Structural Study of the Incorporation of Heavy Metals into Solid Phase Formed during the Oxidation of EDTA by Permanganate at High pH

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Properties of solid phases formed during the oxidation of EDTA by permanganate in a high-pH, high-ionic strength solution, and the retention of Cu²⁺, Ni²⁺, and Zn²⁺ by them were examined. Morphologically, the solids were agglomerates of particles with sizes <100 nm. X-ray absorption spectroscopy (XAS) analysis indicated that these particles were birnessite. Its precipitation was accompanied by the removal of Zn²⁺ and Cu²⁺ released as a result of the breakdown of their complexes with EDTA. However, Ni²⁺ was not removed from the supernatant. Cu²⁺ was strongly bound by birnessite and exhibited little mobility in the pH range from 3 to 14. Zn²⁺ was more mobile, especially at pH > 12. XAS showed that Cu²⁺ binding sites were located within MnO₆ octahedra- comprised sheets that constitute birnessite while Zn²⁺ was positioned between them.

Introduction
Permanganate has been successfully used for in situ degradation of many organic contaminants (1, 2). At Hanford Nuclear Reservation in the state of Washington, it has been utilized to treat high pH, high ionic strength nuclear tank wastes that contain EDTA, NTA, and related ligands. These ligands are oxidized by alkaline permanganate to predominantly CO₂, ammonia, and oxalic acid (1, 3, 4). Following the oxidation, complexes of these ligands with metal cations break down to release uncomplexed metal ions that precipitate, coprecipitate with, or adsorb on solids formed as a result of the reduction of permanganate. This improves the separation of the target elements (transuranic elements, TRU, and Sr⁹⁰) that must be achieved during the treatment (3–6).

Solids formed by the reduction of permanganate at high pH play a crucial role in the removal of TRU and metals present in the wastes. Their nature has not been ascertained, although it has been observed that when the high-pH reduction of permanganate proceeds via the oxidation of H₂O₂ and formate, the reaction yields the mixed Mn(IV) oxide birnessite that belongs to the phyllomanganate group (7). The paucity of information about manganese solids formed in situ in the important case of the permanganate treatment of EDTA-containing nuclear wastes is in contrast with extensive studies of Mn oxides formed ex situ (8–15). These studies have shown that in birnessite and other phyllosilicates the MnO₆ units are arranged in two-dimensional sheets (11–15), whose local structure can be altered by metal cations (Cu²⁺, Ni²⁺, and Zn²⁺) incorporated into them (11, 13).

One practically important question is whether the in situ formed manganese solids bind TRU and other metal cations structurally and thus make them mobilization-resistant as long as the solid substrate remains structurally intact, or are the retained metals sorbed on the surface and can be mobilized without affecting the substrate, for instance, by acidic solutions or in the presence of complexing agents.

To address the above issues, we carried out experiments designed to ascertain the structural properties of solids precipitated during the permanganate oxidation of simulated wastes similar to those in Hanford tanks. EDTA was used as a representative chelating agent that was oxidized by MnO₄⁻. Cu²⁺, Zn²⁺, and Ni²⁺ were chosen as target metals with which to quantify the incorporation of cations present in the wastes into the solid substrate. The structural data were obtained primarily using X-ray absorption spectroscopy (XAS).

Experimental Section
All experiments were conducted at 20 °C using 50 mL polyethylene centrifuge tubes. The concentration of NaOH in the samples was 1 M (pH 14). The initial concentration of EDTA in the solutions was 0.02 M, while that of the metal ions (Cu²⁺, Zn²⁺, and Ni²⁺) was 0.002 M. All chemicals were ACS reagent grade. Reagent water was obtained with a Millipore Super-Q water system. The amount of permanganate added to the solutions corresponded to MnO₄⁻/EDTA molar ratios of 1 and 4. Reactions between EDTA and permanganate caused manganese solids to form immediately, but the reaction was allowed to proceed for 40 h, long after the permanganate was completely consumed. The test tubes were mildly agitated during that time. The precipitated solids and supernatant were separated by centrifugation. Aliquots of the supernatant were collected to determine the concentration of target elements (Mn, Ni, Cu, and Zn) in them.

