<mark>351.8</mark>
COD (mg/L)	<mark>628.4</mark>
Conductivity(mg/L)	<mark>3014.9</mark>
рН	<mark>6.1</mark>
Chloride (mg/L)	<mark>46.8</mark>
Phosphate (mg/L)	<mark>27.5</mark>
Nitrate (mg/L)	143.4
Calcium (mg/L)	43.8
Magnesium (mg/L)	<mark>75.9</mark>
Fe (mg/L)	<mark>138.15</mark>
Zn (mg/L)	<mark>68.4</mark>

Pb (mg/L)	<mark>7.89</mark>
Cu (mg/L)	<mark>8.98</mark>

159 After the soils were subjected to heavy metal analysis, it was observed that iron concentration of the soil increased 160 as the contaminated stream water concentration increased in all the treatments without any inoculation of 161 microorganism (Fig. 1). Treatments 3 and 3d inoculated with P. aeruginosa were lower in concentration of iron 162 compared to treatments 2 and 2d which were inoculated with T. harzianum. Treatment 1d without any inoculation of 163 microorganisms had highest iron concentration followed by treatment 1 also without any inoculation of 164 microorganisms at 5% and 10% contaminated stream water concentration. The order of iron concentration across the 165 treatments with 5% and 10% contaminated stream water concentration was 1d>1>4d>4>2d>2>3d>3 and 166 1d>1>4d>4>2d>2>3d>3 respectively.

167 Treatment 1d had the highest zinc concentration at 10% contaminated stream concentration followed by treatment 1 168 at the same 10% concentration. Treatment 3 at 0% concentration had the lowest iron concentration. Soil samples

169 treated with single or both micro-organisms had the lowest value in zinc compared to soil polluted with

170 contaminated stream water without any treatment with microorganisms (Fig 2). Treatment 3 had the lowest copper

171 level of 2.46 part per million (ppm) at 5% contaminated stream water concentration while treatment 1d had the

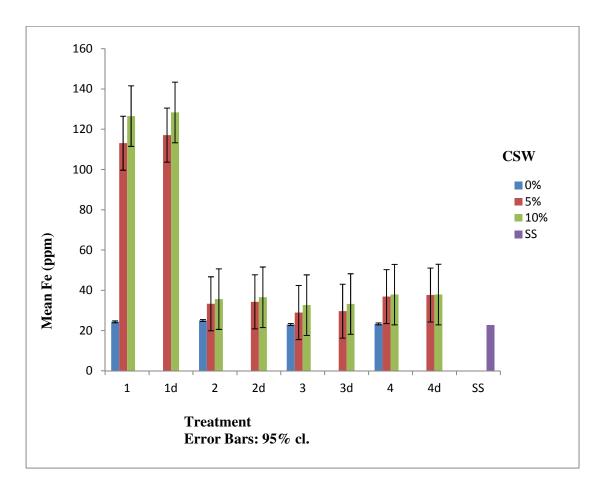
172 highest level of copper with 3.86 ppm at the same concentration (Fig 3). The order of copper concentration in 0%

and 10% was treatment4>2>2>1>3and 1d>1>4d>4>2d>2>3d>3 respectively. Lead analyses in the soil indicated

174 that the order of the concentration in 5% and 10% was 1d>1>4d>4>>2d>2>3d>3 and 1d>1>4d>4>>2d>2>3d>3

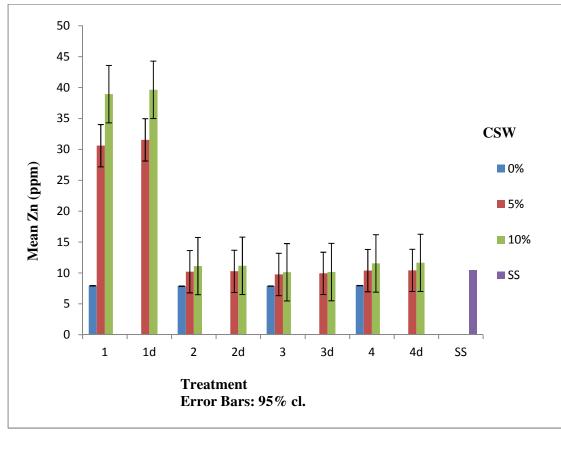
respectively, treatment 1d had the highest level of lead concentration followed by treatment 1 both at 10%

176 contaminated stream water concentration while treatment 2 had the lowest at 0% (Fig. 4).

