

Role of Manganese in Accelerating the Oxidation of Pb(II) Carbonate Solids to Pb(IV) Oxide at Drinking Water Conditions

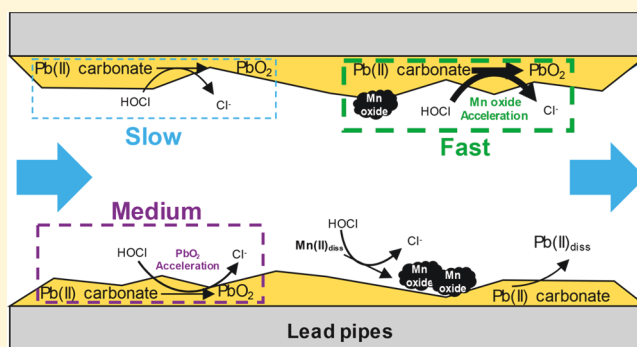
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Supporting Information

ABSTRACT: Pb(II) carbonate solids are corrosion products that form on the inner surfaces of lead service lines (LSLs) and can be oxidized by free chlorine to form Pb(IV) oxide (PbO₂). The formation of PbO₂ can maintain low dissolved lead concentrations in drinking water, but PbO₂ can dissolve if a free chlorine residual is not maintained. Experiments demonstrated that the oxidation of Pb(II) carbonate by free chlorine was faster with manganese (Mn). Without Mn(II), the oxidation of Pb(II) carbonate was an autocatalytic process. With Mn(II), the overall oxidation rate was 2 orders of magnitude faster than without Mn(II). X-ray diffraction and free chlorine consumption profiles indicated that δ -MnO₂ was formed within several minutes of the reaction of Mn(II) with free chlorine, and δ -MnO₂ catalyzed the oxidation of Pb(II) carbonate by free chlorine. Free chlorine consumption profiles for Pb(II) carbonate with and without Mn(II) were interpreted based on the kinetics and stoichiometry of the underlying chemical reactions. These findings highlight the importance of Mn in accelerating the formation of PbO₂ in water with Pb(II) carbonate solids and free chlorine, and it may help explain why PbO₂ is observed on LSLs of some but not all water systems that use free chlorine.



INTRODUCTION

Ingestion of lead-contaminated drinking water is one of the major pathways for human exposure to lead.^{1–3} Lead service lines (LSLs) and premise plumbing are the largest sources of lead to drinking water.² In the United States, the Lead Copper Rule has an action level of 0.015 mg/L lead for the 90th percentile of specific homes' 1-L first draw samples of tap water that triggers additional actions if it is not met.⁴ Although the 1986 Safe Drinking Water Act Amendments banned lead pipes, fittings, fixtures, solder, and flux, millions of partial or whole lead service lines are still present in the United States.^{3,5,6}

Lead concentrations in drinking water are controlled by the solubility of lead corrosion products that form in lead pipes and premise plumbing.^{7–12} These corrosion products include Pb(II) carbonate, oxide, and phosphate solids as well as Pb(IV) oxide (PbO₂).^{9,10,13,14} Previous studies have shown that the oxidation of Pb(II) carbonate, including cerussite (PbCO₃) and hydrocerussite (Pb₃(CO₃)₂(OH)₂), by free chlorine, will lead to the formation of lead(IV) oxide.^{8,15,16} Liu et al.¹⁶ reported an autocatalytic reaction during Pb(II) carbonate oxidation by chlorine. Wang et al.⁸ found that the identity and rate of formation of the PbO₂ phases produced from reaction of various Pb(II) species with free chlorine were affected by the initial solid or dissolved Pb(II) species,

