Heavy Metal Content in Soils under Different Wastewater Irrigation Patterns in Chihuahua, Mexico

V. M. Maldonado
H. Q. Rubio-Arias
R. Quintana
R. A. Saucedo
Melida Gutierrez
Missouri State University

See next page for additional authors

Follow this and additional works at: https://bearworks.missouristate.edu/articles-cnas

Recommended Citation

This article or document was made available through BearWorks, the institutional repository of Missouri State University. The work contained in it may be protected by copyright and require permission of the copyright holder for reuse or redistribution. For more information, please contact BearWorks@library.missouristate.edu.
Heavy Metal Content in Soils under Different Wastewater Irrigation Patterns in Chihuahua, Mexico

V. M. Maldonado¹, H. O. Rubio Arias¹*, R. Quintana¹, R.A. Saucedo², M. Gutierrez³, J. A. Ortega¹ and G.V. Nevarez⁴

¹College of Zootechnology and Ecology, Autonomous University of Chihuahua, Chihuahua, Mexico
²National Institute for Research in Forestry, Agriculture and Animal Production (INIFAP), Chihuahua, Mexico
³Missouri State University, Department of Geography, Geology and Planning, Springfield, MO, USA
⁴College of Chemistry, Autonomous University of Chihuahua, Chihuahua, Mexico
*Correspondence to Dr. Hector Rubio Arias. E-mail: rubia1105@hotmail.com

Received: 18 September 2008 / Accepted: 05 December 2008 / Published: 31 December 2008

Abstract: An area near the city of Chihuahua has been traditionally irrigated with wastewater to grow forage crops. It has been hypothesized that metal levels could be found in these soils high enough to cause potential health problems to the population. The objective of this study was to determine heavy metal concentrations in different soils due to irrigation practices. Four soil types were evaluated; a soil with a past and present history of wastewater irrigation (S1), a soil with a history of wastewater irrigation until 2003 (S2), a soil with no irrigation history (S3), and a soil similar to S1 and adjacent to the river where the wastewater is transported (S11). Three soil depths were evaluated; 0-15, 15-30 and 30-50 cm. Consequently, a total of 150 soil samples were analyzed evaluating pH, EC, OM and the following elements; Na, K, Cd, Pb, Ni, Cr, Cu and Fe. The pH (P=0.000) and EC (P=0.000) were different for each soil type but no differences were noted for soil depth and the interaction. Maximum pH levels were noted in S3 with a value of 8.74 while maximum EC was observed in S1 with a value of 0.850 dSm⁻¹. The OM level was different for soil type (P=0.000), soil depth (P=0.005) and the interaction (P=0.014). S1 and S11 obtained maximum levels of OM while minimum levels were noted in S3. Maximum OM levels were observed at the 0-15 cm depth followed by the 15-30 cm depth and finally at the 30-50 cm depth. The highest concentration of metals was as follows: K in S1 (359.3 mg kg⁻¹); Cd in S11 (4.48 mg kg⁻¹); Pb in S11 (155.83 mg kg⁻¹); Ni in S1 (10.74 mg kg⁻¹); Cu in S1 (51.36 mg kg⁻¹); B in S3 (41.5 mg kg⁻¹); Fe in S3 (20,313.0 mg kg⁻¹), Cr in S3 (44.26 mg kg⁻¹) and Na in S3 (203.0 mg kg⁻¹). The conclusion is that some metals are present in the soils due to anthropogenic activities but others are present in natural forms.

Keywords: Contamination, metals, environment, soil, wastewater, irrigation

Introduction

Soil is an essential natural resource for support of human life; but with time, its degradation has been constantly increasing due to the deposition of pollutants. The background concentration of metals in virgin soil depends primarily on the bedrock type from which the soil parent material was derived [1]. In addition, anthropogenic inputs may increase metal concentrations, especially in highly industrialized parts of the world producing rare and heavy metals [2]. Until recently, most studies concerning soils related to plant nutrition, with most studies published overseas.