The separated solids were split into three subsamples that were washed at three constant pH values (3, 7, and 14, respectively) using two cycles of resuspension in the washing solutions (Milli-Q water with a desired pH) followed by centrifugation. The washed solids were retained on 0.45 μm filters, dried in a desiccator, and used in XAS measurements. Experiments were also carried out with the solids that were first resuspended at pH 14, and then the pH of the system was decreased gradually (with pH increments of 0.3–0.5 units) by adding HClO₄. The supernatant was analyzed for Mn, Cu, Zn, and Ni to determine the influence of pH on the mobilization of the retained metal cations.

Concentrations of metals were measured with a Jobin-Yvon JY-138 Ultratrace inductively coupled plasma (ICP) atomic emission spectrometer. Prior to ICP analysis, all samples were filtered through a 0.45 μm filter and acidified with HNO₃. For structural studies, dried solids were coated with carbon and their morphology was examined with a JEOL JSM-7000F scanning electron microscope (SEM). Brunauer-Emmet-Teller (BET) specific surface areas were determined using a Micromeritics Flow Sorb II 2300 instrument. A Philips PW 1820 X-ray diffractometer was employed to examine the...
structur, having values of 10.3 and 13.5 m²/g for reaction contact with the solids formed during the oxidation of EDTA.

**Results and Discussion**

Measurements of Mn concentrations in the supernatant in contact with the solids formed during the oxidation of EDTA by MnO₄⁻ showed that all the manganese introduced as permanganate was incorporated into them. The surface area of the solids changed only slightly as the reduction of MnO₄⁻ progressed, having values of 10.3 and 13.5 m²/g for reaction times 0.5 and 24 h (Figure S1, Supporting Information). SEM examination showed that the solids lacked specific morphological features and were aggregates with apparent sizes <100 nm. The presence of particles with sizes <10 nm was also notable (Figure S2, Supporting Information).

Fourier transform magnitudes of EXAFS (also known as EXAFS radial distribution functions, ERDFs) were calculated using the K-edge Mn EXAFS for the solids formed in situ at pH 14 (Figure 1). The peak positions in the ERDFs are not corrected for photoelectron phase shifts and, therefore, are shifted by ca. 0.5 Å toward the absorber relative to the actual interatomic distances. EXAFS data for the samples washed at pH 7.0 and 3.0 were indistinguishable from those represented in Figure 1. The first intense peak in the ERDF spectra is located at 1.40 Å. It corresponds to six oxygen atoms that comprise the inner complexation shell of the manganese. The second peak located at 2.41 Å corresponds to the second neighboring manganese atoms, and the small peak at 3.05 Å is likely to be associated with corner-sharing MnO₄ octahedra in birnessite (22, 25).

Mn–O and Mn–Mn distances that were determined via fitting of the experimental K-edge Mn EXAFS data are compiled in Table 1. In agreement with prior research, the coordination numbers for Mn–O and Mn–Mn bonds in these calculations were fixed as 6 (13–15, 20–22). The Mn–O (1.90 ± 0.01 Å) and Mn–Mn distances (2.65 ± 0.01 Å) in both samples were identical. The σ² values for the Mn–O and Mn–Mn distances were small (for instance, 0.0057 and 0.0091 Å², respectively, for particles generated at a 4.0 MnO₄⁻/EDTA ratio). This indicates that, despite the lack of the long-term order in the solids, they are highly ordered microstructurally. The Mn–Mn distance of 2.85 Å determined for our samples corresponds to the edge-sharing hexagonal octahedron

![FIGURE 1. Mn radial density functions and fitting results for manganese solids formed at pH 14 and MnO₄⁻/EDTA molar ratios 1 (A) and 4 (B).](image)
structure (7). This arrangement is characteristic for phyllo-
manganese minerals such as δ-MnO₂ and birnessite (25–
27). Based on the sum of the data of SEM, XRD, XANES, and
EXAFS techniques, as well as results of prior research (7), we
concluded that particles precipitated upon the oxidation of
EDTA by permanganate are birnessite-type solids, although
the presence of minor amounts of other Mn(IV) phases could
not be ruled out completely.