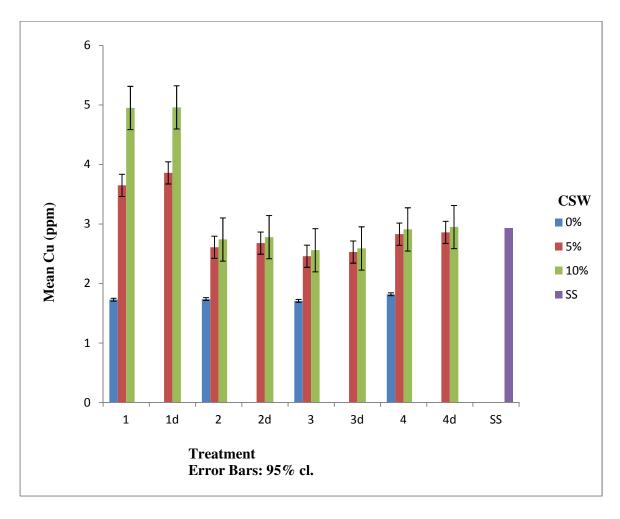


#### 179 Figure 1: Iron (ppm) content of Pre and Post Planting Soil Samples

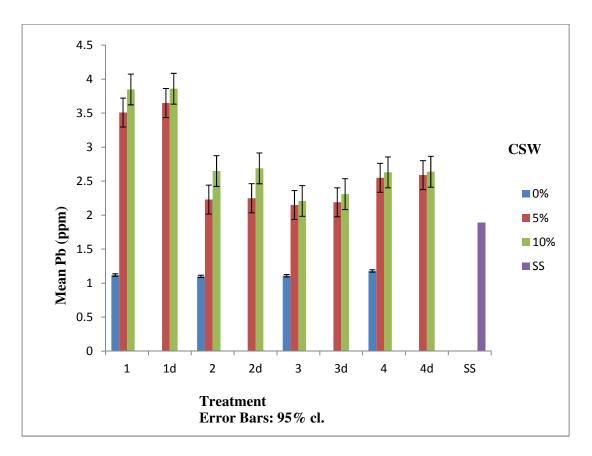
- 180 Legend
- 181 1-SS + TP
- 182 1d-SS + TP
- 183 2-SS + TH+ TP
- 184 2d-SS + TH + TP
- 185 3-SS + PA + TP
- 186 3d-SS + PA + TP
- **187** 4-SS + TH + PA + TP
- 188 4d-SS + TH +PA +TP
- 189 SS Sterilized soil before planting
- 190 d Daily wetting of plants with contaminated stream water
- 191 Cl- Confidence level
- **192** TH *T. harzianum*
- 193 PA P. aeruginosa
- 194 TP Test Plant
- 195 CSW- Contaminated Stream Water



198 Figure 2: Zinc (ppm) content of Pre and Post Planting Soil Samples



204 Figure 3: Copper (ppm) content of Pre and Post Planting Soil Samples



#### 207 Figure 4: Lead (ppm) content of Pre and Post Planting Soil Samples

208

# Heavy metal analysis carried out on plant samples showed that plants from soil samples without inoculation of micro-organisms had the highest heavy metal uptake as the concentration of contaminated stream water increased. For 5% contaminated stream water concentration, treatment 1 had the highest level of iron at 77.78 ppm followed by treatment1d with 77.71 ppm while treatment 3 had the lowest concentration of iron with 13.91 ppm (Fig 5).

