

Removal of CrO_4^{2-} , a Nonradioactive Surrogate of $^{99}\text{TcO}_4^-$, Using LDH– Mo_3S_{13} Nanosheets

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Cite This: *Environ. Sci. Technol.* 2022, 56, 8590–8598



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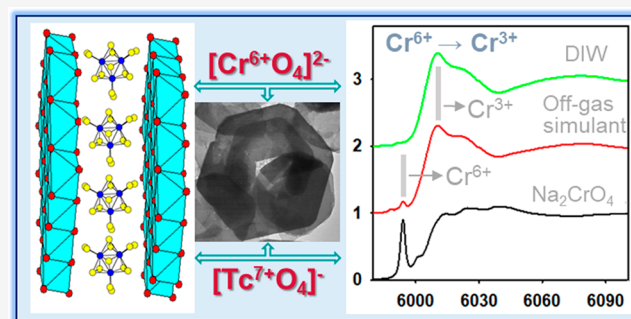
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Supporting Information

ABSTRACT: Removal of chromate (CrO_4^{2-}) and pertechnetate (TcO_4^-) from the Hanford Low Activity Waste (LAW) is beneficial as it impacts the cost, life cycle, operational complexity of the Waste Treatment and Immobilization Plant (WTP), and integrity of vitrified glass for nuclear waste disposal. Here, we report the application of $[\text{Mo}^{\text{IV}}_3\text{S}_{13}]^{2-}$ intercalated layer double hydroxides (LDH– Mo_3S_{13}) for the removal of CrO_4^{2-} as a surrogate for TcO_4^- , from ppm to ppb levels from water and a simulated LAW off-gas condensate of Hanford's WTP. LDH– Mo_3S_{13} removes CrO_4^{2-} from the LAW condensate stream, having a pH of 7.5, from ppm ($\sim 9.086 \times 10^4$ ppb of Cr^{6+}) to below 1 ppb levels with distribution constant (K_d) values of up to $\sim 10^7$ mL/g. Analysis of postadsorbed solids indicates that CrO_4^{2-} removal mainly proceeds by reduction of Cr^{6+} to Cr^{3+} . This study sets the first example of a metal sulfide intercalated LDH for the removal of CrO_4^{2-} , as relevant to TcO_4^- , from the simulated off-gas condensate streams of Hanford's LAW melter which contains highly concentrated competitive anions, namely F^- , Cl^- , CO_3^{2-} , NO_3^- , BO_3^{3-} , NO_2^- , SO_4^{2-} , and $\text{B}_4\text{O}_7^{2-}$. LDH– Mo_3S_{13} 's remarkable removal efficiency makes it a promising sorbent to remediate $\text{CrO}_4^{2-}/\text{TcO}_4^-$ from surface water and an off-gas condensate of nuclear waste.

KEYWORDS: nuclear waste, pertechnetate, chromate, low activity waste, off-gas condensate stream, Hanford's radioactive waste



INTRODUCTION

The Hanford Waste Treatment and Immobilization Plant (WTP), the DOE's largest nuclear waste treatment plant, is designed to process, treat, and immobilize much of the radioactive legacy wastes.^{1–4} The current plan of the Hanford Waste Treatment and Immobilization Plant (WTP) is to separate the tank wastes into high-level waste (HLW) and low-activity waste (LAW) and permanently immobilize the radioisotopes into separately vitrified waste forms.⁵ LAW constitutes a larger fraction of the waste by volume, and it consists of high ionic-strength solutions that contain Na^+ , K^+ , $\text{Al}(\text{OH})_4^-$, Cl^- , F^- , NO_2^- , NO_3^- , OH^- , CO_3^{2-} , organics, other minor ions of Cr, Ni, Cd, and Pb, and radionuclides.⁶ Technetium-99 (^{99}Tc) is a β -emitting long living, $t_{1/2} \sim 2.13 \times 10^5$ years, radionuclide which mostly remains in LAW as TcO_4^- .^{4,7–10} Immobilization of TcO_4^- at the WTP becomes challenging because of its high solubility, complex redox chemistry, and volatility at the vitrification temperature.^{4,7–10} The treatment and vitrification of LAW at the Hanford WTP will generate an aqueous LAW off-gas condensate which is planned to be repeatedly reprocessed and passed through the glass melter until nearly complete immobilization of technetium.^{4,8,11} This procedure results in the increase of the

WTP operational time, waste volume, and vitrification cost. Importantly, this process could be limited for extremely low concentrations, namely ≤ 5 ppb. Also, the off-gas condensate contains halides and chromates which have a detrimental effect on the glass formulation during the vitrification.^{12–14} Therefore, it is important to develop materials, methods, and technologies to redirect the technetium immobilization from LAW.

With a reduction potential of -0.22 V (for the $\text{S}^{2-}/\text{SO}_4^{2-}$ couple), sulfides are capable of reducing the highly soluble TcO_4^- ($\text{TcO}_4^-/\text{Tc}^{4+}$ is $E^\circ \sim +0.74$ V) and CrO_4^{2-} ($\text{CrO}_4^{2-}/\text{Cr}^{3+}$ is $E^\circ \sim +1.23$ V) ions to insoluble Tc(IV) and Cr(III) species.^{6,15–21} Because of the known chemistry of sulfides and other reductants for the reductive precipitation of highly redox-active Tc^{7+} and Cr^{6+} , we introduced $(\text{Cr}^{6+}\text{O}_4)^{2-}$ as a nonradioactive surrogate for a reductive precipitation driven

Received: January 15, 2022

Revised: May 13, 2022

Accepted: May 16, 2022

Published: June 1, 2022



separation.^{1,9,22,23} In addition to being a suitable surrogate for TcO_4^- , CrO_4^{2-} itself is highly toxic²⁴ and also persists within the radioactive waste of the Hanford nuclear waste streams.^{25–27} Because of the leakage in the Hanford underground tanks, chromate has been dispersing in the environment along with various radionuclides.^{28–30} Moreover, during the vitrification process of the LAW, CrO_4^{2-} is transformed into crystalline spinel which in turn jeopardizes the integrity of the glass waste form.^{13,31} Therefore, in addition to pertechnetate, it is crucial to remediate chromate from water and legacy nuclear wastes.