Examination of the high-pH oxidation of EDTA by
preformed MnO₂ showed that EDTA was stable in these
conditions. It was accordingly concluded that reduction of
Mn(IV) to Mn(III) by EDTA did not exceed the level defined
by the precision of the experiments (5%). This indicates that
the high-pH permanganate oxidation of EDTA is likely to
result in the formation of low-sodium birnessite in which the
charge deficit in the MnO₆ octahedra-comprised sheets is
associated primarily with defects in the two-dimensional
layers and lower charge cations such as Cu²⁺ and Zn²⁺ that
are incorporated into them in statu nascendi, as described in
the section that follows. The conclusion that low-sodium

birnessite is formed upon the reduction of MnO₄⁻ by EDTA
is different from that obtained for the reduction of MnO₄⁻
by stronger reduction agents such as H₂O₂ and formate (7).
In that case, the precipitated solids were ascribed the
stoichiometry of the mineral clinobirnessite Na₄Mn₁₄O₂₇·
9H₂O. Despite possible differences in the stoichiometries of
the solid phases formed via the high-pH reduction of
permanganate by EDTA or H₂O₂/formate, in either case the
solids have phyllomanganate-type structures (19).

Precipitation of birnessite particles was accompanied by
effective removal of Cu²⁺ and Zn²⁺ that were present in the
original solution as EDTA complexes (28, 29) and possibly lack of sorption of
Ni²⁺-EDTA complexes on birnessite. The resistance of the
Ni²⁺-EDTA complex during the high-pH permanganate

<table>
<thead>
<tr>
<th>TABLE 1. Structural Characteristics of Manganese in Solids Obtained by Oxidation of EDTA by Permanganate at pH 14*</th>
</tr>
</thead>
<tbody>
<tr>
<td>R (Å)</td>
</tr>
<tr>
<td>-------</td>
</tr>
<tr>
<td>sample R1 (MnO₄⁻/EDTA molar ratio 1)</td>
</tr>
<tr>
<td>sample R4 (MnO₄⁻/EDTA molar ratio 4)</td>
</tr>
</tbody>
</table>

* Data of Mn K-edge EXAFS (CN for Mn–O = 6, Mn–Mn = 6).
treatment is not necessarily a result of a uniquely high thermodynamic stability of Ni$^{2+}$–EDTA complexes. In fact, the log $\beta$ values for Cu$^{2+}$–EDTA and Ni$^{2+}$–EDTA complexes are close, 20.5 and 20.1, respectively (28, 29), while the behavior of these metals is different. It can be hypothesized that the permanganate oxidation of Ni$^{2+}$–EDTA targets not only EDTA but also Ni$^{2+}$ to produce Ni$^{3+}$ that forms highly stable complexes with EDTA (28).

Further exploration of this issue goes beyond the scope of this paper.

Washing at pH 7 and 3 caused a small fraction of copper bound by the solids formed using a 1.0 MnO$_4$–/EDTA molar ratio to be leached out (Figure 2). For solids formed using a 4.0 MnO$_4$–/EDTA molar ratio, the amount of copper bound in the solid phase was as stable as it was at pH 14. Results for zinc showed some variability, but increasing the MnO$_4$–/EDTA molar ratio helped increase both the retention of this metal at pH 14 and its stability at lower pH. Despite the low efficiency of nickel removal during the permanganate oxidation, the amount of nickel that was bound by birnessite was stable during washing at pH 3 and 7 (Figure 2).

Experiments in which the pH of the suspensions was decreased from 14 to 3 showed that virtually no manganese release took place, but the retained Zn$^{2+}$ and Cu$^{2+}$ cations were mobilized to some extent, with systematic differences between the behavior of copper and zinc. The copper retained by the particles formed using both 1 and 4 MnO$_4$–/EDTA molar ratios was stable in the entire range of pH values. Only about 0.2% of it was released into the supernatant at pH > 10 (Figure 3A). At pH < 6, Cu release increased as well but even at pH 3 the percentage of copper mobilized from the solid phase did not exceed 0.2%.

Release of Zn was more prominent. For solids formed using a 1.0 MnO$_4$–/EDTA molar ratio, up to 25% of the zinc initially retained by birnessite was released at pH > 12 (Figure 3B). The mobility of zinc also slightly increased at pH < 4. A similar pattern was observed for solids formed using a 4.0 MnO$_4$–/EDTA molar ratio, but the amount of zinc in contact with it decreased more than 3-fold.

To probe mechanisms that govern the retention of Zn$^{2+}$ and Cu$^{2+}$ by in situ formed birnessite, EXAFS measurements for Cu and Zn were performed. The concentrations of Zn and Cu in the samples are shown in Figure S4 in the Supporting Information. EXAFS measurements for Ni were not carried out.

