



Application of XAFS and XRD methods for describing the copper and zinc adsorption characteristics in hydromorphic soils

Tatiana V. Bauer · David L. Pinskii · Tatiana M. Minkina ·
Victoria A. Shuvaeva · Alexander V. Soldatov · Saglara S. Mandzhieva ·
Victoria S. Tsitsuashvili · Dina G. Nevidomskaya · Ivan N. Semenov

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Abstract Modeling metal sorption in soils is of great importance to predict the fate of heavy metals and to assess the actual risk driven from pollution. The present study focuses on adsorption of HM ions on two types of hydromorphic soils, including calcaric fluvisols loamic and calcaric fluvic arenosols. The individual and competitive adsorption behaviors of Cu and Zn on soils and soil constituents are evaluated comprehensively. It is established that the sorption processes were best described with the Langmuir model. The results suggest that the calcaric fluvic arenosols are more vulnerable to heavy metal input compared to fluvisols loamic. In all cases, Cu had a

higher range of values of the adsorption process parameters relative to Zn. The Zn is likely to be the most critical environmental factor in such soils since it exhibited a decreased sorption under competitive conditions. The retention mechanisms of HM in hydromorphic soils are considered. Based on theoretical calculations of ion activity in soil solutions using solubility diagrams of Cu and Zn compounds, the possibility of precipitation of Cu hydroxide and Zn carbonate in the studied soils is shown. Direct physical methods of nondestructive testing (XAFS and XRD) are applied to experimentally prove the formation of these HM compounds on the surface of montmorillonite, the dominant mineral in hydromorphic soils, and calcite. Thus, the combination of both physicochemical methods and direct physical methods can provide a large amount of real information about the mechanisms of HM retain with solid phases.

T. V. Bauer
Federal Research Centre, The Southern Scientific Centre
of the Russian Academy of Sciences,
Rostov-on-Don 344006, Russian Federation

D. L. Pinskii (✉)
Institute of Physicochemical and Biological Problems of
Soil Science, Russian Academy of Sciences, Institutskaya
str, 2, Pushchino, Pushchino 142290, Russian Federation
e-mail: pinsky43@mail.ru

T. M. Minkina · V. A. Shuvaeva ·
A. V. Soldatov · S. S. Mandzhieva ·
V. S. Tsitsuashvili · D. G. Nevidomskaya
Southern Federal University, Rostov-on-Don 344006,
Russian Federation

I. N. Semenov
Lomonosov Moscow State University, Moscow 119991,
Russian Federation

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Introduction

Heavy metals (HMs) are hazardous elements and well known to impose a threat to human health. Among

HMs, Cu and Zn are the priority pollutants included in the US EPA list (Code of Federal Regulations 2019). As a result of growing ecological awareness about toxic HM globally, it has been recognized that there is a need for improving the protection of public health and the environment due to continuous exposure risks, even in small daily doses (Pirooty and Ghasemzadeh 2013). A high concentration of Cu can exert a negative influence on organisms, followed by destroying the cell metabolism and provoking immunotoxicity, embryotoxicity, and reproductive toxicity (Duan et al. 2016). Exposure to elevated levels of Zn can adversely affect the gastrointestinal, hematological, and respiratory systems. The high level of exposure can negatively alter the function of the cardiovascular and neurological systems in the humans (Radhakrishnan et al. 2016). High concentrations of Zn^{2+} ions are known to trigger cell death by apoptosis or necrosis in a variety of cell types in the brain (Nriagu 2019).

The excessive concentration of HM in soils can result from different sources. Fertilizers and agrochemicals products, mining-related industries, and waste incineration are among the contributing sources (Janoš et al. 2010). Adsorption is a major process responsible for the accumulation of HM. The study of the sorption capacity of soils relative to HM is important because it affects key ecological issues, such as prediction of the risk related to the bumping of metals in soil, remediation of polluted soils, and utilization of disposal (Vidal et al. 2009). In a broad sense, the sorption of HM includes diverse adsorption processes, e.g., the formation of extra- and intraspheric complexes and ion exchange, as well as surficial sedimentation and other mechanisms (McBride 1994). The sorption capacity of the soil is influenced by pH values and the presence of active adsorption centers in different soil components, such as carbonates, phosphates, organic matter, silicates, and (hydr)oxides (Sipos et al. 2008; Vidal et al. 2009; Fisher-Power et al. 2016). Clay minerals prevail in soil particles less than 0.002 mm in diameter. Since the negatively charged layered minerals dominate in the soil surface, they are characterized by a significant sorption capacity relative to HM (Plyaskina and Ladonin 2005). The adsorption and precipitation of Cu in soil carbonates play an important role in metal fixation because of the high dispersion of soil carbonates and the small solubility constant of carbonates of HM (Jinren and Weiling 2003; Burachevskaya et al.

2018). Assessment of the pollutants sorption by soil is usually based on laboratory experiments to quantitatively estimate the coefficient of metal distribution between the solid and liquid phases of soil (Vidal et al. 2009). By measuring the aqueous and adsorbed concentration of the contaminant at adsorption equilibrium, adsorption isotherms and related parameters can be derived (Mizutani et al. 2017). This simple approach can be expanded based on additional mechanisms, such as desorption, competition for adsorption sites in the case of polymetallic pollution, and heterogeneity of exchange sites in the solid phase of soil (Ding et al. 2009). The chemical and energetic heterogeneity in the surface of soil particles leads to irregular strength of the HM fixation and dependence of the interaction energy on the degree of the surface filling by the adsorbed cations and their individual properties (Saha et al. 2002).