Numerous metal sulfides, such as pyrrhotite (Fe_{1-x}S) and pyrite (FeS_2),^{32,33} chalcocite (Cu_2S),^{29,30} and stibnite (Sb_2S_3),³⁴ and layered metal sulfides⁶ are known to immobilize TcO_4^- by reductive precipitation.^{15,16} Layered double hydroxides (LDHs) are anionic clays which can also remove $\text{TcO}_4^-/\text{CrO}_4^{2-}$ but with ion-exchange and surface sorption mechanisms.^{8,15,24,35} Considering the efficiency of LDH and metal sulfides for $\text{TcO}_4^-/\text{CrO}_4^{2-}$ remediation, we hypothesized that a hybrid structure of metal sulfide intercalated LDH will boost the removal of pertechnetate from water and legacy nuclear waste. Altogether, it becomes important to investigate LDH-metal sulfide based materials, evaluate the remediation of TcO_4^- , and delineate the sorption mechanisms involved in this process.

Herein, we report a modified synthesis method for LDH- Mo_3S_{13} ³⁶ to achieve nanosheets and the investigation of CrO_4^{2-} , which also acts as a TcO_4^- surrogate, sorption from naturally contaminated surface water, and a simulated off-gas condensate of the LAW melter. We show that LDH- Mo_3S_{13} exhibits ultrahigh efficient sorption of CrO_4^{2-} from water and an off-gas condensate from ppm to ppb levels (<1 ppb) with K_d values of up to $\sim 10^7$ mL/g. For LDH- Mo_3S_{13} , we show that reductive precipitation is the dominant mechanism; however, ion-exchange and/or surface sorption can provide parallel paths for the sequestration of CrO_4^{2-} . These integrated multimode sorption processes make LDH- Mo_3S_{13} a promising sorbent of $\text{CrO}_4^{2-}/\text{TcO}_4^-$ from polluted water and nuclear waste.

■ EXPERIMENTAL SECTION

Material Synthesis and Chromate Uptake Study.

$\text{MgAl}-\text{CO}_3$ (LDH- CO_3), $\text{MgAl}-\text{NO}_3$ (LDH- NO_3), and $(\text{NH}_4)_2\text{Mo}_3\text{S}_{13}\cdot\text{H}_2\text{O}$ were synthesized as previously described.^{37–40} LDH- Mo_3S_{13} nanosheets were synthesized with a two-step method: first, exfoliation of LDH nanosheets of the LDH- NO_3 and then the settlement of positively charged LDH sheets in the presence of dissolved $\text{Mo}_3\text{S}_{13}^{2-}$ anions. More specifically, 200 mg of LDH- NO_3 was added to 20 mL of formamide and stirred for 24 h to exfoliate the positively charged LDH nanosheets. Afterward, a solution of 200 mg of $(\text{NH}_4)_2\text{Mo}_3\text{S}_{13}\cdot\text{H}_2\text{O}$ in 2 mL of formamide was slowly added to the exfoliated LDH solution and stirred for 30 min. Later, the solution was left at ambient conditions for 2 h, filtered, then washed with water and acetone, and subsequently dried at ambient conditions to give a brown solid of LDH- Mo_3S_{13} . Thus, this process of synthesis differs from the previously reported synthesis of LDH- Mo_3S_{13} .³⁶

Chromate Uptake Study. LDH- Mo_3S_{13} was added to the CrO_4^{2-} spiked solutions at different concentrations, stirred for different periods of time, and centrifuged to separate the supernatant solution from the solids. For the Hanford's LAW off-gas condensate simulated streams,¹ the sorption study was

conducted by the batch method by spiking the simulant with 9.086×10^4 ppb of Cr^{6+} as CrO_4^{2-} (CrO_4^{2-} equivalent concentrations $\sim 2.027 \times 10^5$ ppb), using variable loading of adsorbent, LDH- Mo_3S_{13} (10 to 100 mg), and at variable time scales (~ 1 h to 7 d), in 10 mL of simulated solutions, as described in Table S1.^{1,9} This study mostly focuses on the application of the LDH material for the remediation of $\text{CrO}_4^{2-}/\text{TcO}_4^-$ from the high ionic strength simulated off-gas condensate streams of Hanford LAW waste at pH ~ 7.5 . Hence, the variation of the solution pH and ionic strengths was not monitored or controlled during the Cr uptake studies since these are not known for the real off-gas condensate streams as the WPT has not started this operation yet.

After the sorption experiments, the residual CrO_4^{2-} concentrations in the supernatant were analyzed by inductively coupled plasma-mass spectrometry (ICP-MS). The adsorption capacity was determined from the difference in the concentrations of Cr before and after sorption.

The distribution coefficient (K_d) for the sorption of CrO_4^{2-} was determined in accordance with the equations, $K_d = (V[(C_0 - C_f)/C_f])/m$, where V is the solution volume (mL), C_0 and C_f are the initial and the final concentrations of CrO_4^{2-} in ppm, and m is the mass of the solid sorbent (g).⁴¹ In this work, we used the K_d values to compare the Cr^{6+} removal performance with other adsorbents. The removal rate of CrO_4^{2-} was computed using the equation of $100 \times (C_0 - C_f)/C_0$. The removal capacity, q_m (mg/g), can be obtained from the following equation: $10^{-3} \times (C_0 - C_f)V/m$. The adsorption experiments were carried out with $V:m$ ratios of 100–1000 mL/g, at room temperature (RT) and at different time scales ranging from minutes to days.