In the particular case of Mexico, one of the environmental problems that require immediate attention is that of metal contaminated soils [3]. Due mainly to industrial activity in recent years, the deposition of a variety of elements in the natural environment has increased. As a result, the concern over water contaminated from human activities has also increased [4]. This concern takes into account a variety of elements referred to in the scientific literature as toxic metals. These metals might be found in waste materials, which in the future will likely be incorporated to the soil resource. The final outcome is an accumulation in the biota, followed by a transfer to humans through the food chain, and causing a potential risk to human health [5, 6]. According to the USDA [7], the accumulation of heavy metals in soil is toxic for humans as well as animals in general.

In agricultural areas, the addition of metals into the system is primarily due to commercial fertilizer
and includes a very dynamic human population. Several Chihuahua, a city which has grown in terms of industry has been producing crops and irrigating the land with reactions [21]. Therefore, the metals in soil may such as adsorption-desorption, occlusion, and precipitation factors [20]. The latter factors are related to processes depended not only on its concentration, but it was affected that depended largely on physical and chemical soil mobility and bioavailability of heavy metals were factors other study, Cuevas and Walter [19] concluded that the transferring metals held high precipitation events might be responsible for soil temperature and soil events over the quantity of environmental parameters such as residue amendments and Adeli et al. [16] who concluded that broiler litter application did not threaten the ecosystem. Regardless, once in the soil, contaminants may be further transported throughout the soil profiles to groundwater.

Holmgren et al. [17] created a database after analyzing 3,045 soil samples collected from 307 soil series in the United States and found differences between land regions. Dudal et al. [18] calculated the influence of some environmental parameters such as residue incorporation, soil temperature and soil events over the quantity of metals that the organic matter could hold. They found that high precipitation events might be responsible for transferring metals held in the soluble organic matter. In other study, Cuevas and Walter [19] concluded that the mobility and bioavailability of heavy metals were factors that depended largely on physical and chemical soil characteristics. Also, the mobility of metals in soil depended not only on its concentration, but it was affected by soil properties, metal properties and environmental factors [20]. The latter factors are related to processes such as adsorption-desorption, occlusion, and precipitation reactions [21]. Therefore, the metals in soil may accumulate in different forms [20] and are the foremost factors impeding the soil microbial process [22].

Ejido Tabalaopa, an area near the city of Chihuahua, has been producing crops and irrigating the land with water from the Chuviscar River. This river, a tributary of the Río Conchos, contains the wastewater of the city of Chihuahua, a city which has grown in terms of industry and includes a very dynamic human population. Several studies like those of Gutierrez et al. [23], Rubio et al. [24], Holguin et al. [25] and Gutierrez and Borrego [26] reported heavy metal concentrations in the water of the Río Conchos; thus, we expected contamination of the soil resource being irrigated with this water. Yet, in the Tabalaopa area, there is no information concerning the level of soil contamination as an outcome of wastewater irrigation. The objective of this study was to determine the level of contamination from heavy metals, in four types of soil and at three soil depths. The results offer an overall assessment of the contamination degree of the soil due to irrigation with wastewater. It is also an attempt to understand the accumulation of some elements in soil that are not biodegradable and immobile [27] and its possible transfer to the food chain and persistence in the environment as a whole.

Materials and Methods

The study was conducted during 2006, within Ejido Tabalaopa, an area of 1,094 ha located near the southeastern part of the city of Chihuahua, Mexico. The soil was an Orthid Aridisol with well developed pedogenic horizons, low organic matter in the virgin soil, and dry more than six months a year. An ejido is a community where Mexican law has established that the land must be used collectively or shared. The Ejido Tabalaopa was traditionally used to grow various crops, among them vegetables, until 1996, and using wastewater from the city of Chihuahua to irrigate the land. In 1996, the production of vegetables or any edible crop using wastewater irrigation throughout Mexico was prohibited. At present, this agricultural area is producing different sorts of forage for cattle production and some irrigated land has been abandoned.