Soil pollution in industrially developed regions takes place simultaneously with a wide range of chemicals (Linnik et al. 2019). However, the polyelemental character of contamination is rarely considered in the investigation due to mutual competition between cations for exchange sites, aggravated by specific properties of metals and heterogeneity of the adsorbent.

The formation of sediments of the low-soluble HM compounds is an important mechanism for the HM immobilization. Copper can be precipitated as $Cu(OH)_2$, CuO , and $Cu_2CO_3(OH)_2$, Zn, as ZnO , $ZnCO_3$, and $Zn_5(OH)_6(CO_3)_2$ (Madrid and Diaz-Barrientos 1992). Sophisticated analytical techniques such as X-ray powder diffraction (XRD), X-ray absorption near edge structure (XANES), and extended X-ray absorption fine structure (EXAFS) can be used to identify the HM-resistant compounds in soils (Nevidomskaya et al. 2016).

Being on the way of the chemical elements flow into water bodies, floodplain soils act as barriers and accumulators of not only vital elements but also pollutants while performing a protective function in valley landscapes. A feature of the soils of river floodplains is their dependence on frequently changing environmental conditions associated with the features of the relief, hydrological conditions, the dynamics of alluvial deposition, the variable nature of moisture, and the composition of vegetation. For this reason, the floodplain soils are in particular interest at the study of the HM absorption mechanisms. The complex process

of soil formation, its high dynamism, and the specificity of water nutrition are responsible for a poorly study of the absorption capacity of floodplain soils relative to HM.

The aim of the present paper is to study mechanisms of Cu^{2+} and Zn^{2+} adsorption by the main types of hydromorphic soils in the Rostov region (Russia).

Materials and methods

Soil sampling

The objects of study were the hydromorphic soils of the Lower Don basin (Rostov region, Russia). The soils were classified as meadow heavy loamy soils (calcaric fluvisols loamic) on alluvial deposits and alluvial sandy soils (calcaric fluvic arenosols). Soils of both types were selected in the areas without anthropogenic impact, located in the floodplain of the Seversky Donets River, a large right-bank tributary of the Don. The absence of high concentrations of HM, including Zn and Cu (Table 1) in the natural soils, was shown by our previous studies (Konstantinova et al. 2020).

The calcaric fluvisols are located in the central part of a floodplain meadow on the left bank of the Seversky Donets River (N 48°21'06.67", E 40°14'09.57"). The calcaric arenosols are located at the edge of the floodplain on the right bank of the Seversky Donets River (N 48°20'12.91", E 40°14'10.89"). The soil samples were taken from the surface layer (0–20 cm), dried at room temperature, ground and sieved (1 mm mesh) to remove residual, such as visible stones and plants, and then stored at 4 °C until further use. The studied soils differ

from each other according to physicochemical parameters (Table 1).

The mineralogical composition of the clay fraction in hydromorphic soils is represented by mixed-layer mica–smectite formations with a high content of smectite packets (59–80%), hydromica (17–26%), kaolinite and chlorite (5–12%). An X-ray diffractometry using DRON-7 diffractometer in Cu-K α -filtered radiation is used to determine the mineralogical composition of the clay fraction of soils with a size less than 1 μm . The soil particle size distribution analysis was performed according to the pipette method with pyrophosphate procedure of soil preparation (ISO 13,317–2 2001). Soil properties were determined using standard methods: pH of soil suspension in water by ISO 10,390 (2005); total organic carbon (TOC) content by sulfochromic oxidation, ISO 14,235 (1998); carbonates content by a Scheibler apparatus, ISO 10,693(1995) , and exchangeable cations by hexaminecobalt trichloride solution, ISO 23,470(2011) . The total content of Zn in the soils was determined by X-ray fluorescent (XRF) scanning spectrometer SPECTROSCAN MAKC-GV.

Heavy metal adsorption procedure

Air-dried aliquots (5 g) of the studied soil samples were flushed with 50 ml of metal solutions ($\text{Cu}(\text{NO}_3)_2$ and $\text{Zn}(\text{NO}_3)_2$) with the following Cu^{2+} and Zn^{2+} concentrations (mM L^{-1}): 0.05, 0.08, 0.1, 0.3, 0.5, 0.8, and 1.0. The metal solutions were prepared from $\text{Cu}(\text{NO}_3)_2 \times 3\text{H}_2\text{O}$ and $\text{Zn}(\text{NO}_3)_2 \times 6\text{H}_2\text{O}$ salts of the ‘chemically pure grade.’ The HMs were added separately or simultaneously. The resulting suspensions were stirred for 1 h and then left in the calm state for 24 h. This period was sufficient for the adsorption to reach equilibrium. The time required for the system