The analysis of the Zn EXAFS data was initially done assuming that the immobilized Zn$^{2+}$ was bound at the defect sites located in the layers formed by MnO$_6$ octahedra. However, it was observed that the Zn–Mn distance that was
TABLE 2. Structural Characteristics of Zinc in Solids Formed via the Oxidation of EDTA by Permanganate at pH 14a

<table>
<thead>
<tr>
<th>sample</th>
<th>Zn–O</th>
<th>Zn–Mn</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>R (Å)</td>
<td>σ2 (Å²)</td>
</tr>
<tr>
<td>sample R1, pH 14</td>
<td>1.93 ± 0.02</td>
<td>0.010 ± 0.002</td>
</tr>
<tr>
<td>sample R1, pH 7</td>
<td>1.97 ± 0.01</td>
<td>0.011 ± 0.001</td>
</tr>
<tr>
<td>sample R1, pH 3</td>
<td>1.91 ± 0.02</td>
<td>0.009 ± 0.002</td>
</tr>
<tr>
<td>sample R4, pH 14</td>
<td>1.94 ± 0.02</td>
<td>0.010 ± 0.002</td>
</tr>
<tr>
<td>sample R4, pH 7</td>
<td>2.01 ± 0.02</td>
<td>0.012 ± 0.002</td>
</tr>
<tr>
<td>sample R4, pH 3</td>
<td>1.99 ± 0.01</td>
<td>0.011 ± 0.002</td>
</tr>
</tbody>
</table>

a Data of Zn K-edge EXAFS (CN for Zn–O = 6, Zn–Mn = 6).

TABLE 3. Structural Characteristics of Copper in Solids Formed via the Oxidation of EDTA by Permanganate at pH 14a

<table>
<thead>
<tr>
<th>sample</th>
<th>equatorial Cu–O (4)</th>
<th>axial Cu–O (2)</th>
<th>Cu–Mn (6)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>R (Å)</td>
<td>σ2 (Å²)</td>
<td>R (Å)</td>
</tr>
<tr>
<td>sample R1, pH 14</td>
<td>1.95 ± 0.08</td>
<td>0.007 ± 0.008</td>
<td>1.97 ± 0.08</td>
</tr>
<tr>
<td>sample R1, pH 7</td>
<td>1.93 ± 0.06</td>
<td>0.005 ± 0.008</td>
<td>1.98 ± 0.08</td>
</tr>
<tr>
<td>sample R1, pH 3</td>
<td>1.91 ± 0.03</td>
<td>0.002 ± 0.009</td>
<td>2.00 ± 0.09</td>
</tr>
<tr>
<td>sample R4, pH 14</td>
<td>1.93 ± 0.04</td>
<td>0.004 ± 0.004</td>
<td>2.01 ± 0.08</td>
</tr>
<tr>
<td>sample R4, pH 7</td>
<td>1.89 ± 0.05</td>
<td>0.001 ± 0.011</td>
<td>2.02 ± 0.13</td>
</tr>
<tr>
<td>sample R4, pH 3</td>
<td>2.00 ± 0.03</td>
<td>0.004 ± 0.004</td>
<td>1.99 ± 0.13</td>
</tr>
</tbody>
</table>

a Data of Cu K-edge EXAFS (CN for Cu–O = 6, including 4 equatorial and 2 axial bonds; Cu–Mn = 6).

Assumed to exist for this structure was inconsistent with the observations. Because of that, the local structure around the Zn2+ ions was hypothesized to be similar to that of chalcopyrite ZnMn3O7·3H2O. The structure of chalcopyrite is formed by sheets of layered MnO3 octahedra with one vacant site per every seven cation sites. The Zn2+ ions are coordinated in chalcopyrite above the vacant sites located in the sheets of MnO6 octahedra (12, 30).

Processing of the Zn EXAFS data using this hypothesis resulted in an acceptable level of goodness-of-fit. The results for pH 3, 7, and 14 for a 4.0 MnO4-/EDTA ratio are shown in Figure 4. Additional Zn EXAFS data for solids formed at a 1.0 MnO4-/EDTA ratio are represented in Figure S6 in the Supporting Information.