dissolved inorganic carbon (DIC), pH, and free chlorine concentration.¹⁷ PbO₂ has two phases, scrutinite (α -PbO₂) and plattnerite (β -PbO₂), both of which have a low solubility compared with Pb(II) corrosion products. Triantafyllidou et al.¹⁸ reported that lead release from lead pipes coated with Pb(IV) oxides was consistently much lower than from pipes coated with Pb(II)-containing solids and hence concluded that formation of PbO₂ in LSLs could be an effective lead corrosion control strategy.¹⁸ The depletion of free chlorine, lower oxidation reduction potential (ORP) disinfectants (such as monochloramine), and the presence of reductants (natural organic matter, Br[−], I[−], Fe(II), Mn(II), etc.) may lead to the dissolution of PbO₂.^{19–25} The increases in lead concentrations in Washington, DC tap water that began in late 2000 were triggered by a change in disinfectant from higher ORP free chlorine to lower ORP monochloramine.²⁶ While PbO₂ has been observed on many LSLs from water systems that use free chlorine,^{10,12,14,18} PbO₂ has not always been observed in the scales of LSLs for such systems.^{10,12,18} The limited range of conditions over which PbO₂ is stable may limit its observation.

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While PbO_2 can be formed by reaction with free chlorine, it is only stable when a free chlorine residual is present, and PbO_2 can be reduced by reductants such as natural organic matter, I^- , Br^- , Fe(II) , Mn(II) , and even water (Figure 1) when free chlorine is depleted.^{19,21–25}

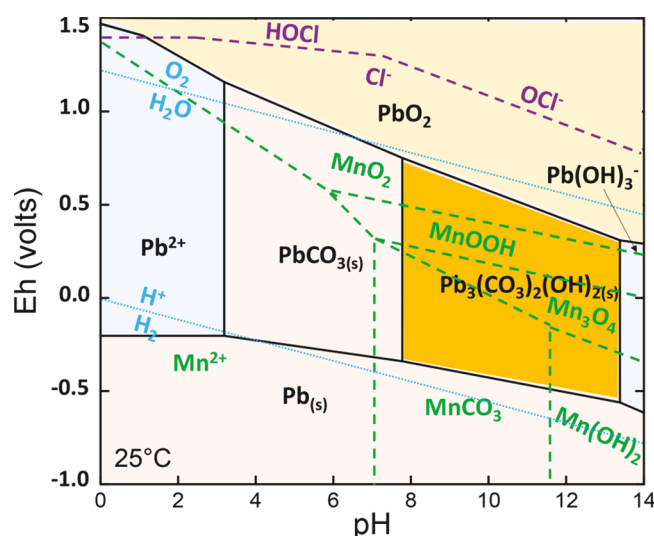


Figure 1. Eh–pH diagram of Pb–Mn–Cl at 25 °C and 1 bar total pressure. Predominance areas were calculated for conditions with 2 mM Pb, 0.1 mM Mn, and 0.2 mM inorganic carbon. Boundaries of predominance areas are shown for chlorine (purple), lead (black), and manganese (green) species. Blue lines indicate the stability limits of water.

Manganese accumulation has been reported in different water distribution systems.^{11,27–30} The accumulated Mn in water distribution systems is generated from oxidation of soluble Mn(II) by free chlorine, oxygen, and other oxidants to form insoluble Mn(III) and Mn(IV) solids.^{27,28,31} In addition, the activity of manganese depositing microbes can also give rise to the formation of manganese oxides in drinking water networks.^{27,32–35} Gerke et al.³³ reported that birnessite (Mn(III) and Mn(IV)), hollandite (Mn(II) and Mn(IV)), and braunite (Mn(II) and Mn(IV)) were the main Mn oxides/oxyhydroxide in water distribution systems. The interactions between Pb and Mn may represent a sink or source for lead and Mn in drinking water, and previous studies have investigated the adsorption of Pb(II) on Mn(III, IV) oxides^{36–38} as well as the oxidation of Mn(II) by lead(IV) oxide.^{25,39} Matocha et al.³⁹ revealed that Pb(II) was adsorbed as inner-sphere complexes on both birnessite and manganite, and they found no evidence of Pb(II) oxidation by birnessite and manganite based on X-ray absorption fine structure (XAFS) spectra. Shi and Stone²⁵ studied the redox reaction between PbO_2 and Mn(II), and they found that Mn(II) could be oxidized to Mn(III) or Mn(IV) (hydr)oxides (MnOOH and $\sigma\text{-MnO}_2$) by PbO_2 . This result is consistent with Figure 1, which illustrates that PbO_2 is a stronger oxidant than Mn(IV) oxide.²⁵