The sorption kinetics of LDH- Mo_3S_{13} was studied to determine the rate of removal of CrO_4^{2-} and to understand the sorption mechanism. In general, the adsorption rate is determined by two different rate equations, known as pseudo-first-order and pseudo-second-order mechanisms. Here, we used these mechanisms to analyze the adsorption phenomena of LDH- Mo_3S_{13} . The comparison was then drawn between the experimental and calculated data in accordance with the rate equations, as follows⁴²

Pseudo-first-order:

$$\ln(q_e - q_t) = \ln q_e - k_1 t \quad (1)$$

Pseudo-second-order:

$$\frac{t}{q_t} = \frac{1}{k_2 q_e^2} + \frac{t}{q_e} \quad (2)$$

where q_e (mg/g) is the amount of adsorbed element per unit mass of adsorbent at equilibrium, and q_t (mg/g) is the adsorbed amount at time t , while k_1 (min^{-1}) and k_2 ($\text{g}/\text{mg} \cdot \text{min}^{-1}$) are equilibrium rate constants of pseudo-first-order and pseudo-second-order adsorption interactions, respectively.⁴³ The k_1 value can be obtained by plotting $\ln(q_e - q_t)$ against t , and the k_2 value can be obtained by plotting t/q_t against t .

To understand the adsorbent and adsorbate interactions, we demonstrate the experimentally obtained adsorption data using the Langmuir isotherm model as given in eq 3. According to this model, the adsorbate moieties undergo monolayer type coverage on the surface of the adsorbent materials. It also predicts, once an adsorption site is occupied, no further adsorption can occur at the same site.⁴⁴ The Langmuir isotherm model for heterogeneous models is shown as eq 3

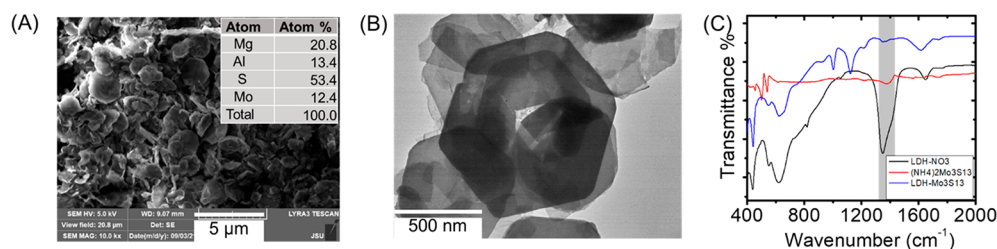


Figure 1. SEM image of the as-prepared LDH-Mo₃S₁₃ (Inserts show the average atomic abundance in percentage that was obtained by EDS.) (A), TEM image showing the ultrathin platelike morphology of the crystallites (B), and infrared spectrum showing the absence of NO₃⁻ peaks from the LDH-Mo₃S₁₃ indicating ion exchange of nitrate by the Mo₃S₁₃ (C).

$$\text{Langmuir isotherm: } q = q_m \frac{bC_e}{1 + bC_e} \quad (3)$$

where C_e (mg/L) is the concentration at equilibrium, q (mg/g) is the equilibrium sorption capacity of the adsorbed CrO₄²⁻, q_m (mg/g) is the theoretical maximum sorption capacity, and b (L·mg⁻¹) is the Langmuir constant which is related to the interaction energy of LDH-Mo₃S₁₃ and CrO₄²⁻.

Characterization. Samples were analyzed by scanning electron microscopy (SEM), transmission electron microscopy (TEM), Energy Dispersive Spectroscopy (EDS), X-ray Powder Diffraction (XRD) and Infrared (FT-IR) Spectroscopy, Inductively Coupled Plasma-Mass Spectrometry (ICP-MS), and X-ray photoelectron spectroscopy (XPS) (see the Supporting Information for experimental details).

Synchrotron X-ray Absorption Spectroscopy. The solid samples after the chromate interactions in DIW and simulated solutions were dried at ambient conditions and analyzed by synchrotron X-ray absorption near edge structure (XANES) and extended X-ray absorption fine structure (EXAFS). S K-edge and Mo L₃-edge XANES and Cr K-edge EXAFS measurements were performed at the Soft X-ray Microcharacterization Beamline (SXRMB) of the Canadian Light Source (CLS), Saskatoon, Canada. SXRMB is a bending-magnet-based beamline that utilizes InSb(111) and Si(111) crystals for monochromatization to cover an energy range of 1.7–10 keV. Samples were mounted onto double-sided, conductive carbon tape and loaded into the vacuum chamber with a vacuum of 10⁻⁷ Torr. Na₂CrO₄ and HgSO₄ were used as references and for energy calibration. A 7-element SDD detector was used to record the fluorescence yield (FY) of the powder samples. The total electron yield (TEY) by recording the drain current of the sample was also recorded. The collected data were processed and analyzed using the Demeter software package including Athena and Artemis.⁴⁵ Data from multiple scans were processed using Athena by aligning and merging the spectra followed by background subtraction using the AUTOBK algorithm. Chromium K-edge EXAFS data analysis was conducted on the merged and normalized spectra using Artemis. Theoretical models were constructed with the program FEFF7. Cr₂O₃ was used as a reference structural model.⁴⁶ Fits to the Cr EXAFS data were made in R space (R from 1 to 3.2 Å) and obtained by taking the Fourier Transform (FT) of $\chi(k)$ (k from 1.5 to 10.5) with a k weighting of 3.

RESULTS AND DISCUSSION

Synthesis and Characterization of LDH-Mo₃S₁₃ Nanosheets. The LDH-Mo₃S₁₃ hybrid nanosheets were synthesized using a novel two-step method starting from LDH-NO₃ that involves (i) the chemical exfoliation of LDH-

NO₃ and (ii) the subsequent treatment of the exfoliated ultrathin LDH nanosheets by the [Mo₃S₁₃]²⁻ anion (Figure 1). This modified synthesis is advantageous over the previously reported method³⁶ because it reduces ~90% of the organic solvents and the time to produce LDH-Mo₃S₁₃. A shorter interaction time in the solution can help to produce LDH nanosheets, retain the integrity of [Mo₃S₁₃]²⁻ anions, and avoid the side reactions. SEM and TEM images reveal the platelike morphology of LDH-Mo₃S₁₃ nanosheets (Figures 1A and 1B). EDS determines the average abundance of Mg, Al, Mo, and S is 20.8, 13.4, 12.4, and 53.4 in atomic percentage, respectively. Here, the Mo:S ratio is ~1:4.3 which is aligned to the composition of [Mo₃S₁₃]²⁻ anions. Elemental mapping of SEM micrographs further confirms the uniform distributions of Mg, Al, Mo, and S across the LDH-Mo₃S₁₃ crystallites (Figure S1). Infrared spectrum of the LDH-NO₃ shows a very strong peak at about 1370 cm⁻¹ that corresponds to the stretching vibration of NO₃⁻ anions,^{47,48} while for LDH-Mo₃S₁₃, the intensity of the 1370 cm⁻¹ peak is markedly diminished (Figure 1C). This demonstrates the exchange of NO₃⁻ with [Mo₃S₁₃]²⁻ anions between the positively charge nanosheets of LDH.