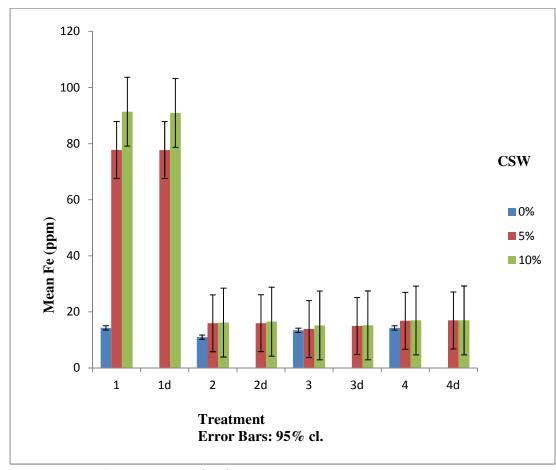
The order of concentration in iron at 10% was sample 1>1d>4d>4>2d>2>3d>3. Zinc at 10% contaminated stream water concentration had the highest concentration in treatment 1d and the lowest at treatment 3 at same 10%. The order of zinc concentration at 5% was 1d>1>4d>4>2d>2>3d>3 while 10% was 1d>1>4d>2>2d>3d>3 (Fig. 6).

Copper in treatment 1d without any inoculation had the highest concentration at 5% and 10% followed by treatment

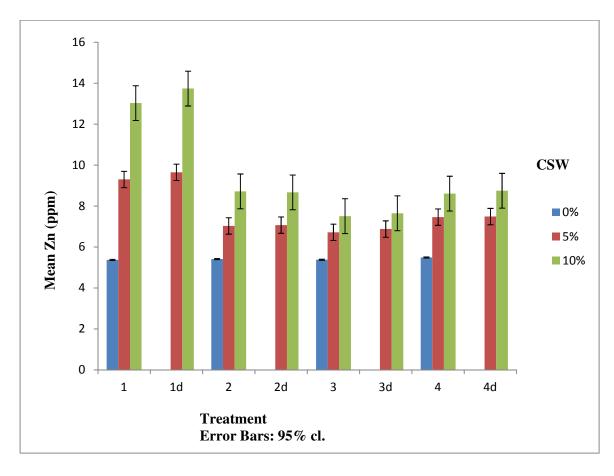
1 at same concentrations with treatment 3 inoculated with *P. aeruginosa* having the lowest value (Fig.7).

Lead content in the plant samples was highest in treatment 1d, followed by those from treatment 1 but lowest in treatment 3. Order of increase of lead is treatment 1d>1>4d>4>2d>2>3d>3 (Fig 8). Treatments 2 and 2d inoculated with *T. harzianum* had more of the heavy metal in plant tissue compared to treatments 3 and 3d treated with *P. aeruginosa*.

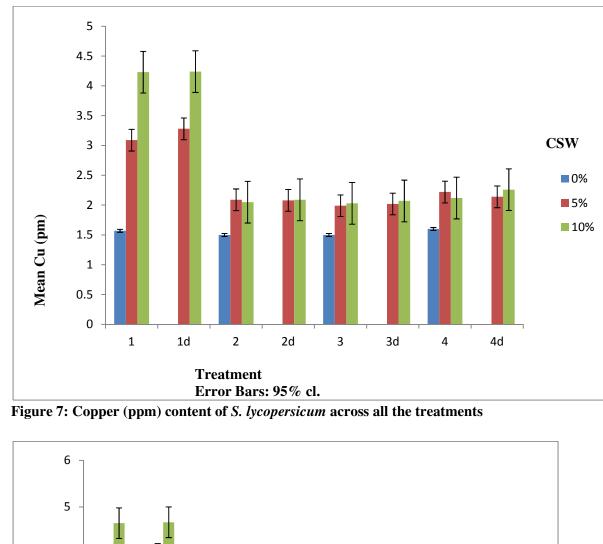
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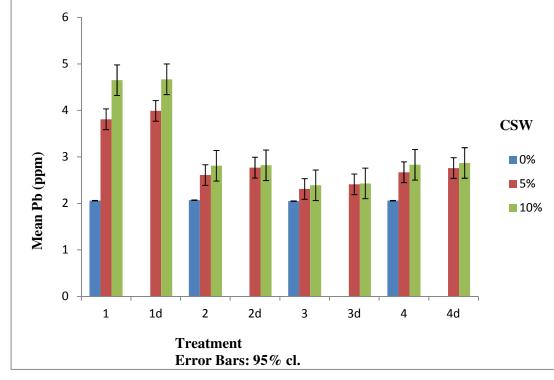
226 227 Figure 5: Iron (ppm) content of S. lycopersicum across all the treatments