Although recent investigations have examined the fundamental surface reactions between Pb- and Mn-containing solids, the nature of these interactions at conditions relevant to drinking water supply and their implications for tap water quality had not been sufficiently explored. Such interactions can be important in affecting lead concentrations in drinking

water. Recently, Trueman et al.⁴⁰ reported that without free chlorine, Mn(II) increased Pb release mainly because of the oxidation of lead by oxygen on MnO_2 deposits or by Mn(IV) itself. However, this may not be the scenario when free chlorine is present in the system. The goal of this work was to identify the products and pathways of the reaction of dissolved Mn(II), Pb(II) carbonate, and free chlorine. Specific attention was paid to the heterogeneous catalytic effect of Mn solids on hydrocerussite and cerussite oxidation by chlorine. The oxidation of Pb(II) is a necessary step to form Pb(IV) oxide. This process may take place during drinking water distribution at locations where lead-containing materials and dissolved or solid Mn coexist. In this study, we investigated the oxidation of Pb(II) from Pb(II) carbonate solids with and without the presence of Mn(II) at various pH values. We performed batch experiments to probe the products and dynamics of the reactions, and these were supplemented by kinetic models based on mass balances and the stoichiometry of the reactions. Flow-through experiments provided additional information on reaction rates with a lower but sustained supply of free chlorine that is more directly relevant to the actual conditions in LSLs.

MATERIALS AND METHODS

Materials. Hydrocerussite and cerussite were purchased from Sigma-Aldrich. A free chlorine stock solution was prepared using NaOCl (Fisher Chemical, 4–6% w/w) and ultrapure water. Plattnerite ($\beta\text{-PbO}_2$) was purchased from Acros Organics. Scrutinyite ($\alpha\text{-PbO}_2$) was synthesized in a 500 mL polypropylene batch reactor with hydrocerussite as a precursor with the pH maintained at 10. The synthesis was conducted at carbonate-free conditions by sparging the solution with air that had been scrubbed of CO_2 using a 1 M NaOH solution. During the scrutinyite synthesis, the free chlorine concentration was maintained at 50 mg/L (0.70 mM) as Cl_2 . The XRD pattern (Figure S1) confirmed the high purity of synthesized scrutinyite. Synthetic $\delta\text{-MnO}_2$ (Figure S2) was prepared by reacting KMnO_4 with MnCl_2 at basic pH following a published method.^{41,42} The identities of the solids synthesized were confirmed by X-ray diffraction (XRD). Reagent grade NaHCO_3 , NaNO_3 , NaOH, MnCl_2 , and concentrated HNO_3 were purchased from Fisher Scientific. All solutions were prepared using ultrapure water (resistivity >18.2 M Ω -cm, Milli-Q, Millipore Corp., Milford, MA).

Experimental Design. Three sets of batch experiments were conducted. To study cerussite or hydrocerussite oxidation in isolation, the first set of experiments was conducted with 200 mL of 2 mM cerussite or hydrocerussite suspensions with a DIC concentration of 0.2 mM (2.4 mg/L as C), 0.28 ± 0.03 mM free chlorine (20 ± 2 mg/L as Cl_2), and pH 6.5 to 8.5 ± 0.2 maintained by 0.1 M HNO_3 and 0.1 M NaOH. The high chlorine concentration was chosen to facilitate the development of a kinetic model for oxidation of Pb(II) in Pb(II) carbonate solids by free chlorine. A lower free chlorine concentration will yield a relatively short reaction time, and we may not be able to capture some crucial features of the reactions in the batch experiment. To investigate the effect of an initial amount of scrutinyite and plattnerite on lead carbonate oxidation, the second set of experiments had certain amounts of scrutinyite or plattnerite added to the suspensions of cerussite or hydrocerussite with the same DIC, free chlorine, and pH as in the first set of experiments. In support of this second set of experiments, control experiments were conducted with just scrutinyite or plattnerite and free chlorine