Our study area was divided into four zones; a zone having soils with a past and present history of wastewater irrigation (S1), a zone with a history of wastewater irrigation until 2003 (S2), a soil with no irrigation history (S3), and a soil similar to S1 and adjacent to the river where the wastewater is transported (S11). This division was performed using the geographic package Idrisi Kilimanjaro. Using the map included in the package, a total of fifty points were randomly selected resulting in 18 sampling points for S1, 12 sampling points for S11, 10 sampling points for S2 and 10 sampling points for S3. At the Tabalaopa stakeholder’s office, using Autocad software, each site was properly located. At each point, three composite soil samples were taken at 0-15, 15-30, and 30-50 cm depths. Hence, fifty-four soil samples were taken in S1, 36 soil samples in S11, 30 soil samples in S2 and 30 soil samples in S3, totaling 150 soil samples.

The soil was collected in plastic bags and transported to the lab. The samples were dried, ground and passed through a 0.355 mm sieve to remove rocks, roots and any larger particles. The digestion of the soil samples was accomplished with aqua regia following the Analysis protocol of Canada (MAF). The metal concentrations were determined using an Inductively Coupled Plasma-Optical Emission Spectrometry (ICP-OES) Perkin Elmer 2100. The pH and EC values were determined by a saturated paste using a standard glass electrode for pH (Hanna) and a conductivity meter for EC (Hanna), The OM was calculated following the standard methodology. Soil texture was calculated using the Boyoucous method.

The statistical model included the zone as factor A with three levels (0-15, 15-30 and 30-50 cm). The soil was collected in plastic bags and transported to the lab. The samples were dried, ground and passed through a 0.355 mm sieve to remove rocks, roots and any larger particles. The digestion of the soil samples was accomplished with aqua regia following the Analysis protocol of Canada (MAF). The metal concentrations were determined using an Inductively Coupled Plasma-Optical Emission Spectrometry (ICP-OES) Perkin Elmer 2100. The pH and EC values were determined by a saturated paste using a standard glass electrode for pH (Hanna) and a conductivity meter for EC (Hanna), The OM was calculated following the standard methodology. Soil texture was calculated using the Boyoucous method.

The statistical model included the zone as factor A with four levels (S1, S2, S3 and S11) and the soil profile as factor B with three levels (0-15, 15-30 and 30-50 cm).

Results and Discussion

Concerning parameter pH, ANOVA showed statistical differences for soil type (P=0.000) but no differences were found for soil depth and the interaction. It is well known that soil pH plays an important role in the mobility of metals as in their bioavailability for plants [28, 29]. Figure 1 shows the pH levels in the four types of soil and its noted that high pH values with an average of 8.74 were in S3. Minimum pH values were noted in S11 with 7.93 as well as in S1 with 7.80. In S2 soil, the pH was 8.41. It is also apparent that in the four soil types the pH was moderately alkaline. Additionally, a tendency of the pH to increase with depth was observed even though no
statistical differences were found. This tendency of the pH to increase below soil profile disagrees with the findings of Ilg et al. [30] which noted that pH values diminished with increasing depth after phosphorous fertilizations.

Figure 1: Soil pH in four different types of soil and three depths

The average pH value in the above soil profile (0-15 cm) was 8.11 followed by the 15-30 depth with 8.14 and profile 30-50 with a pH average of 8.18. These results concur with those presented by other researchers [31]. It is generally accepted and well documented that organic matter applications reduce pH levels [32-34] although Percival [9] did not find this effect in a soil amended with sewage sludge as did Lee et al. [35] who found an increase in the surface soil with manure compost applications. Moreover, He and Singh [36] discovered that organic matter applications reduced the pH levels while McLaren et al. [37] found a decrease in the pH levels of four soil types after applications of sewage sludge.

In this study, the minimum pH levels found in S1 and S11 may be explained by the fact that organic matter, in its initial phase forms organic acids; but with time, these acids are consumed or transformed and so the pH tends to increase [38]. Even so, other authors like Lee et al. [35] noted that pig litter applications to a soil in Taiwan tended to increase the pH in the upper soil horizons. These tests confirmed the results of Shen and Shen [39] who also noted pH increases in soil with pig compost applications. The results obtained by Adeli et al. [16] found that broiler litter applications increased the pH values. Interestingly, these researchers also determined that the pH values decreased in soil receiving only commercial fertilizing applications.