Sorption of CrO₄²⁻ from Water. The sorption experiments with Cr⁶⁺ show that LDH-Mo₃S₁₃ nanosheets are highly efficient for the sequestration of CrO₄²⁻ from water under acidic, neutral, and alkaline conditions (Table S2). At neutral pH, from 1000 ppb of a Cr⁶⁺ (CrO₄²⁻) spiked deionized water (DIW) solution, LDH-Mo₃S₁₃ can capture over 99.99% of Cr⁶⁺ leaving the final concentration below 1 ppb with a K_d value of $\sim 1.0 \times 10^7$ mL/g in 24 h (Tables S2 and S3). This concentration is well below the tolerance limits provided by the U.S. EPA (100 ppb) and WHO (50 ppb) for drinking water.^{49,50} Also, it should be noted that the value of K_d is higher to or comparable to other high performing materials known in the literature.^{18,24} At pH ~ 11, LDH-Mo₃S₁₃ can sequester over 98.7% of Cr⁶⁺ decreasing the final concentration to 13 ppb with the $K_d \sim 7.7 \times 10^4$ mL/g in 48 h. Conversely, at pH ~ 2, LDH-Mo₃S₁₃ can sequester over 74% of Cr⁶⁺ ions. The lower effectiveness at this pH may be attributed to the slow decomposition of LDH-Mo₃S₁₃. A similar experiment with a 10-fold higher Cr⁶⁺ concentration revealed similar adsorption efficiencies of over 79.2, 99.9, and 91.9% at pH ~ 2, 7, and 11, respectively (Table S2). Noticeably, despite the 10-fold increase of Cr⁶⁺ concentrations in the solutions, the residual concentrations, especially at neutral pH, remain below 10 ppb with K_d values of $\geq 10^6$ mL/g. Overall, our experiments show that LDH-Mo₃S₁₃ is highly efficient to remove Cr⁶⁺ from neutral and alkaline media.

The kinetic study of LDH-Mo₃S₁₃ for the removal of Cr⁶⁺ was conducted using initial concentrations of 1.0×10^3 and 1.0

$\times 10^4$ ppb of Cr^{6+} in DIW (Figures 2A and 2B, Tables S3 and S4). The experimental data were fitted with the pseudo-

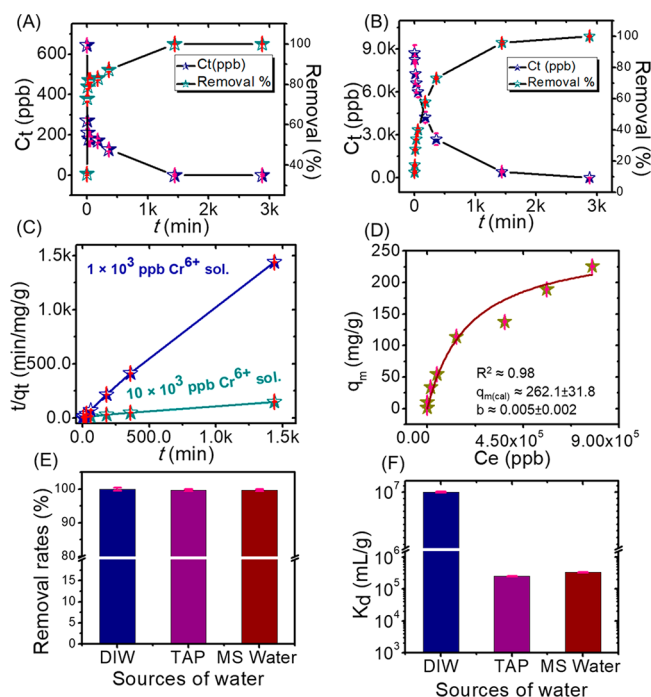


Figure 2. Adsorption kinetics for the residual concentrations (C_t) and removal rates of Cr^{6+} for 1.0×10^3 ppb (A) and 1.0×10^4 ppb (B) solutions in DIW; a comparison of the pseudo-second-order adsorption kinetics for 1×10^3 and 1×10^4 ppb solutions (C); adsorption capacity (q_m) vs equilibrium adsorption concentrations (C_e) (D); and a comparable study of the sorption of Cr^{6+} from CrO_4^{2-} spiked DIW, Tap, and Mississippi River water showing the removal rate (E) and distribution coefficient (F). These experiments were conducted using 10 mg of $\text{LDH-Mo}_3\text{S}_{13}$ in 10 mL ($V/m \sim 1000$ mL/g) of solutions at room temperature and pressure for a period of 48 h for (A–D) and 24 h for (E and F). The blue and blue-green stars in A and B represent C_t and the removal percentage, respectively. In C, the blue and blue-green stars represent 1.0×10^3 and 1.0×10^4 ppb of Cr^{6+} concentration; golden yellow stars in D represent equilibrium concentrations, and the red bars in A–F represent standard deviations.

second-order rate equation that yielded the coefficient of determination, $R^2 > 0.99$ (Figure 2C, Table S5). The pseudo-second-order rate constants were determined to be 0.0886 ± 0.0572 and 0.0015 ± 0.0003 g/mg·min $^{-1}$ for 1.0×10^3 and 1.0×10^4 ppb of Cr^{6+} solutions, respectively. The difference in rate constants is indicative of different sorption mechanisms at different initial concentrations. Apart from this, the higher rate constant indicates faster sorption kinetics of Cr^{6+} for the 1.0×10^3 ppb solution.