and no Pb(II) carbonate solids. In the third set of experiments, the effect of Mn(II) was studied by adding 0.1 or 1 mL of 0.25 M MnCl_2 solution to the cerussite or hydrocerussite suspensions to reach 0.1 mM or 1 M initial Mn(II) concentrations. All batch experiments were performed in 250 mL glass bottles with their contents stirred with polypropylene-coated stir bars at a speed of 500 rpm on a multiposition stirrer (VARIOMAG, Thermo, U.S.A.) at room temperature ($21 \pm 1^\circ\text{C}$).

For all batch experiments, 5 mL samples of suspension were collected at selected times. Reactions between solutes and the surfaces of solids were quenched by immediately filtering suspensions through 0.05 μm pore diameter poly(ether sulfone) (PES) membranes (Tisch Scientific, U.S.A.) with a collection of 2 mL of filtrate. Samples after this additional filtration step were acidified to 2% HNO_3 and preserved for dissolved lead analysis by ICP-MS. Samples of the suspension that included solids were collected at the beginning of the experiment and after 5 and 24 h of reaction. The suspensions were centrifuged and freeze-dried to yield solids for further analysis.

Flow-through experiments with suspensions of lead carbonate reacting with free chlorine were conducted in continuously stirred tank reactors (CSTRs). In contrast to the batch reactors in which free chlorine was added at a relatively high initial concentration, the flow-through reactors could provide a steady supply of free chlorine at a lower concentration that is more directly relevant to the drinking water supply. The reactions were examined both in the absence and presence of 0.1 mM $\delta\text{-MnO}_2$ at the ambient room temperature ($21 \pm 1^\circ\text{C}$). The $\delta\text{-MnO}_2$ solid was employed instead of Mn(II) solution in the flow-through experiments because any Mn(II) in the solution that was not oxidized by the initial influx of free chlorine would have been flushed out of the reactors (Figure S3). The volume of each reactor was 40 mL, and lead carbonate suspensions were loaded to a concentration of 1 mM as Pb. Specific amounts of $\delta\text{-MnO}_2$ that provided the same Pb:Mn ratio as a set of the batch experiments were added to CSTRs. Mixing was achieved by continuous stirring with a polypropylene-coated magnetic stir bar at 250 rpm. A 0.22 μm mixed cellulose filter membrane at the reactor outlet prevented the loss of solids from the reactor. Influent was pumped into the reactor at 2 mL/min using a peristaltic pump (Cole-Parmer), so the reactor hydraulic residence time (τ) was 20 min. The pH and free chlorine concentrations of the effluents were monitored, and aqueous samples were periodically collected for dissolved Pb(II) concentration analysis. The influents were prepared in 4-L polypropylene bottles covered with aluminum foil to minimize CO_2 exchange with the atmosphere and prevent photo-degradation of free chlorine. Addition of an aliquot of 1 M NaHCO_3 solution to the bottles provided the DIC concentration of 0.02 mM (2.4 mg/L) as C. An aliquot of a NaOCl stock solution was added to reach a concentration of 0.034 mM (2.4 mg/L) as Cl_2 . The pH was adjusted to the target values by addition of concentrated HNO_3 or freshly prepared 0.5 M NaOH solutions to reach $\text{pH } 7.5 \pm 0.2$. The flow-through experiments were conducted for 2000 min (i.e., 100 residence times).

Analytical Methods. The residual free chlorine concentration was measured with the standard DPD method.⁴³ Dissolved lead was analyzed by inductively coupled plasma-mass spectrometry (ICP-MS) (PerkinElmer ELAN DRC II).