Similar to the results for soil pH, ANOVA detected statistical differences for parameter EC for soil type (P=0.000) but no differences were found for soil depth or the interaction. In Figure 2, the main effects of the EC values are observed, and it is evident that S11 obtained maximum levels while the minimum values were noted in S3 with a mean of 0.44 dSm⁻¹. It is well documented that OM may increase the levels of EC in soils and the results of this study showed the maximum amount of the OM in S1. Abbaspour et al. [40] reported an increase in levels of EC from 2.3 to 2.7 dSm⁻¹ when 50 mmol NaCl⁻¹ was applied to the soil. It is important to acknowledge that the four zones showed acceptable levels of salinity. The literature has reported that high salinity levels may have increased the metal mobility in soils. This process is complete when some cations like Na and K, substitute the heavy metal cations in the absorption places.

Figure 2: Levels of electrical conductivity (dSm⁻¹) in four different types of soil and three depths

Soil OM is a complex and heterogeneous medium that consists of both fractions; the humus fraction (humic acid, fulvic acid, humin) and the particulate fraction, known as particulate organic matter [41]. In this study, there were statistical differences in OM for soil type (P=0.000), soil depth (P=0.005) and for the interaction (P=0.014). OM is critical because of its tendency to form metal composite OM-metals [42, 43]. Previous studies have demonstrated that OM contains functional groups like –COOH and –OH that serve as a union of OM with metals [44, 45]. Other studies have exemplified the increment of OM levels in soils with the application of organic matter [46-49], resulting in the improvement of the soil structure [50]. Figure 3 identifies the different levels of OM, according to soil type. As expected, S1 and S11 obtained maximum levels of OM with means of 3.31 and 2.19 mg kg⁻¹ in comparison with S2 and S3, which presented means of 1.52 and 0.89 mg kg⁻¹, respectively. For S1 and S11, a negative linear tendency of OM content as affected by soil profile was clear. The maximum amount of OM was found in the 0-15 cm soil profile with a mean of 2.65 mg kg⁻¹, followed by the soil profile 15-30 cm with an OM mean of 2.21 mg kg⁻¹, and finally, the soil profile 30-50 cm with a mean of 1.75 mg kg⁻¹. It is essential to point out that OM has proven the most important soil component controlling the sorption and desorption of metals, being most clear this effect in the metal cadmium [51].
The K concentration was different for soil type (P=0.000) and no differences were found for soil depth or the interaction. Maximum K levels were noted in S1 with 359.3 mg kg\(^{-1}\) and S11 with 349.2 mg kg\(^{-1}\) when compared to levels of 321.8 mg kg\(^{-1}\) for S2 and 331.0 mg kg\(^{-1}\) for S3. Even though there were no statistical differences in soil depth, a decrease in K was observed in the way the soil horizon was lower. Lee et al. [35] in a study with pink compost applications, found the maximum K concentration in the higher soil horizons. The results of these authors agree with those of Bulluck et al. [52] who recorded K increments in the above soil profile after bovine cattle manure applications. In this research, soils S1 and S11 are obviously receiving this element in a dynamic form, hence their presence in the above strata.

The amount of Na was different for soil type (P=0.000) but no differences were noted for soil depth or the interaction. The highest level of this element was found in S3 with 203.0 mg kg\(^{-1}\) while the lowest amount was noted in S1 with 197.6 mg kg\(^{-1}\) and S11 with 197.8 mg kg\(^{-1}\). The Na levels in soil depth were similar and no tendency was noted, obtaining 199.7 mg kg\(^{-1}\) in the 0-15 cm depth, 199.6 mg kg\(^{-1}\) in the 15-30 cm depth and 198.9 mg kg\(^{-1}\) in 30-50 cm soil profile.