To evaluate Cr^{6+} uptake capacity and adsorption isotherm, we investigated the sorption of Cr^{6+} for a broad range of concentrations, 1.0×10^3 to 1.0×10^5 ppb (Figure 2D, Table S6). This study reveals that the sorption capacity increases with the increase of Cr^{6+} concentrations until it reaches an equilibrium. Our experiment determines that the maximum adsorption of Cr^{6+} is about 225 mg/g. This value of sorption capacity is superior to other high performing materials, namely $\text{LDH-MoS}_4 \sim 130$ mg/g and $\text{CoAl-LDH} \sim 93.5$ mg/g,^{18,24} cationic aluminum oxyhydroxides ~ 105.4 mg/g,⁵¹ and anion exchange resin (IRN78 ~ 63.5 mg/g),⁵¹ and comparable to

metal organic frameworks ($\text{MOR-1-HA} \sim 242$)⁵² and non-LDH cationic layered material (TJU-1 ~ 279 mg/g).²⁶ The experimentally obtained Cr^{6+} sorption isotherm data are fitted with the Langmuir model which yielded $R^2 \sim 0.98$ (Figure 2D). The Langmuir constant, b , was obtained as $0.005(2)$ L/mg and is comparable to other Cr^{6+} adsorbents.⁵³

To determine the effectiveness of $\text{LDH-Mo}_3\text{S}_{13}$ for Cr^{6+} removal in the presence of highly competitive ions in purified and naturally contaminated water, we investigated Cr^{6+} removal efficiency from tap water and Mississippi River water (collected from Vidalia, Louisiana) by spiking them with 1000 ppb of Cr^{6+} (CrO_4^{2-}) (Figures 2E and 2F, Table S7). This study showed that despite the presence of numerous ions, such as Ca^{2+} , Mg^{2+} , Na^+ , K^+ , HCO_3^- , SO_4^{2-} , and Cl^- , and other organic or inorganic constituents, $\text{LDH-Mo}_3\text{S}_{13}$ can remove $\geq 99.6\%$ of Cr^{6+} . Such a high removal rate is indicative of the higher efficiency of $\text{LDH-Mo}_3\text{S}_{13}$ for CrO_4^{2-} removal. Such an ultrahigh removal decreased its final concentration to <5 ppb with $K_d \sim 10^5$ mL/g. These values are close to the values obtained for DIW as discussed above. This finding suggests that $\text{LDH-Mo}_3\text{S}_{13}$ is effective at sequestration of Cr^{6+} even in the presence of highly competitive ions.

Removal of CrO_4^{2-} , a Surrogate of TcO_4^- , from the Simulated off-Gas Condensate of Hanford's LAW. To evaluate the removal efficiency of $\text{LDH-Mo}_3\text{S}_{13}$ for chromate, a nonradiogenic surrogate of $^{99}\text{TcO}_4^-$, we investigated the simulated LAW melter off-gas condensate of Hanford's WTP (Figure 3, Tables S8–S12). The simulant was prepared following a procedure described by Taylor-Pashow et al.^{4,9} The chemical composition of the simulant consists of high

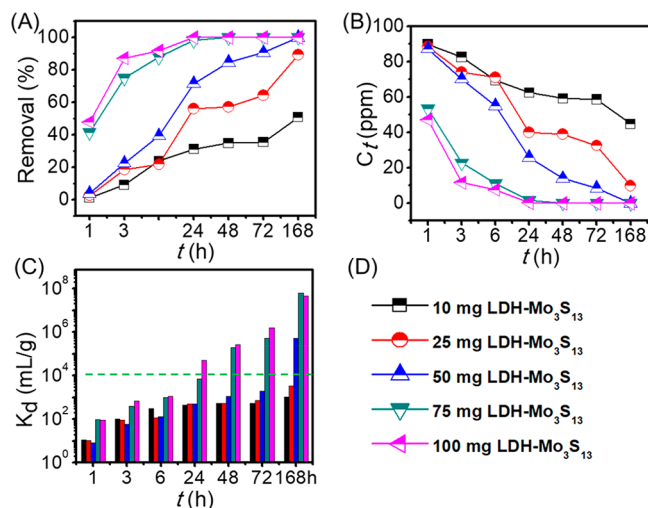


Figure 3. Time-dependent sorption of CrO_4^{2-} with an initial concentration of 9.086×10^4 ppb of Cr^{6+} as CrO_4^{2-} (CrO_4^{2-} equivalent concentrations $\sim 2.027 \times 10^5$ ppb) as a nonradioactive surrogate of TcO_4^- from the simulated off-gas condensate of Hanford's LAW melter with respect to various amounts of loading of the sorbents showing the removal (%) of CrO_4^{2-} (A), residual chromate concentration in the simulant (B), and the variation of K_d (mL/g) (C). Panel (D) represents the symbols that demonstrate the various amounts of $\text{LDH-Mo}_3\text{S}_{13}$ used for this experiment. These experiments were conducted using 10 mg of $\text{LDH-Mo}_3\text{S}_{13}$ in 10 mL ($V/m \sim 1000$ mL/g) of simulant solutions at room temperature and pressure for a period of 1 h through 68 h. Each experiment was replicated four times, and the average was considered to plot the diagram and analyze the results.