Table 1 Physicochemical properties of the studied soils

Soil	pH of water	TOC percent (%)	CaCO ₃	Physical clay (particle size < 0.01 mm)	Clay (particle size < 0.001 mm)	Exchangeable Ca ²⁺ + Mg ²⁺ , cmol(+)/kg	Cu mg/kg	Zn mg/kg
Calcaric fluvisols	7.5	4.3	0.6	55.8	32	38.1	37	110
Calcaric fluvic arenosols	7.5	0.9	0.1	2.8	1.6	6.6	15	85

to reach equilibrium was in good agreement with the results of our kinetic adsorption experiments (Pinskii et al. 2010). The studies were carried out with a change in the ion strength from 0.0036 to 0.87 mol/L. The pH values were measured by potentiometry of the equilibrium solutions. Then, the suspension was centrifuged and filtered. The metal content in filtrate was determined by the atomic absorption spectrophotometer with electrothermic atomization and polarization Zeeman effect correction for the nonselective adsorption using an MGA-915MD spectrometer (Lumex Scientific-Production Company, St. Petersburg, Russia). The amount of metal adsorbed by the solid phase was found by the eliminating of the added amount from the measured concentration in the equilibrium solution (Du et al. 1999):

$$C_{ad} = \frac{(C_i - C_s) \cdot V}{m} \quad (1)$$

where C_{ad} is the amount of metal adsorbed by a unit sorbent mass, mM kg^{-1} ; C_i is the initial metal concentration in solution, mM L^{-1} ; C_s is the equilibrium metal concentration in solution, mM L^{-1} ; V is the solution volume, ml; and m is the air-dry sorbent mass, g. The obtained data were used to draw the C_s - C_{ad} plot, where C_s is the equilibrium concentration of metal added to the solution, mM L^{-1} ; and C_{ad} is the specific content of metal in the sorbent phase, mM kg^{-1} . The multi-solute competitive sorption experiments were conducted in the same manner as in the single-solute sorption experiments. The corresponding sorption isotherms for each metal were quantitatively described by parameters through fitting the experimental data to the Langmuir isotherm described by the following equations, respectively:

$$C_{ad} = \infty K_L C_e / (1 + K_L C_e) \quad (2)$$

where C_{ad} is the amount of adsorbed cations; C_{∞} is the maximum metal adsorption, mM kg^{-1} ; C_e is the equilibrium metal concentration, mM L^{-1} ; and K_L is the Langmuir constant, L mM^{-1} .

The Langmuir isotherm enables us to calculate the maximum adsorption capacity (C_{∞}) of each metal on the solid phase and to quantify the metal affinity with soil (K_L). The Langmuir model assumes monolayer coverage in a homogeneous adsorption mode occurred on a surface with binding sites having equal energies (Musso et al. 2014).

Adsorption mechanisms

The components of the mineral phases were used to study the mechanisms of HM immobilization in the soils. The montmorillonite sample and calcite were saturated with Cu^{2+} and Zn^{2+} ions using a saturated solution of metal nitrate for 1 week (Nevidomskaya et al. 2016). The calcite sample isolation from the studied fluvisols was carried out under MBS-10 binoculars in the magnification range from 5 to 32. The montmorillonite sample is a standard reference material (GSO 8694–2005).

X-ray powder diffraction

The X-ray diffraction based on synchrotron radiation was used for the analysis of the reference samples (Nevidomskaya et al. 2016). The structural accordance of montmorillonite mineral phase was compared to the diffraction patterns obtained at the Structural Materials Science beamline of the Kurchatov Center for Synchrotron Radiation (Chernyshov et al. 2009). A 1.7 T bend magnet of the Siberia-2 storage ring is the source of synchrotron radiation. The electron beam energy is 2.5 GeV; the average current is 120 mA.

Diffraction studies of monochromatic synchrotron X-ray radiation ($\lambda = 0.68886 \text{ \AA}$, Si monochromator) were performed in transmission geometry using a Fujifilm Imaging Plate two-coordinate detector at 0.68886 \AA . X-ray diffraction patterns were recorded in integrated mode at $20 \text{ }^\circ\text{C}$. The time of sample exposure was about 15 min. A silicon standard (NIST SRM 640C) was used for the angular calibration of the scale. Compared to the conventional X-ray diffractometry, the use of high-intensity monochromatic synchrotron radiation significantly improves the intensity and resolution of diffraction patterns once it is combined with a two-coordinate detector and a Si monochromator. The diffraction patterns were compared with the ICSD (Inorganic Crystal Structure Database) (<https://icsd.products.fiz-karlsruhe.de/>).

X-ray absorption spectroscopy

The XANES spectra at the K-edge of Zn (9659 eV) of the carbonate were obtained by spectrometer Rigaku R-XAS Looper in the fluorescence regime (Nevidomskaya et al. 2014). The zinc nitrate ($\text{Zn}(\text{NO}_3)_2$) was

used as a reference compound. The XANES spectra first derivatives were calculated. This calculation is a means to obtain and identify detailed information on the state of Zn^{2+} ions in the studied sample (Nevidomskaya et al. 2014).

Statistical analysis

The SigmaPlot 12.5 software packages with a confidence coefficient of 0.95 were used for the statistical process. Each experimental isotherm point was fixed after three replications. For each sample analyzed through the X-ray absorption spectroscopy method, ten scans were collected and averaged to achieve a higher signal-to-noise ratio. For energy calibration, copper foils were used as standard samples. For statistical purposes, the 60-s exposure time was considered to collect the data for each point in the spectrum. For the X-ray powder diffraction analysis, each of the studied samples was examined in three replications to scrutinize the homogeneity of the fraction composition.

Results and discussion

Monoelement adsorption of HM

Figure 1 presents the isotherms of Cu^{2+} and Zn^{2+} adsorption by the calcaric fluvisols and calcaric fluvic arenosols during the separate input of metals from nitrate salt solutions. In all cases, adsorption of metals in the studied soils is described by Eq. (2) with a high

degree of reliability ($R^2 = 0.99$). Experimental data on Cu^{2+} and Zn^{2+} adsorption by the studied soils revealed some discrepancies in their adsorption capacity due to differences in chemical properties of these elements and between the soils themselves. Both soil types show a higher affinity to Cu^{2+} rather than Zn^{2+} , as suggested by the shape of isotherms and their location near the y-axis. Isotherms of Cu^{2+} and Zn^{2+} adsorption by the fluvisols and arenosols have a typical Langmuir shape, indicating the metal adsorption by one mechanism or several mechanisms weakly distinguished from each other. The values of Langmuir parameters computed with the linearization of experimental isotherms are presented in Table 2.