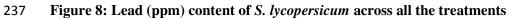
230 Figure 6: Zinc (ppm) content of *S. lycopersicum* across all the treatments











#### 239 **DISCUSSION**

Heavy metals are elements that exhibit metallic properties such as ductility, malleability, conductivity, cation stability, and ligand specificity (Opaoluwa, 2010). They are characterized by relatively high density and high relative atomic weight with an atomic number greater than 20. Industrial effluents are usually considered as undesirable for arable soil, plants, animals and human health. This is due to the contained heavy and trace metals like Cr, Mn, Fe, Cu, Co, Zn, Ni, As, Cd and Pb that are discharged continuously into water source (streams/ nullahs,

- canals and rivers). These are allowed to spread on agricultural lands. The unplanned disposal of these effluents has
- increased the threat of environmental pollution (Gulfraz *et al.*, 2003). Soils, whether in urban or agricultural areas
- represent a major sink for metals released into the environment from a wide variety of anthropogenic sources (Niragu, 1991).
- 249 Su et al. (2014) reported that low concentration of heavy metals could stimulate microbial growth and increase 250 microbial biomass, while high concentration could decrease soil microbial biomass significantly. The 251 microorganisms used in this study (T. harzianum and P. aeruginosa) were highly effective in transforming heavy 252 metals. The bio-sorption potential of the organisms used in this study showed that T. harzianum and P. aeruginosa 253 posses effective heavy metal absorption capacity. It was discovered in this study that at higher concentrations of 254 these metals, there were reductions in plant growth. This may be due to the decrease in growth parameters of S. 255 lycopersicum as the contaminated stream water concentration increased in this study. Heavy metals of soil in all the 256 soil samples showed an increase as the contaminated stream water increased in concentration. Treatments inoculated 257 with P. aeruginosa were found to have lower concentration of heavy metals (Fe, Zn, Cu and Pb) followed by 258 treatments inoculated with T. harzianum. Due to a change in their oxidation state, heavy metals can be transformed 259 to become either less toxic, easily volatilized, more water soluble (and thus can be removed through leaching), less 260 water soluble (which allows them to precipitate and become easily removed from the environment) or less 261 bioavailable (Marques et. al., 2009).
- 262 The biodegrading ability of *P. aeruginosa* which showed the most efficient heavy metal uptake from the soil is in 263 agreement with report of Lewis et al. (2002) and Odeyemi et al. (2011) which stated that Psedomonas spp have a 264 high biodegrading ability. Report from Jankiewicz et al. (2000) also support the findings from this study which 265 noted that *P. aeruginosa* cells grown in biofilms accumulate higher amounts of heavy metals. Also, many species of 266 soil fungi including *Trichoderma* are able to dissolve through the release of chelating compounds of organic acids. 267 The fungus releasing organic acids causes acidification of the environment, which helps increase the mobility of 268 heavy metals (Barea et al., 2005; Ledin, 2000; Wang and Chen, 2009). This study confirms this reports. Treatments 269 inoculated with dual inoculation of T. harzianum and P. aeruginosa were found to have slightly higher 270 concentration of heavy metals than treatments inoculated with P. aeruginosa or T. harzianum. However treatments 271 with no inoculation of one or two microorganisms showed very high concentration of heavy metals in the soil in comparison with treatments with dual microorganisms. This confirms that the microorganisms used in this study 272 273 biotransformed the heavy metals in the soil. This also revealed that there is positive and productive interaction 274 between T. harzianum and P. aeruginosa in bioremediation of heavy metals polluted soil.