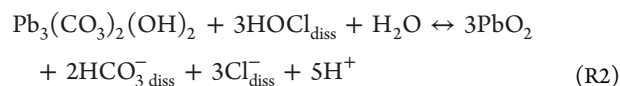
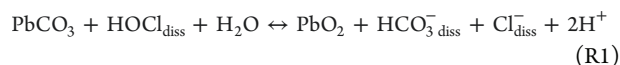
Solution pH was measured with a glass pH electrode and pH meter (Accumet). XRD patterns were collected using Cu $K\alpha$ radiation (Bruker d8 Advance X-ray diffractometer). A Scanning Electron Microscopy (SEM) examination was performed with FEI Nova NanoSEM 230. The specific surface areas of dry solids were determined by the Brunauer–Emmett–Teller- N_2 (BET-N_2) adsorption (Quantachrome Instruments) method.

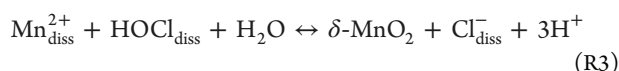
Model Development. All the models were developed using constraints of mass balances and the stoichiometry of the oxidation–reduction reactions. The specific rate equations used are presented in the Results and Discussion section, and a brief overview of the modeling approach is provided here. The conceptual model of the processes occurring is presented in Figure 2. The formation of PbO_2 included noncatalytic



Figure 2. Conceptual model of PbO_2 formation from cerussite considering PbO_2 autocatalytic and MnO_2 catalytic effects. Both the hypochlorite ion and hypochlorous acid can oxidize Pb(II) to Pb(IV), and only the hypochlorite ion is shown here for the sake of simplicity.

oxidation of Pb(II), autocatalytic oxidation of Pb(II) with PbO_2 formed *in situ*, and catalytic oxidation of Pb(II) with Mn oxides. The consumption of free chlorine included noncatalytic oxidation of Pb(II), autocatalytic oxidation of Pb(II) with PbO_2 formed *in situ*, oxidation of Mn(II) to $\delta\text{-MnO}_2$, and catalytic oxidation of Pb(II) with $\delta\text{-MnO}_2$ formed *in situ*. In this study, *in situ* means in the reactor. For cerussite and hydrocerussite oxidation, the consumption of free chlorine was assumed to equal the formation of PbO_2 in molar concentration (reactions R1 and R2). For cerussite and hydrocerussite reaction with free chlorine in the presence of Mn, the consumption of free chlorine included oxidation of Pb(II) as well as oxidation of Mn(II) (reaction R3). Mole balance equations were developed for free chlorine and PbO_2 that included terms for the rates of oxidation of Pb(II) carbonate directly, catalyzed by PbO_2 , and catalyzed by MnO_2 . The pH dependence was not accounted in model development since the data examined in comparison with the model were all collected at $\text{pH } 7.5 \pm 0.2$. For the flow-through experiment, the free chlorine in the influent and $\delta\text{-MnO}_2$ concentrations were treated as constants since it was continuously supplied from a well mixed 4 L reservoir. Although catalytic reactions were surface-related reactions, the surface area was incorporated into reaction rate constants in our model development. Differential equations obtained from the mass balance equations were solved by MATLAB R2018a. Optimization of the fit of data to model by adjusting multiple parameters at the same time to obtain the minimum coefficient of determination (R^2).





RESULTS AND DISCUSSION

The Catalytic Effect of PbO₂ on Lead Carbonate Oxidation. The trends in the consumption of free chlorine for oxidation of cerussite provided insights into an autocatalytic reaction in which the production of an initial amount of PbO₂ accelerated the rate of further PbO₂ formation (Figure 3). We