The levels of Cd were different for soil type (P=0.000) but not differences were noted for soil depth and the interaction. The lowest levels of Cd were in S2 with 1.62 mg kg\(^{-1}\) while maximum levels were obtained in S1 (4.88 mg kg\(^{-1}\)) and S3 (4.41 mg kg\(^{-1}\)). The Cd level found in S3 which identified a soil under natural condition is higher than the results presented by Rubio et al. [53] in a study of natural soil also in the state of Chihuahua. As well, the results obtained in the present study were higher than those presented in soils of India with levels of 0.37 mg kg\(^{-1}\) [54], England with 0.8 mg kg\(^{-1}\) [55], China with 0.07 mg kg\(^{-1}\) [56], Spain with amounts in the range of 0.4-0.8 mg kg\(^{-1}\) [57], Germany with levels in a range of 0.6-1.4 mg kg\(^{-1}\) [58] and Japan with 0.41 mg kg\(^{-1}\) [59]. Although no statistical differences were noted for soil depth, a slight decrement of Cd amounts in lower soil profiles was observed. These results may be explained by the fact that Cd is more soluble and mobile in soil when compared with other elements [60, 61]. Plus, OM levels in soil are the most important component that controls the absorption and desorption processes [51]. Yet, researchers like McLaren et al. [62] reported that Cd was the element less mobile when compared to other elements like Cr, Cu and Pb. These researchers explained their results by which OM influences the soil environment. Our original hypothesis was that differences will be evident in S1 and S3; one with wastewater irrigation and the other natural soil without any irrigation background. It was surprising that both soils presented similar levels of Cd. So, it is assumed that the Cd in the natural soil is coming from natural sources or due to anthropogenic pollution from sources other than irrigation. Near the study area, one of the larger smelter/refinery complexes in the world, named Avalos, operated for approximately 90 until the past several years when it ceased to operate, and we assumed that the amount of Pb in S3 might be explained by that fact. To date, there are no geological studies to acknowledge the Cd levels in the zone; however, the results of this study are higher than other results overseas (<1.0 mg kg\(^{-1}\)) for a mineral soil [63].

Figure 3: Soil OM in four different types of soil and three depths

![Figure 3](image-url)  
Figure 3: Soil OM in four different types of soil and three depths

The K concentration was different for soil type (P=0.000) and no differences were found for soil depth or the interaction. Maximum K levels were noted in S1 with 359.3 mg kg\(^{-1}\) and S11 with 349.2 mg kg\(^{-1}\) when compared to levels of 321.8 mg kg\(^{-1}\) for S2 and 331.0 mg kg\(^{-1}\) for S3. Even though there were no statistical differences in soil depth, a decrease in K was observed in the way the soil horizon was lower. Lee et al. [35] in a study with pink compost applications, found the maximum K concentration in the higher soil horizons. The results of these authors agree with those of Bulluck et al. [52] who recorded K increments in the above soil profile after bovine cattle manure applications. In this research, soils S1 and S11 are obviously receiving this element in a dynamic form, hence their presence in the above strata.

The amount of Na was different for soil type (P=0.000) but no differences were noted for soil depth or the interaction. The highest level of this element was found in S3 with 203.0 mg kg\(^{-1}\) while the lowest amount was noted in S1 with 197.6 mg kg\(^{-1}\) and S11 with 197.8 mg kg\(^{-1}\). The Na levels in soil depth were similar and no tendency was noted, obtaining 199.7 mg kg\(^{-1}\) in the 0-15 cm depth, 199.6 mg kg\(^{-1}\) in the 15-30 cm depth and 198.9 mg kg\(^{-1}\) in 30-50 cm soil profile.