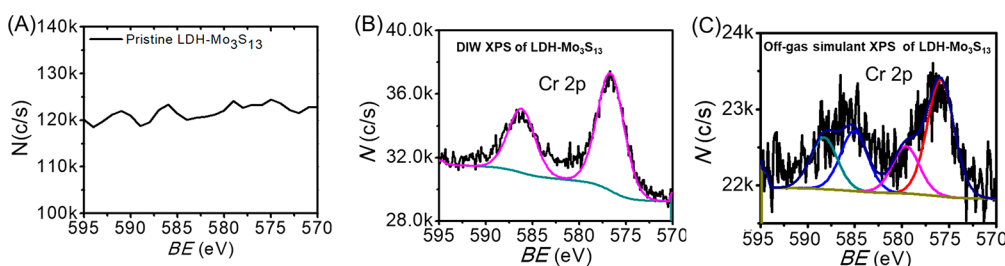


Figure 4. XPS spectra were collected for the pristine LDH–Mo₃S₁₃ showing the absence of Cr (A), and the DIW XPS data (B) and the simulant XPS (C) show the presence of Cr. The DIW data was collected after the sorption experiments for 1×10^5 ppb of Cr⁶⁺ using 10 mg of LDH–Mo₃S₁₃ in DIW, and the simulant XPS was collected for 9.086×10^4 ppb of the Cr⁶⁺ containing solution after treating the solutions using 100 mg of LDH–Mo₃S₁₃. Each experiment was conducted with 10 mL of solutions.

concentrations of anions, e.g., F[−], Cl[−], CO₃^{2−}, NO₃[−], BO₃^{3−}, NO₂[−], SO₄^{2−}, and B₄O₇^{2−} at pH $\sim$ 7.5.

We observed that for the 10 and 25 mg loading of LDH–Mo₃S₁₃, the maximum removal of Cr⁶⁺ was about 50.7 and 89.3%, respectively, from a 9.086×10^4 ppb Cr⁶⁺ spiked simulated solution after 7 days of interactions (Figures 3A and 3B, Tables S8 and S9). This value of removal rate and K_d are remarkably higher than that of the LDH–NO₃ and (NH₄)₂Mo₃S₁₃, suggesting the importance of the LDH–Mo₃S₁₃ for the ultrahigh removal of Cr⁶⁺ ions from the simulated solution (Table S13). Conversely, for an interaction of 50 mg of LDH–Mo₃S₁₃, the Cr⁶⁺ removal reached >99.7% in 7 d. This leads to the residual concentration of Cr⁶⁺ to $\sim$ 35 ppb and K_d values of $\sim 5 \times 10^5$ mL/g (Figure 3C, Table S10). With a higher loading of 75 and 100 mg of LDH–Mo₃S₁₃, the removal of Cr⁶⁺ reached over 99.9 and 99.7% in only 2 and 1 d, respectively (Figure 3C, Tables S11 and S12). Importantly, after 3 days, the residual concentrations reach trace levels, 23 (75 mg) and 6 ppb (100 mg), and the K_d values reach $\geq 10^5$ mL/g. Besides, after 7 d of exposure, removal percentages reach $\sim$ 100%, residual concentrations become as low as <0.1 ppb, and the K_d reach $\sim 10^7$ mL/g. Herewith, LDH–Mo₃S₁₃ demonstrates itself as a highly efficient sorbent of CrO₄^{2−} and by extension also its TcO₄[−] surrogate from the simulated off-gas condensate of the LAW melter.

Understanding the Adsorption Mechanisms. The EDS and elemental mapping of the solid samples after the reaction with CrO₄^{2−} in DIW and simulated solution show the presence chromium across the sorbents (Figures S2 and S3). The SEM and TEM images of the solid sorbents after the adsorption study show the retention of the platelike morphology (Figures S2 and S3). The TEM image of the postinteracted solids shows the presence of hexagonal crystallites similar to pristine LDH–Mo₃S₁₃. For the simulant treated samples, the crystallites' surface becomes deteriorated which could be the impact of highly concentrated anions of F[−], Cl[−], and others.

XPS analysis of the pristine and loaded LDH–Mo₃S₁₃ revealed the chemical state of the Mo, S, and Cr ions (Figure 4, Table S14). The post-treatment solid samples of LDH–Mo₃S₁₃ show the doublet of the peaks in the range of 570–595 eV corresponding to Cr 2p orbitals. As expected, these peaks are absent in the pristine LDH–Mo₃S₁₃. Specifically, the postadsorbed samples in water show the peaks at 586.2 eV/576.7 eV corresponding to the Cr³⁺ oxidation state, probably the formation of Cr₂O₃. Hence, the doublet of peaks is the result of the spin–orbit splitting of the Cr³⁺ 2p orbitals.^{18,54} For the samples that were treated with simulated solutions, the XPS show two pairs of peaks at 577.5 eV/585.5 eV and 589.1 eV/579.5 eV. The bands at 577.5 eV/585.5 eV can be originated

from Cr³⁺, while the peaks at higher energy are attributed to Cr⁶⁺ ions of the CrO₄^{2−}.^{54,55} The S 2p peaks that are observed in the range of 169.2–171.2 eV can be attributed to the oxidation of sulfides to sulfate anions (Figure S4).^{56–58} Similarly, for the Mo 3d, the weak peaks that are observed at about 236 eV are the results of the partial oxidation of Mo⁴⁺ to Mo⁶⁺ of the Mo⁴⁺₃S₁₃ moieties of LDH–Mo₃S₁₃ (Figure S4).^{59,60}

Furthermore, ICP analysis shows the presence of molybdenum ions in the postinteraction solutions. A quantitative analysis by ICP MS determined that $\sim$ 21% of Mo of LDH–Mo₃S₁₃ dissolves. The presence of molybdenum ions can be attributed to oxidation dissolution of Mo⁴⁺ of the [Mo⁴⁺₃S₁₃]^{2−} to Mo⁶⁺ ions. The dissolution of Mo⁶⁺ results in the concurrent dissolution of sulfide which is likely to oxidize to sulfate/sulfite anions. Hence, this study reveals that a cooperative oxidation of both sulfur and molybdenum likely enhances the reduction of Cr⁶⁺ to Cr³⁺.