It has been established that $C_{\infty}(Cu) > C_{\infty}(Zn)$ and the value of C_{∞} for the fluvisols is always higher than for the calcaric fluvic arenosols. Langmuir constants (K_L) characterizing the energy of interaction between the adsorbed cations and SAC are also always higher than for the fluvisols. One of the reasons responsible for a higher absorption capacity of the fluvisols (Loamic) is related to the high content of organic matter, clay, and silt, relative to the arenosols, in which the TOC content is just 0.9%. Discrepancies between K_L values for the studied Cu^{2+} and Zn^{2+} cations are 17–27 times higher depending on the soil type. Hence, the properties of the metal cation exert a significant influence on the energy of interaction with the soil particle surface (Jalali, Moharrami 2007; Pinskii et al. 2014; Huang et al. 2018). The metal hydrolysis properties affect the retention tendency in a particular way. In this context, Cu^{2+} has more tendency to be retained compared to Zn^{2+} (Saha

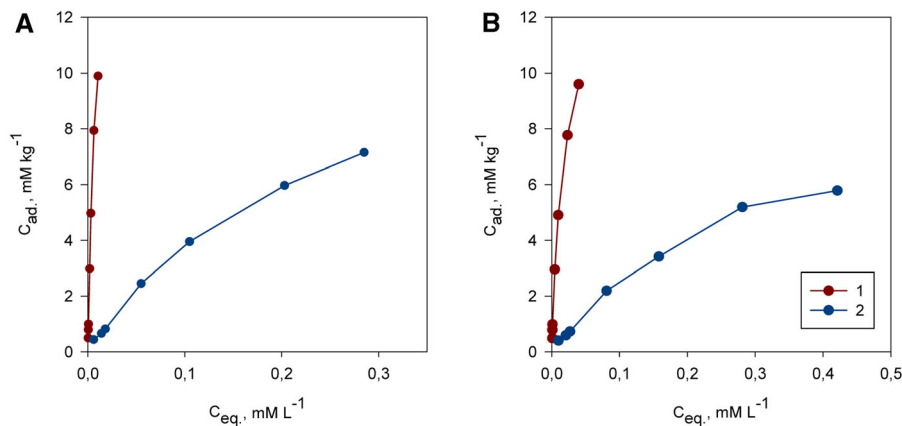


Fig. 1 Isotherms of the monoelement HM adsorption by the calcaric fluvisols a and calcaric fluvic arenosols b: (1) Cu, (2) Zn

Table 2 Parameters of HM adsorption by the hydromorphic soils during the monoelement and polyelement pollution

Variation	Calcaric fluvisols			Calcaric fluvic arenosols		
	C_{∞} , mM kg ⁻¹	K_L , l mM ⁻¹	R^2	C_{∞} , mM kg ⁻¹	K_L , l mM ⁻¹	R^2
<i>Monoelement pollution</i>						
Cu	19.2 ± 1.7	105.8 ± 16.3	0.99	13.8 ± 0.3	57.1 ± 3.0	0.99
Zn	13.5 ± 0.5	4.0 ± 0.3	0.99	10.1 ± 0.8	3.4 ± 0.5	0.99
<i>Polyelement pollution</i>						
Cu	12.3 ± 1.3	126.7 ± 33.5	0.98	9.9 ± 0.4	64.0 ± 8.3	0.99
Zn	5.6 ± 0.3	11.6 ± 1.9	0.99	4.0 ± 0.2	8.3 ± 1.2	0.99

et al. 2002; Agbenin and Olojo 2004). In the more alkaline pH range, the influence of the metal hydrolysis on metal adsorption becomes dominant (Jalali and Moharrami 2007). The high electronegativity of metal leads to its strong interaction with the soil particle surface (Pinskii et al. 2014). Electronegativity is an essential contributing factor to determine the level of preference in which the HM tends to chemisorb with the highest order: Cu > Ni > Co > Pb > Cd > Zn > Mg > Sr (McBride 1994).

The obtained Misono softness parameter predicts the metal affinity sequences or preferential retention by soil components (Misono et al. 1967). The softness parameter is an index to determine the affinity of a metal to form covalent bonds with colloids (Sposito 1989). This parameter considers both the ionic charge and the ionization potential (which is higher for Cu than Zn) (Agbenin and Olojo 2004). Thus, the preferential retention of Cu to Zn is caused by differences in electronegativity, first hydrolysis constants, and the softness parameter.

Variation of pH during the monoelement adsorption of HM

To estimate discrepancies in Cu²⁺ and Zn²⁺ adsorption by soils and elucidate possible mechanisms of the interaction of metals with them, we accomplished an experiment with the determination of pH values in the equilibrium and corresponding initial solutions.