The information obtained based on Zn EXAFS data is summarized in Table 2. The data show that the average Zn–O and Zn–Mn distances in all samples are close, with two trends related to the effects of pH and MnO4-/EDTA dose. First, the Zn–O and Zn–Mn distances at pH 7 (2.01 and 3.44 Å respectively for a 4.0 MnO4-/EDTA ratio) are somewhat longer than those for pH 3 and 14. Second, these distances are slightly longer in the samples obtained using a 4.0 MnO4-/EDTA ratio. The values of the Debye–Waller factor (σ2) indicating the disorder of the Zn–O bonds are similar for all conditions, while those for the Zn–Mn distances are somewhat lower at pH 14.

Despite these changes, the local structure around the Zn2+ ions immobilized in the birnessite particles remains similar to that of chalcopyrite in all cases. It is likely that during the in situ formation of the birnessite, their structure undergoes a change induced by interactions with the decomplexed Zn2+ to produce, locally, chalcopyrite-type structures for sites at which Zn2+ is incorporated. This is in agreement with the data of prior studies of the incorporation of Zn2+ into preformed birnessite (13, 15).

Attempts to fit the Cu EXAFS data assuming a structural arrangement similar to that of chalcopyrite were unsuccessful. Accordingly, it was hypothesized that Cu2+ was bound at sites located within the two-dimensional sheets comprised by MnO6 octahedra. It was also assumed that the number of nearest neighbors in the inner shell of the bound Cu2+ ions was six, including four equatorially and two axially positioned oxygen atoms with potentially different Cu–O distances (Jahn–Teller effect) (15, 22, 31–33). The number of manganese atoms that affect the EXAFS spectra of incorporated Cu2+ ions was six.

This hypothesis allowed obtaining a good quality of fitting for the Cu EXAFS data. Figure 5 compares the results of the fitting that utilized these assumptions and experimental Cu EXAFS data for the sample generated using a 4.0 MnO4-/EDTA ratio. Additional Cu EXAFS data for solids formed at a 1.0 MnO4-/EDTA ratio are represented in Figure S6 in the Supporting Information. The structural data for Cu2+ bound in birnessite are summarized in Table 3. They indicate that neither the pH of the washing solution nor MnO4-/EDTA molar ratio at which these solids were formed affects the structural status of the bound copper. The equatorial Cu–O distances (ca. 1.89–1.95 Å, except one sample washed at pH 3, Table 3) determined for all samples are close to those determined for equatorially bound oxygen atoms in aqueous Cu2+ complexes and many sorbed Cu2+ species (15, 22, 31–33). Both the axial Cu–O distances for Cu2+ incorporated into birnessite (1.97–2.02 Å) and their respective Debye–Waller factors are close to those for the equatorial bonds. This indicates a nearly complete elimination of the Jahn–Teller effect. This is indicative of strong interactions between the incorporated Cu2+ ions and MnO6 octahedra that surround them.

The Debye–Waller factors for the Cu–Mn distances are higher than those for the Cu–O bonds, and the scatter of the Cu–Mn distances exhibit an increase when the pH is decreased from 14 to 3 (Table 3). This is likely to indicate that although Cu2+ remains tightly bound in the birnessite at pH values as low as 3, the strength of bonds between this metal and the substrate becomes weaker.

The Cu–Mn distances in all samples are similar (2.80–2.89 Å). This value is close to that for the Mn–Mn distance (2.85 Å) in unsubstituted birnessite (Table 1). This provides a strong indication that that little distortion of the layers formed by MnO6 octahedra takes place when Cu2+ is incorporated into them.

The above structural data indicate that the birnessite-type solids formed during the oxidation of high-pH, high-ionic nuclear wastes containing EDTA and similar organic ligands are likely to resist mobilization of heavy metals such as Cu and Zn incorporated into them in a wide range of
environmental conditions, as long as the reduction of Mn-(IV) that precipitates in the solid is prevented. The issue of whether the representative TRU is bound in a manner similar to that established in this study for Zn²⁺ and Cu²⁺ needs further exploration.

Acknowledgments

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Supporting Information Available

Six figures demonstrating changes of the specific surface area of birnessite solids precipitated during the reduction of permanganate by EDTA at varying reaction times, the morphology of these solids and their Mn XANES spectra, and concentrations of and EXAFS radial density functions for Cu and Zn retained by birnessite formed using a low permanganate/EDTA molar ratio. This material is available free of charge via the Internet at http://pubs.acs.org.

Literature Cited


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