The Cr K-edge XANES spectra of LDH–Mo₃S₁₃ samples were collected after reaction with CrO₄^{2−} in DIW and the simulated LAW off-gas condensate, together with Na₂CrO₄ as a model compound (Figure 5A). For Na₂CrO₄, a sharp edge peak at 5994.0 eV is observed which is characteristic of the four coordinated character of the hexavalent chromate ion (CrO₄^{2−}).^{61,62} Conversely, this peak at 5994.0 eV nearly completely disappeared for the DIW interacted sample, suggesting that Cr⁶⁺ was hardly detectable and the dominant species was Cr³⁺ reduced from CrO₄^{2−}. Additionally, there was another weak peak at 5991.2 eV, which is a typical pre-edge peak for Cr₂O₃ or Cr(OH)₃.^{54,63} The solid sorbent obtained from the simulant experiment shows the presence of the peak at 5994.0 eV representative of CrO₄^{2−} ions. It is estimated from the Cr K-edge XANES there is $\sim$ 15% CrO₄^{2−} and $\sim$ 85% Cr³⁺ in this sample. In addition, there was a new peak at 5988.2 eV, together with a bump at 5991.2 eV corresponding to the pre-edge peak observed for the DIW treated sorbent. These two features are attributed to the split of the pre-edge and are assigned to Cr 1s transitions to 3d (*t*_{2g}) and 3d (*e*_g) electronic states, respectively, in the polyhedra of Cr³⁺ ions.⁶⁴ Overall, the Cr K-edge of XANES spectra of Na₂CrO₄ and postinteracted LDH–Mo₃S₁₃ in DIW and the simulant illustrate features that are consistent with the Cr valence state and change in coordination number. Notably, the reductive precipitation of Cr⁶⁺ $\rightarrow$ Cr³⁺ could be attributed to the presence of highly redox active sulfide anions in LDH–Mo₃S₁₃. Hence, the higher redox potentials of Tc⁷⁺ (TcO₄[−]/Tc⁴⁺ is $E^\circ \sim +0.74$ V)²¹ and Cr⁶⁺ (CrO₄^{2−}/Cr³⁺ is $E^\circ \sim +1.23$ V)²⁰ enable their reduction by the mono- and disulfides species, S^{2/1−} $\rightarrow$ S^{0/+} + *ne*[−] (e.g., E° of S^{2−}/SO₄^{2−} is ~ -0.22 V),²³ of LDH–Mo₃S₁₃. This

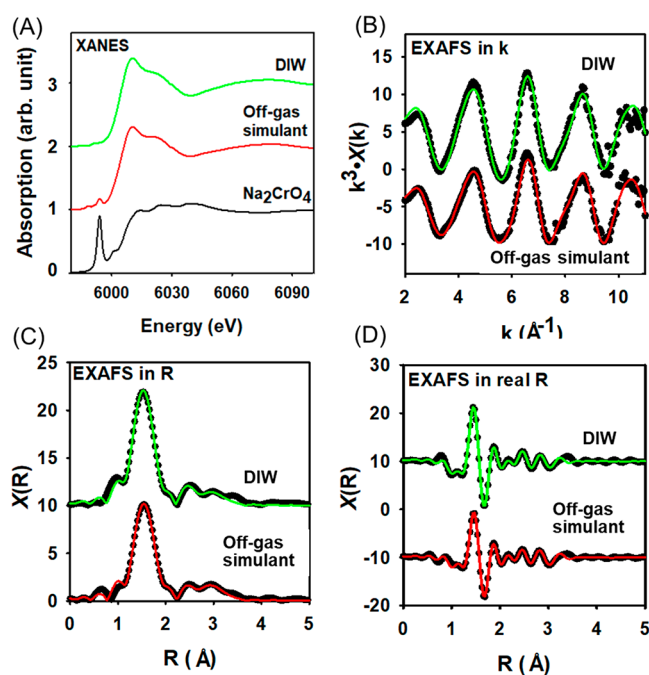


Figure 5. Cr K-edge XANES and EXAFS spectra of two LDH– Mo_3S_{13} samples after Cr treatments in DIW and an off-gas simulant, XANES, together with the spectrum of Na_2CrO_4 (A), EXAFS data in k space (B), EXAFS data in R space (C), and EXAFS data in real R space (D). Dotted and solid-colored lines in EXAFS represent experimental and fitted data, respectively.

reduction immobilizes $\text{Tc}^{4+}/\text{Cr}^{3+}$ salts by the deposition of technetium and chromium containing solids on the sorbent. Since this reductive precipitation is the dominant mechanism for the sequestration of Cr^{6+} and the sorbents are contaminated with the precipitated solids, this material is unlikely to be reusable.

XPS and XANES predominantly show the removal of chromium involves reduction of Cr^{6+} to Cr^{3+} . The CrO_4^{2-} anion adopts a tetrahedral geometry, while the Cr^{3+} species remains predominantly in an octahedrally coordinated environment. This change in coordination number and the geometry are further demonstrated by Cr K-edge EXAFS data (Figure 5B–D). The fitted EXAFS parameters of these samples are summarized in Table 1, in comparison with the X-ray

diffraction data of Cr_2O_3 .⁴⁶ The Cr K-edge EXAFS data of the sorbent sample exposed to CrO_4^{2-} in DIW was fitted with octahedral Cr–O paths at a Cr–O distance of 1.971 ± 0.005 Å with a fitted coordination number of 5.10 ± 0.39 and three Cr–Cr paths at 2.54 ± 0.03 , 3.09 ± 0.03 , and 3.38 ± 0.03 Å, respectively, with the corresponding fixed coordination numbers obtained from Cr_2O_3 X-ray diffraction data.⁴⁶ The Cr–O octahedron in Cr_2O_3 displays three Cr–O bonds at a distance of 1.964 Å and three Cr–O bonds at a distance of 2.013 Å, but the EXAFS data fitting was kept with the same Cr–O bond distance to limit the fitting parameters. The fitted Cr–O bond distance of 1.971 Å and the coordination number of 5.1 are comparable to those in Cr_2O_3 , but the fitted bond distances for two of three Cr–Cr paths show a larger variation compared to those in Cr_2O_3 . However, the overall EXAFS data fitting of this sample was acceptable as measured by the R factor of 0.0026 (Table 1). Thus, the Cr K-edge EXAFS spectra of the sorbent exposed to CrO_4^{2-} in DIW confirmed that the Cr species associated with the sorbent was likely Cr_2O_3 or $\text{Cr}(\text{OH})_3$.