The dissolution of salts Cu(NO₃)₂ and Zn(NO₃)₂ provokes variation of pH values in the initial solutions due to processes of hydrolysis (Fig. 2). With increasing solution concentration, the pH value in the initial Cu(NO₃)₂ solution decreases gradually from 6.2 to 3.9; in the Zn(NO₃)₂ solution, from 6.4 to 5.9. In general, the pH value in the initial solutions at different amounts of the adsorbed Cu is much lower

relative to Zn. In the calcaric fluvisols, the pH value varies from 7.4 to 6.6 for Cu and 7.4 to 7.1 for Zn. In equilibrium solutions of the calcaric fluvic arenosols, the pH value decreases from 7.4 to 6.4 and from 7.4 to 6.8, for Cu and Zn, respectively.

The contact of initial solutions with soil leads to two effects: (1) abrupt change of pH because of the buffer capacity of soil related to its composition, composition of the added solution, and properties of the adsorbed cation; (2) subsequent gradual decrease in pH in equilibrium solutions with the increase in adsorbed HM cations. The nature of the dependence of pH from the adsorbed metal amount is governed mainly by the composition of the solid phase: the presence of carbonates and hydrogen, which can exchange with HM cations. Analogous reasons for the pH variation in solutions resulted from sorption processes in soils were described in Mesquita and Silva (2002). Certain influence can also be exerted by processes taking place in the liquid phase during its compositional change due to the adsorption of HM cations and the transition of other cations into the solution. If the amount of adsorbed HM is close to the maximum adsorption value, the discrepancy between pH values in the initial and final solutions can reach 2.5–3.0 units and persist for a long time (Davis and Leckie 1978).

Displacement of the exchangeable hydrogen during the adsorption of HM cations, probably, plays an important role in the acidification of equilibrium solutions. Its presence in soils is related, first of all, to weakly acid exchange centers in the solid phases (Pinskii et al. 2018). A decrease in pH values can also be related to the interaction of HM hydroxo complexes with the soil mineral particles, leading to the completion of octahedral layer and the expulsion of hydrogen ions into the solution (Ladonin 1997).

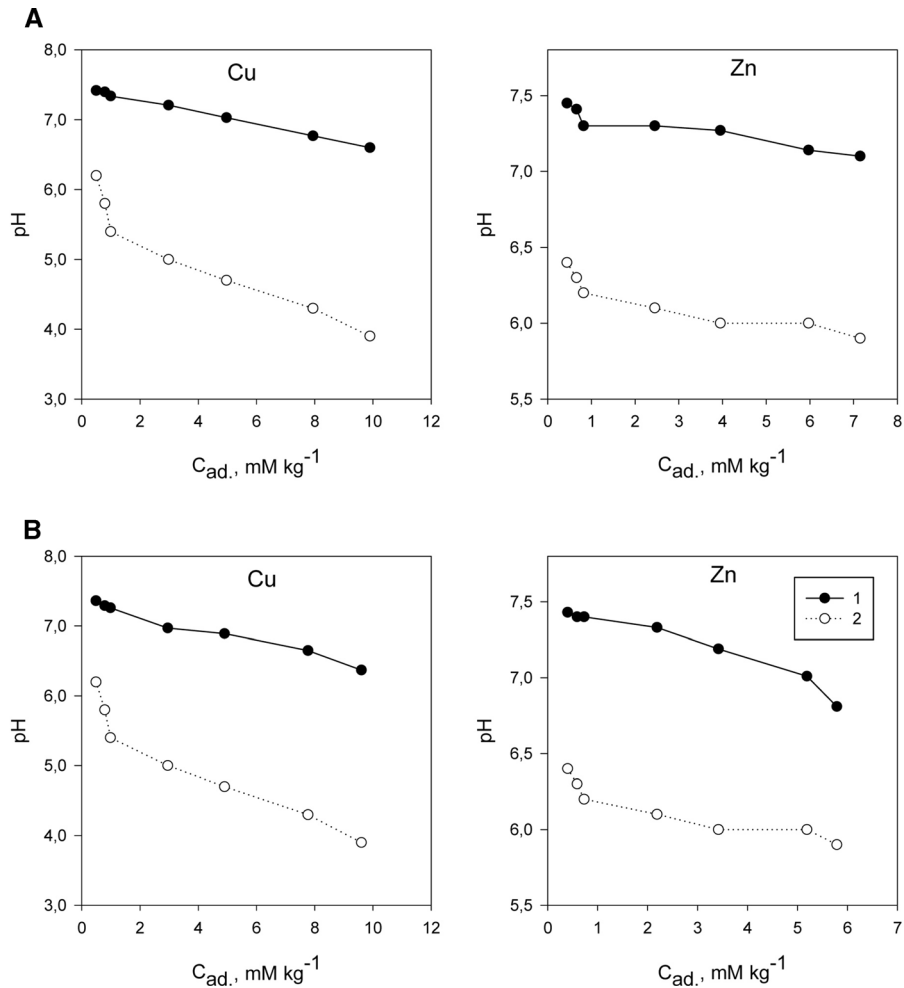


Fig. 2 Variation of pH equilibrium (1) and respective initial solutions (2) at HM adsorption by the calcaric fluvisols **a** and calcaric fluvic arenosols **b** from nitrate salt solutions

The addition of Cu^{2+} salts to the soil provoked a notable decrease in pH value in the equilibrium solution, relative to the addition of Zn^{2+} salt (Fig. 2). Copper can be absorbed more firmly by soils, and, probably, this metal has a greater capacity to displace the exchangeable hydrogen from the solid phase into the solution. Adsorption of Zn by the studied soil types is characterized not only by lesser values of parameters C_{∞} and K_L , but also by least variations of pH values, probably, because of its lower capacity to complex formation. A positive correlation between the formation of hydroxo complexes of HMs and the decrease in pH values in the soil–solution system is reported in McBride (1989).