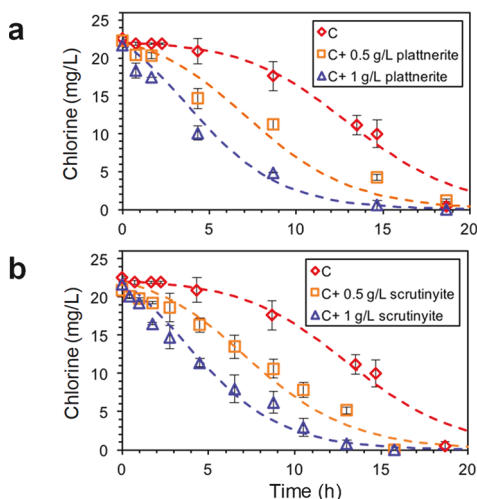


Figure 3. Oxidation of cerussite (C) by chlorine with and without (a) plattnerite and (b) scrutinyite. Conditions: pH = 7.5 ± 0.2, initial free chlorine = 0.31 mM (22 mg/L as Cl₂), initial cerussite = 2 mM as Pb, DIC = 0.2 mM as C. Experimental results are shown as points, and modeling results are shown as dashed lines. The average values from duplicate experiments are shown, and the error bars represent the standard deviation from the duplicate experiments.

observed the oxidation occurring with a lag stage followed by a rapid stage for both cerussite and hydrocerussite. These two stages of oxidation for lead carbonate were also observed by other researchers.¹⁶ The addition of PbO₂ (either plattnerite or scrutinyite) accelerated cerussite oxidation by free chlorine and dramatically shortened the lag stage of the reaction (Figure 3). With the increasing amount of PbO₂ (from 0.5 to 1 g/L), the free chlorine consumption speed increased, indicating the increasing rate of cerussite oxidation. A shortened lag stage was still observed with high PbO₂ concentration (1 g/L). Control experiments determined that plattnerite and scrutinyite by themselves did not accelerate chlorine degradation (Figure S4). While the data presented in Figure 3 are for cerussite, similar trends were observed for hydrocerussite and are shown in the Supporting Information (Figure S6).

When comparing the rate of chlorine consumption associated with oxidation of cerussite and hydrocerussite, the rate was faster with hydrocerussite (Figure S6), which was consistent with previous findings.^{16,44} The lower oxidation speed of cerussite by free chlorine may be due to cerussite's lower specific surface area (3.28 m²/g versus 10.70 m²/g for hydrocerussite) that may be related to lower concentration of active surface sites for oxidation or to a slower release of dissolved Pb(II) from the Pb(II) carbonate solid.

Kinetic models were developed to interpret the rates of cerussite and hydrocerussite oxidation under the experimental conditions. For cerussite and hydrocerussite oxidation with or

without initial PbO₂ (scrutinyite or plattnerite), the consumption of free chlorine included noncatalytic oxidation of Pb(II) and autocatalytic oxidation of Pb(II) with PbO₂ added or formed *in situ* (eq 1). The moles of PbO₂ formed were the same as the moles of free chlorine consumed (eq 2). For Pb(II) carbonate oxidation without any added PbO₂, the consumption of free chlorine and the formation of PbO₂ were described as simplified forms of eqs 1 and 2 with a value of zero for [Pb(IV)_{added}] which eliminated particular terms in the equations. The term [HOCl] indicates all free chlorine (both HOCl and OCl⁻). Good fits of the model to the data required different rate constants for Pb(II) oxidation catalyzed by PbO₂ formed *in situ* versus PbO₂ added as presynthesized solids. An initial modeling effort attempted to interpret the data using a single rate constant for PbO₂, but this provided much poorer fits (eqs S1–S4 and Figure S5). The kinetic constants k_1^* and k_2^* correspond to the noncatalytic and autocatalytic reaction proposed in other studies.^{8,16,45} The catalytic reaction rate constants (k_2^* , k_3^* , and k_4^*) were the value of apparent rate constants k_2 , k_3 , and k_4 multiplied by the amount of PbO₂ in the system (eqs 3 and 4). The added scrutinyite and plattnerite were assumed to have different catalytic reaction rate constants (k_3^* and k_4^* , respectively). Eqs 1–5 were employed for kinetic model development with scrutinyite, and eqs S5–S9 were used with plattnerite. The four unknown constants were estimated by finding the values that provided the optimal fit of five sets of experimentally measured chlorine consumption data (Figure 3 and Figure S6). The definition and unit of each constant are listed in Table 1. These constants in Table 1 are apparent rate