The levels of Cd were different for soil type (P=0.000) but not differences were noted for soil depth and the interaction. The lowest levels of Cd were in S2 with 1.62 mg kg\(^{-1}\) while maximum levels were obtained in S1 (4.88 mg kg\(^{-1}\)) and S3 (4.41 mg kg\(^{-1}\)). The Cd level found in S3 which identified a soil under natural condition is higher than the results presented by Rubio et al. [53] in a study of natural soil also in the state of Chihuahua. As well, the results obtained in the present study were higher than those presented in soils of India with levels of 0.37 mg kg\(^{-1}\) [54], England with 0.8 mg kg\(^{-1}\) [55], China with 0.07 mg kg\(^{-1}\) [56], Spain with amounts in the range of 0.4-0.8 mg kg\(^{-1}\) [57], Germany with levels in a range of 0.6-1.4 mg kg\(^{-1}\) [58] and Japan with 0.41 mg kg\(^{-1}\) [59]. Although no statistical differences were noted for soil depth, a slight decrement of Cd amounts in lower soil profiles was observed. These results may be explained by the fact that Cd is more soluble and mobile in soil when compared with other elements [60, 61]. Plus, OM levels in soil are the most important component that controls the absorption and desorption processes [51]. Yet, researchers like McLaren et al. [62] reported that Cd was the element less mobile when compared to other elements like Cr, Cu and Pb. These researchers explained their results by which OM influences the soil environment. Our original hypothesis was that differences will be evident in S1 and S3; one with wastewater irrigation and the other natural soil without any irrigation background. It was surprising that both soils presented similar levels of Cd. So, it is assumed that the Cd in the natural soil is coming from natural sources or due to anthropogenic pollution from sources other than irrigation. Near the study area, one of the larger smelter/refinery complexes in the world, named Avalos, operated for approximately 90 until the past several years when it ceased to operate, and we assumed that the amount of Pb in S3 might be explained by that fact. To date, there are no geological studies to acknowledge the Cd levels in the zone; however, the results of this study are higher than other results overseas (<1.0 mg kg\(^{-1}\)) for a mineral soil [63].

Figure 4: Soil Pb levels in four different types of soil and three depths

Figure 4 shows the main effects of Pb in the four soil types and in the three depths. ANOVA detected statistical differences for soil type but not differences for soil depth and the interaction. In Figure 4, it is evident that the greatest amount of Pb was in S11 with a mean of 155.83 mg kg\(^{-1}\). Contrary to our original hypothesis, S2 obtained the lowest Pb concentration with a mean of 68.47 mg kg\(^{-1}\) followed by S3 with 98.75 mg kg\(^{-1}\). The results in S2 might be explained by the lixiviation of the element in soil due to precipitation events after wastewater irrigation ceased [64, 65]. The presence of Pb in S3 may be explained by the background of the smelter/refinery complex (Avalos); in fact, Ornelas et al. [66] reported values of 400 ppm of this element in the vicinity of the smelter/refinery complexes (Avalos), operated for approximately 90 until the past several years when it ceased to operate, and we assumed that the amount of Pb in S3 might be explained by that fact. To date, there are no geological studies to acknowledge the Cd levels in the zone; however, the results of this study are higher than other results overseas (<1.0 mg kg\(^{-1}\)) for a mineral soil [63].
accumulation of this metal in soils even far from the source [69]. For instance, Billett et al. [70] found significant increments in forestry soils where the contamination source was about 120 km away. Ultimately, the levels of Pb reported in this study must be regarded as high, when considering that a normal soil contains approximately 10 mg kg⁻¹, but some contaminated soils may contain 1,000 mg kg⁻¹ [71]. A tendency of the Pb levels to decrease with depth was observed. These results support the hypothesis proposed by Miller and Friedland [72] which established that colloidal particles are of great importance in the Pb movement from the upper to lower soil horizons. This study has noted that all levels of OM were higher in the upper soil horizon and possibly, the levels in the lower soil horizons are a result of this movement [73]. The concentrations obtained in this study are higher than those reported by Rubio et al. [53] in Chihuahua, Mexico; likewise, in soils of India with 10.4 mg kg⁻¹ [54] and Japan with 21 mg kg⁻¹ [59]. It must be known that high levels of Pb in soils may produce a significant decrement on the enzymatic activity, meaning that this metal has a toxic effect in the different biochemical reaction occurring in the soil environment [22, 74-76]. Accordingly, plant productivity and microbiological communities in the soil environment decrease [77].

In the particular case of Ni, ANOVA detected statistical differences by soil type (P=0.000) but no differences were found for soil depth or the interaction (Figure 5). Again, the highest level of Ni was observed in the 30-50 cm depth (9.81 mg kg⁻¹) with the exception of soil S3. These results are consistent with those of McLaren et al. [62] who found that about 57% of Ni applications were lixiviated from upper soil horizons to lower soil horizons. The lowest Ni concentrations were detected in S2 with a mean of 7.45 mg kg⁻¹ and S11 with a mean of 8.86 mg kg⁻¹.