The Cr K-edge EXAFS data of the sorbent sample exposed to CrO_4^{2-} in the off-gas simulant was also fitted using the Cr_2O_3 structure model. Although its Cr K-edge XANES indicated that 15.4% of Cr was CrO_4^{2-} , the EXAFS data fitting did not permit more parameters with the addition of another phase. In general, the fitted EXAFS parameters for the sorbent exposed to CrO_4^{2-} in the off-gas simulant were similar to those of the sorbent exposed to CrO_4^{2-} in DIW (Table 1), and the overall fitting of this sample was acceptable as measured by the R factor of 0.0039 (Table 1). However, the fitted Cr–O coordination number for this sample was 3.72 ± 0.35 (Table 1), which was significantly smaller than the Cr–O coordination number of 5.1 in the sample retrieved from DIW and the Cr–O coordination number of 6 in Cr_2O_3 . Except for the common error of up to 20% for the fitted coordination number from the EXAFS data fitting, this discrepancy might also indicate that a small portion of Cr in this sample was tetrahedral CrO_4^{2-} . Thus, the Cr K-edge EXAFS spectra of the sorbent exposed to CrO_4^{2-} in the off-gas simulant confirmed that the dominant Cr species associated with the sorbent was likely Cr_2O_3 or $\text{Cr}(\text{OH})_3$, but a small amount of CrO_4^{2-} was also present, consistent with its Cr K-edge XANES data discussed above.

Table 1. Cr K-Edge Fitting Data for the Cr Species after Interactions of LDH– Mo_3S_{13} with CrO_4^{2-} in Water and in the Simulated LAW off-Gas Condensate

samples	path	bond distance (Å)	coordination number	Debye–Waller factor, σ^2 (Å ²)	ΔE_0 (eV)	R-factor
DIW	Cr–O2	1.971 ± 0.005	5.10 ± 0.39	0.0026 ± 0.0007	3.2 ± 0.8	0.0026
	Cr–Cr1	2.54 ± 0.03	1	0.0149 ± 0.0047		
	Cr–Cr2	3.09 ± 0.03	3	0.0155 ± 0.0039		
	Cr–Cr3	3.38 ± 0.03	3	0.0111 ± 0.0027		
off-gas simulant	Cr–O2	1.976 ± 0.005	3.72 ± 0.35	0.0011 ± 0.0009	3.7 ± 1.0	0.0039
	Cr–Cr1	2.56 ± 0.05	1	0.0185 ± 0.0072		
	Cr–Cr2	3.09 ± 0.05	3	0.0216 ± 0.0072		
	Cr–Cr3	3.39 ± 0.02	3	0.0108 ± 0.0025		
Cr_2O_3	Cr–O1	1.964	3			
	Cr–O2	2.013	3			
	Cr–Cr1	2.650	1			
	Cr–Cr2	2.888	3			
	Cr–Cr3	3.425	3			

Mo L3-edge XANES reveals the presence of the dipole allowed $2p \rightarrow 4d$ transition of Mo^{4+} at 2523.8 eV, indicating that Mo predominantly remains as Mo^{4+} in the postreacted solids, in agreement with XPS results (Figure S5).⁶⁵ The XANES of S K-edge shows peaks at about 2471.4, 2471.6, and 2471.7 eV for the pristine, water, and simulant treated samples, respectively. These peaks demonstrate the presence of sulfide species. The peak at 2481.9 eV can be ascribed to S^{6+} oxidation states which suggests concomitant oxidation of some of the sulfide groups of $\text{LDH-Mo}_3\text{S}_{13}$. Importantly, the intensity of the S^{6+} peaks sharply increases from the pristine to the simulant treated $\text{LDH-Mo}_3\text{S}_{13}$, indicating a higher propensity of sulfide oxidation in the simulated solutions which further supports the reductive precipitation of Cr^{6+} to Cr^{3+} .

Altogether, an insight of the sorption mechanisms demonstrates that reductive precipitation is the dominant mechanism for the removal of $\text{TcO}_4^-/\text{CrO}_4^{2-}$; nevertheless, a small contribution of ion-exchange and/or surface adsorption cannot be ruled out, especially for the simulated off-gas condensate of the LAW melter. The reductive precipitation mechanism requires the use of S^{n-} ($n = 1, 2$) and Mo^{4+} of the $\text{LDH-Mo}_3\text{S}_{13}$ as the reductants to precipitate out the Cr^{3+} species. Overall, this study reveals that $\text{LDH-Mo}_3\text{S}_{13}$ is a high performing sorbent of $\text{TcO}_4^-/\text{CrO}_4^{2-}$ from defense nuclear legacy waste. We believe this finding advances our knowledge and points to design principles of materials that can accomplish excellent efficiency in complex high ionic strength solutions for the sequestration of pertechnetate and chromate anions from aqueous solutions.

■ ASSOCIATED CONTENT

SI Supporting Information

The Supporting Information is available free of charge at <https://pubs.acs.org/doi/10.1021/acs.est.1c08766>.

Details on uptake study as well as characterization of pristine and postadsorption $\text{LDH-Mo}_3\text{S}_{13}$ (PDF)

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<https://pubs.acs.org/doi/10.1021/acs.est.1c08766>

Notes

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The authors declare no competing financial interest.

■ ACKNOWLEDGMENTS

This work was partially supported by the US Department of Energy Minority Serving Institution Partnership Program (MSIPP) managed by the Savannah River National Laboratory under SRNS contract (RFP No. 0000542525 and 0000458357). S.C.R. is thankful to the NSF Division of Chemistry (NSF-2100797). All of the ICP-MS analysis was conducted at JSU's core research centers supported by RCMI (NIH grant: 1U54MD015929). X-ray Absorption Spectroscopy was performed at the SXRMB beamline of the Canadian Light Source, a national research facility of the University of Saskatchewan, which is supported by the Canada Foundation for Innovation (CFI), the Natural Sciences and Engineering Research Council (NSERC), the National Research Council (NRC), the Canadian Institutes of Health Research (CIHR), the Government of Saskatchewan, and the University of Saskatchewan. This work made use of the Keck-II facility of Northwestern University's NUANCE Center, which has received support from the SHyNE Resource (NSF ECCS-2025633), the IIN, and Northwestern's MRSEC program (NSF DMR-1720139).

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