Competitive adsorption of heavy metals

Figure 3 presents the isotherms of Cu^{2+} and Zn^{2+} adsorption by the calcaric fluvisols and calcaric fluvic arenosols during their combined addition to the studied soil. As in the case of monoelement adsorption, the isotherms are described well by Eq. (2). However, peculiarities of the interaction are influenced considerably by the competition for sorption sites (Table 2). Adsorption of Cu^{2+} by soils is much more intense relative to Zn^{2+} .

The presented data show a decrease in C_{∞} values for both cations. Moreover, the sum of C_{∞} (Cu) and C_{∞} (Zn) during the joint adsorption is equal within the error limit to mean C_{∞} values during the monoelement adsorption of Cu and Zn from each soil. The presence

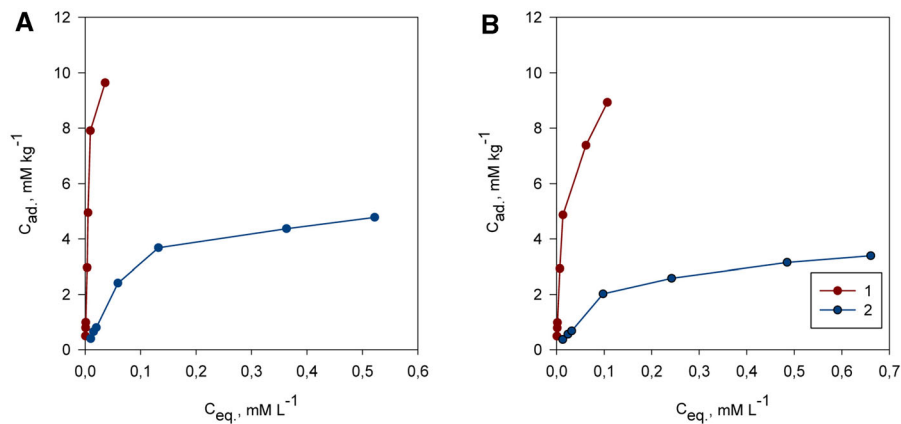


Fig. 3 Isotherms of the competitive HM adsorption by the calcaric fluvisols **a** and calcaric fluvic arenosols **b**: (1) Cu, (2) Zn

of both metal cations in the system is reflected mostly in the Zn adsorption, with C_{∞} decreasing 2.4–2.5 times. For example, the adsorption capacity of Kastanozems decreases drastically during the polyelement contamination: Pb is absorbed 5 times less, Cu 2 times less, and Zn 4 times less (Panin and Siromlya 2005). Hence, the two cations compete for the same sites. At the same time, K_L values in the Langmuir equation during the polyelement adsorption increase, relative to the process of monoelement adsorption.

Thus, adsorption sites in the studied soils are heterogeneous in terms of energy. Under the influence of competition from other ions, ions of each metal interact ultimately with reaction centers characterized by the maximum affinity (Ladonin 1997; Pinskii et al. 2010). Under conditions of competition, Cu occupies a dominant position and interacts with adsorption centers characterized by the maximum affinity. As a result of the polyelement adsorption, the amount of each adsorbed metal will decrease, while the strength of its bond with the pedogenic adsorption complex will increase.

Processes of the sedimentation of HM ions in soil solutions

The formation of sediments of the low-solubility compounds can also influence the behavior of Cu and Zn cations in the studied systems. It is often challenging to separate adsorption from precipitation in the experiments. To this end, the sorption is used to confirm the declining trend of solute in the liquid phase, which is resulted from adsorption and

sedimentation or polymerization process (Jinren and Weiling 2003). Based on the formation of aluminum orthophosphate, the possibility of the formation of sediments of the low-soluble compounds during adsorption was demonstrated by Sposito (1984). In this case, the sedimentation equation is identical to the Langmuir equation. Thus, both processes become experimentally indiscernible.

The possibility of the formation of the low-solubility Cu–Zn carbonate and hydroxide sediments was studied in the floodplain soils based on the solubility diagrams (Fig. 4). Figure 4 also shows logarithms of the concentrations of these metals in equilibrium solutions of the calcaric fluvisols and calcaric fluvic arenosols obtained during the study of the cations adsorption. Computations were accomplished using the data on constants of the stability of associated HM species and their solubility constants (Table 3).

Analysis of the presented data shows that the concentration of Cu^{2+} ions in equilibrium solutions during absorption by the calcaric fluvisols and calcaric fluvic arenosols lies on a straight line corresponding to the solubility of sediment $\text{Cu}(\text{OH})_2$, suggesting the possibility of the formation of a compound, in addition to its adsorption by hydromorphic soils. Data points of Zn^{2+} concentration are grouped slightly lower, relative to the more soluble ZnCO_3 . Obviously, carbonate can also form near $\text{pH} = 7.3$. In the field of higher and lower pH values, the formation of this sediment is less likely, because Zn concentration in the equilibrium solution is lower than the values needed for sedimentation.