275 Many species of plants have been successful in absorbing contaminants such as lead, cadmium, chromium, arsenic,

and various radionuclides from soils. Some metals with unknown biological function (Cd, Fe, Zn, Cu, Cr, Pb, Co,

- Ag, Se, Hg) can also be accumulated (Cho-Ruk *et al.*, 2006). Contaminant uptake by plants and its mechanisms have
  been being explored by several researchers. It could be used to optimize the factors to improve the performance of
- 279 plant uptake. According to Sinha *et al.* (2008), the plants act both as "accumulators" and "excluders". Accumulators
- survive despite concentrating contaminants in their aerial tissues. They biotransform the contaminants into inert
- forms in their tissues. The excluders restrict contaminant uptake into their biomass. Plant has a lot of consequences
- from heavy metal pollution in soil (Liao 1993, Su *et al.*, 2014, Wu *et al.*, 1998), plants were also seen to be polluted
- by heavy metals (Yin et al., 1999), which consequently threatens the health of animals and human beings via the
- 284 food chain (Wang *et al.*, 2001).

285 Heavy metals such as cadmium and lead are non-essential elements for plants. Microbial populations are generally 286 higher in the rhizosphere than in the root-free soil. This is due to a symbiotic relationship between soil 287 microorganisms and plants. This symbiotic relationship can enhance some bioremediation processes. Plant roots also 288 may provide surfaces for sorption or precipitation of metal contaminants (Sas-Nowosielska et al., 2008). This study 289 were found to show reduction in growth parameters as heavy metals increased which is brought by increase in 290 contaminated stream water concentration. Iron, Zinc, Copper and Lead level were higher in plant tissues from soil 291 samples containing no inoculation of microorganisms at 5% and 10% contaminated stream water concentration. This 292 was discovered to affect the growth of the plants. Su et al., (2014) reported that dicots, leafy vegetable crops are 293 sensitive to Zn toxicity, especially spinach and beet; because of their inherent high Zn uptake capacity. However soil 294 samples containing P. aeruginosa was generally the lowest in plant heavy metal uptake of iron, zinc, copper and 295 lead followed by samples containing T. harzianum. This may be an indication that the heavy metals in the soil had 296 been transformed by the microorganisms used which also showed there is low amount of heavy metals in soil left for 297 the plant to absorb. This result was found to be consistent with the work of Soumitra et al. (2014) which 298 demonstrated that P. aeruginosa reduced heavy metal uptake in Orvza sativa L. and increase its growth. Also 299 Trichoderma spp. produces organic acids such as gluconic acid, fumaric acid, and citric acid, which can decrease the 300 pH of the soil and allow for the dissolution of phosphate, as well as macro- and micronutrients such as iron, 301 manganese, and magnesium, which are necessary for plant metabolism (Ociepa, 2011; Cao et al., 2008). Treatments 302 inoculated with a combination of T. harzianum and P. aeruginosa in this study had lower concentration of heavy 303 metals in their plant tissue compared to treatments without inoculation of microorganisms. This may insinuate that 304 there is positive and effective interaction between T. harzianum and P. aeruginosa in the reduction of heavy metals

build up in plant cultivated on heavy metals polluted soil. Concentrations of metals were attributed to the contaminated stream water irrigation. The results from this study indicates that there is a serious potential health risk

307 associated with heavy metals in tomato by using contaminated water for irrigation by farmers for tomato production.

#### 308 CONCLUSION

309 It is obvious from the result of this study that biodegradation of heavy metals is an environmental friendly and easy approach to transform heavy metals in polluted soils. Pseudomonas aeruginosa showed a higher ability in 310 311 biotransforming the heavy metals in the soil than Trichoderma harzianum. The combination of the two 312 microorganisms showed a better improvement in the transformation of the heavy metal polluted soil and enhanced 313 crop production in polluted soil than soil with no inoculation of either T. harzianum or P. aeruginosa. It was also 314 observed in this study that both microorganisms enhance crop production in soil without contaminated stream water pollution. This study was able to observe the morphological and chemical differences that took place under the 315 316 different experimental treatments. It showed that use of P. aeruginosa and/or T. harzianum in the soil were able to 317 tolerate physiological stress as a result of the heavy metal pollute environment. The presence of P. aeruginosa and 318 T. harzianum were able to effectively bioaccumulate the heavy metals in the soil and increase the growth and yield 319 of S. lycopersicum. The use of fungi and bacteria in biodegradation is relatively economical and effective because it 320 is inexpensive and easy to multiply these organisms.

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