![Figure 5: Soil Ni levels in four different types of soil and three depths](image)

The Cr level was superior in S3 with a mean of 44.27 mg kg⁻¹ while the lowest values were noted in S2 with a mean of 37.10 mg kg⁻¹. Intermediate values were observed in S1 with 40.06 mg kg⁻¹ and S11 with 38.74 mg kg⁻¹. It was clear that Cr has a tendency to be increased with depth.

Contrary to the findings of Ni and Cr, the lowest concentrations of Cu were detected in S3 and S2 with means of 7.17 and 10.6 mg kg⁻¹, respectively, while maximum values were noted in S1 with 51.35 mg kg⁻¹ and S11 with 38.61 mg kg⁻¹. Li et al. [78] reported that levels of Cu in soil above 10-25 mg kg⁻¹ might reduce plant productivity. Thus, S1 and S11 might prove toxic for plant production. Besides, the Cu concentration has a tendency to increase with depth. This tendency agrees with a report by Hu et al. [79] that found Cu concentrations in a range of 0.01 to 0.26 mg L⁻¹ in the 20 cm depth and of 0.01 to 0.33 mg L⁻¹ in the 50 cm depth. In other study, McLaren et al. [62] mentioned that Cu soil applications have certain mobility from upper horizons to lower horizons. These authors specified that Cu, Cr and Pb had a mobility of 35% from the above soil profile to deeper soil profiles. These results also agree with Dowdy et al. [80] which speculated that metals could be transported from upper zones to lower zones through macropores present in soils. For this reason, most of the metals evaluated in this research have a tendency to increase in lower horizons and coincide with other studies previously reported [81, 82].

Fe level was different due to soil type (P=0.000) but no differences were found for soil depth or the interaction. The highest levels of Fe were noted in S3 with 20,313.0 mg kg⁻¹ while the lowest amount was observed in S2 with a mean of 17,865.33 mg kg⁻¹. As noted in the other evaluated metals, the levels of Fe also increased with depth. The B concentration was different due to soil type (P=0.000) but no differences were found for soil depth and the interaction. The S3 showed the highest amount of this element with a mean of 41.47 mg kg⁻¹, while the lower amount was detected in S2 with a mean of 14.70 mg kg⁻¹. In contract with other elements, B levels were superior in the upper soil horizon.

Conclusions and Recommendations

The analysis demonstrated that among the variables, soil type was the only showing a statistical difference, which indicates that the resulting concentrations can be largely explained by the type of irrigation the soil had at the time. It was noted that concentrations of nickel, chromium, copper, iron and boron concentrated in deeper soil layers while potassium, sodium, cadmium, and lead showed the opposite effect.

In areas where elements were expected to be present in lesser concentrations as in the case of S3, the opposite effect was observed with respect to S1, S11 and S2. Instead, sodium, cadmium, chrome, iron and boron, showed higher concentrations in S3, which is contradictory to the established hypothesis for being an area lacking in irrigation. This may be explained by natural concentrations of said elements or, in the case of cadmium, by airborne contamination from the Avalos smelter. It was also noted that organic material is an important variable and that it can influence the mobility of
metals in those areas where high concentrations such as S1 and S11, which coincide with constant irrigation. Clearly, the area has been constantly exposed to certain health hazardous metals. More attention is recommended, even though at this time a wastewater treatment plant has been built and part treated water is used to irrigate the crops. Still, residual water is used for forage irrigation, and this practice could generate illnesses in the population near the study area.

Acknowledgments: We are deeply grateful with the National Committee of Science and Technology of Mexico (CONACYT) which provided a grant for carrying out the research reported here.

References


78. Li, G. C.; Wang, Y. P.; Chang, J. M.: Heavy metals concentrations in soils of Taiwan. Publication by the Environmental Protection Administration (EPA) of Taiwan. Taipei, Taiwan. 1987.

79. Hu, N.; Luo, Y.; Longhua, W.; Song, J.: A field lysimeter study of heavy metals movement down to the profile soils with a multiple metal pollution during