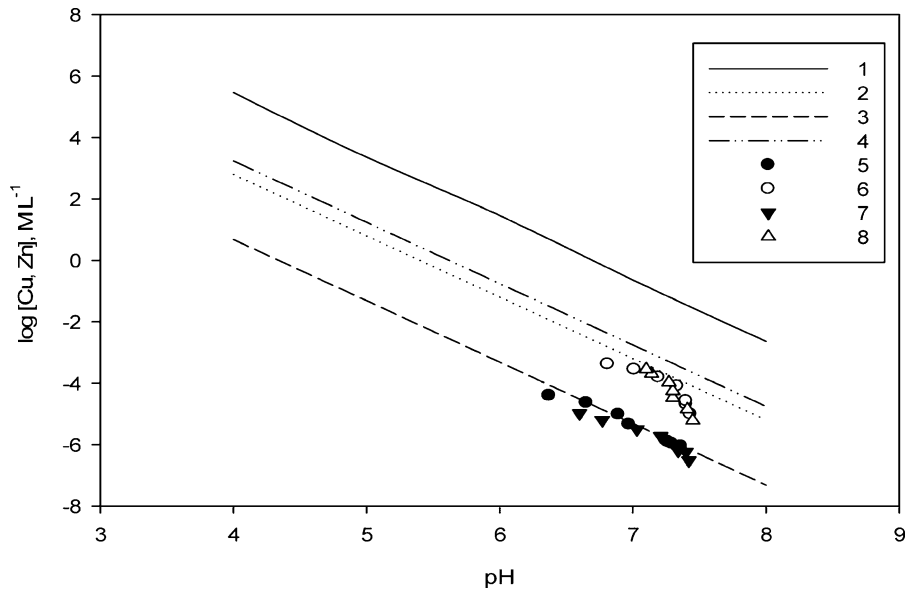


Fig. 4 Diagrams of the solubility of CuCO_3 (1), ZnCO_3 (2), Cu(OH)_2 (3), and Zn(OH)_2 (4) and logarithms of Cu and Zn concentrations in equilibrium solutions at adsorption by the calcaric fluvic arenosols (5, 6) and calcaric fluvisols (7, 8)

Table 3 The logarithms values of the constants of stability ($\log \beta$) and solubility ($\log K_{sp}$) of some associates and low-solubility Cu and Zn compounds (Lur'e 1979; Sillen and Martell 1970)

Associates	$\log \beta$	Low-solubility compounds	$\log K_{sp}$
ZnOH^+	6.31	Zn(OH)_2	-16.76
Zn(OH)_2^0	11.19	ZnCO_3	-10.84
ZnCO_3^0	6.63	Cu(OH)_2	-19.32
$\text{Zn(CO}_3)_2^{2-}$	9.63	CuCO_3	-8.18
ZnHCO_3^+	2.07		
ZnNO_3^+	0.40		
$\text{Zn(NO}_3)_2^0$	-0.30		
CuOH^+	6.20		
Cu(OH)_2^0	14.50		
$\text{Cu}_2(\text{OH})_2^{2+}$	17.50		
CuNO_3^+	0.50		
CuCO_3^0	6.73		

According to Sparks (2001), the HM sorption by soil and soil components produces a new phase composed of sediment Me(OH)_2 (low-solubility compound). In this case, the surficial precipitation is observed at ion concentrations and pH values much lower than those when the solution becomes saturated relative to the solid phase. A local pH heterogeneity

related to protonation of the surface can be observed near the surface of mineral or organomineral particles of soil. The pH values near the surface can be much higher than in the solution volume (Kurochkina et al. 2014). This phenomenon also promotes sedimentation of the low-soluble HM hydroxides and carbonates.

The comparison of diffraction patterns for the layered silicate phases of montmorillonite is performed before and after modification with a saturated $\text{Cu(NO}_3)_2$ solution. The results showed appreciable changes in the diffraction pattern of the contaminated phases of montmorillonite compared to the original samples (Fig. 5). Additional diffraction peaks corresponding to a new crystalline phase appeared in the diffraction patterns of montmorillonite saturated with Cu^{2+} ions.

A peculiar feature of layered silicate phases is the chemical and energetic heterogeneity of their surface characterized by the presence of structural defects and different functional groups, which can act as active centers during metal adsorption (Minkina et al. 2018). These active centers on the surface of montmorillonite can include exchangeable cations, surface hydroxyl groups, and oxygen atoms of the tetrahedral lattice. Some active centers occur on the lateral faces of minerals formed during the splitting of minerals.

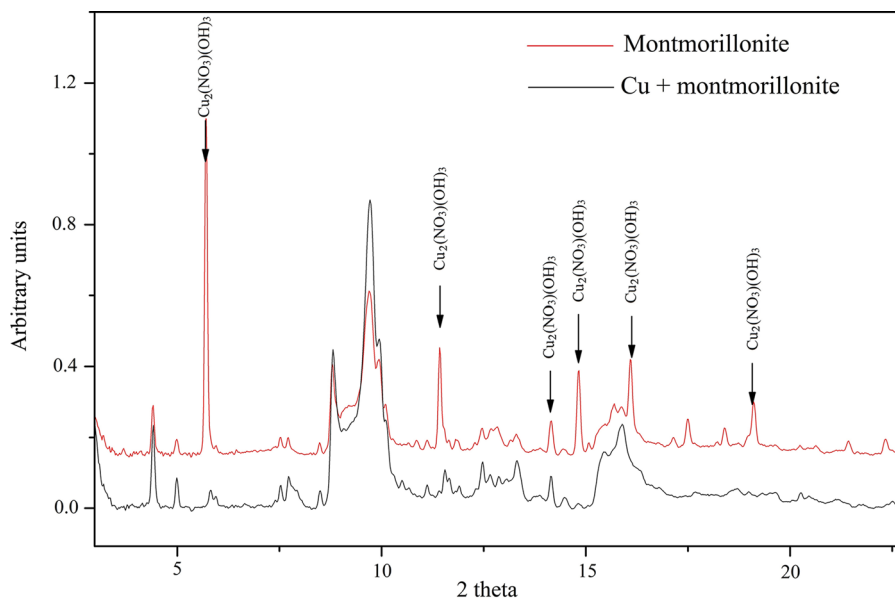


Fig. 5 Comparative analysis of X-ray powder diffractograms of the initial sample of montmorillonite mineral phase and after saturation with $\text{Cu}(\text{NO}_3)_2$

The saturation of phases of layered silicates with a $\text{Cu}(\text{NO}_3)_2$ solution at a constant pH of 3.9 increases the share of active acidic centers, which affects the proportions of acid–base active centers on the surface of mineral phases (Ponizovskii and Mironenko 2001). Hydrolysis processes, which shift the system equilibrium, also contribute. During the initial saturation period, Ca^{2+} cations are desorbed from the interlayer positions of layered silicate into the contacting solution.

X-ray diffraction data showed that Cu^{2+} ions are sorbed from the saturated solution by active centers on the internal surface of the lattice of dioctahedral aluminosilicates, and surface hydroxyls at the octahedrally coordinated aluminum atom play the main role (Furnare et al. 2005). X-ray diagnostics revealed that excess Cu^{2+} ions are removed from the system due to the formation and precipitation of coarsely crystalline $\text{Cu}_2(\text{NO}_3)(\text{OH})_3$ (Fig. 3).

Comparison of the XANES spectra of the initial Zn $(\text{NO}_3)_2$ compound and the saturated Zn phase of carbonates revealed a shift (more than 1 eV) by the position of the main energy maximum, which means a change in the charge state as a result of electron transfer from Zn. Analysis of the K-edge XANES spectra of Zn adsorbed by soil carbonates revealed that the metal ions substitute Ca at octahedral sites of

calcite (Fig. 6). This observation is followed by $1s \rightarrow 4p$ electron transition, and metal ions coordinate with carbonate ions as ligands to make up absorption complexes on the surface of the mineral calcite, first of all, at the defect and fracture sites. The extended X-ray absorption fine structure spectroscopy was used to confirm Zn incorporation into calcite. Zn occupies the Ca position in octahedral coordination, with Zn–O distances only slightly larger than in pure ZnCO_3 (Elzinga and Reeder 2002).

Conclusion

Regularities in the absorption of Zn and Cu by the main types of hydromorphic soils in southern Russia are studied. Using combination of physicochemical and direct physical methods provides detailed information about the nature of the soil components and possible interactions with HM. Isotherms of sorption by the calcareous fluvisols (loamic) demonstrate a high affinity of the soil particle surface to the studied metals, relative to calcareous fluvic arenosols. This is due to the higher content of organic matter, clay, and silt in the fluvisols, compared to the arenosols. Experiments with the sorption of Cu and Zn by the studied soils during their joint presence demonstrated that the soil

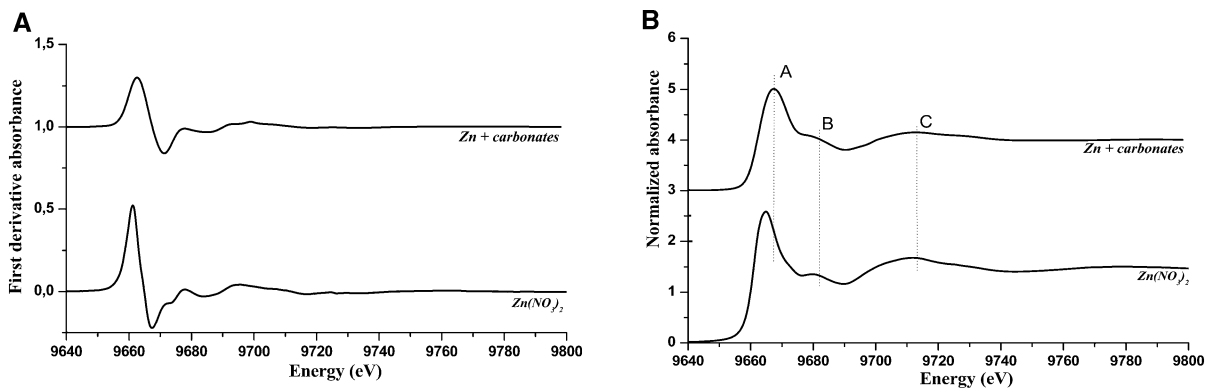


Fig. 6 Experimental Zn K-edge XANES spectra in samples of carbonates and reference compounds $Zn(NO_3)_2$ (a); first derivative of the Zn K-edge XANES spectra (b)

particle surface includes reaction centers with different selectivity relative to metals. Under conditions of competition, Cu manifests a higher affinity to adsorption centers than Zn. Analysis of the solubility diagrams of metal compounds revealed a possibility of the formation of sediments of the low-solubility Cu hydroxides and Zn carbonates in the studied soils. Analysis of the XANES spectra and XRD identification confirmed the results based on simulations of the solubility diagram of metal compounds. In addition to ion exchange processes, the system is also marked by the formation of a Zn carbonate adsorbed on the surface of soil carbonates (calcite), and Cu ions are precipitated on the surface of the main layered minerals as the coarse crystalline phase $Cu_2(NO_3)(-OH)_3$. The obtained results imply that montmorillonite and calcite can be used as an efficient material for the retention of Zn and Cu in soils.

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