Table 1. Definitions and Units for Constants in Kinetic Model

constants	definitions	units
$^*[\text{HOCl}]_0$	initial free chlorine concentration	mM
Pb(II)_0	initial Pb(II) carbonate concentrations	mM
$\text{Pb(IV)}_{\text{add}}$	concentration of Pb(IV) solids added	mM
$\delta\text{-MnO}_2$	concentration of Mn(IV) formed in the batch experiments	mM
k_1^*	rate constant of homogeneous Pb(II) carbonate oxidation by chlorine	M ⁻¹ ·h ⁻¹
k_2^*	rate constant for heterogeneous Pb(II) carbonate oxidation by chlorine catalyzed by PbO ₂ generated <i>in situ</i>	M ⁻¹ ·h ⁻¹
k_3^*	rate constant for heterogeneous Pb(II) carbonate oxidation by chlorine catalyzed by scrutinyite	M ⁻¹ ·h ⁻¹
k_4^*	rate constant for heterogeneous Pb(II) carbonate oxidation by chlorine catalyzed by plattnerite	M ⁻¹ ·h ⁻¹
k_5^*	rate constant for heterogeneous Pb(II) carbonate oxidation by chlorine catalyzed by manganese oxide	M ⁻¹ ·h ⁻¹
k_2	apparent constant for heterogeneous Pb(II) carbonate oxidation by chlorine catalyzed by PbO ₂ generated <i>in situ</i>	M ⁻² ·h ⁻¹
k_3	apparent constant for heterogeneous Pb(II) carbonate oxidation by chlorine catalyzed by scrutinyite	M ⁻² ·h ⁻¹
k_4	apparent constant for heterogeneous Pb(II) carbonate oxidation by chlorine catalyzed by plattnerite	M ⁻² ·h ⁻¹
k_5	apparent constant for heterogeneous Pb(II) carbonate oxidation by chlorine catalyzed by manganese oxide	M ⁻² ·h ⁻¹

constants for oxidation reactions. The dissolution of lead carbonate solids and adsorption of Pb(II) onto PbO₂ or $\delta\text{-MnO}_2$ were not considered in the model development because they are much faster compared with the oxidation reactions so they can be ignored.

Table 2. Constants for Modeling the Dynamics of HClO during Pb(II) Carbonate Oxidation at pH 7.5

constants	cerussite	hydrocerussite	refs
k_1^*	$20.0 \pm 4.0 \text{ M}^{-1}\cdot\text{h}^{-1}$	$30.0 \pm 2.4 \text{ M}^{-1}\cdot\text{h}^{-1}$	our study
k_2	$(5.5 \pm 1.1) \times 10^5 \text{ M}^{-2}\cdot\text{h}^{-1}$	$(7.5 \pm 0.6) \times 10^5 \text{ M}^{-2}\cdot\text{h}^{-1}$	our study
k_3	$(6.0 \pm 1.3) \times 10^3 \text{ M}^{-2}\cdot\text{h}^{-1}$	$(6.0 \pm 1.0) \times 10^3 \text{ M}^{-2}\cdot\text{h}^{-1}$	our study
k_4	$(6.0 \pm 1.2) \times 10^3 \text{ M}^{-2}\cdot\text{h}^{-1}$	$(4.0 \pm 0.7) \times 10^3 \text{ M}^{-2}\cdot\text{h}^{-1}$	our study
k_5	$(3.0 \pm 0.3) \times 10^6 \text{ M}^{-2}\cdot\text{h}^{-1}$	$(5.5 \pm 0.6) \times 10^6 \text{ M}^{-2}\cdot\text{h}^{-1}$	our study

$$-\frac{d[\text{HOCl}]}{dt} = k_1^* \cdot [\text{Pb(II)}]_{\text{lead carbonate}} \cdot [\text{HOCl}] + k_2^* \cdot [\text{Pb(II)}]_{\text{lead carbonate}} \cdot [\text{HOCl}] + k_3^* \cdot [\text{Pb(II)}]_{\text{lead carbonate}} \cdot [\text{HOCl}] \quad (1)$$

$$\frac{d[\text{Pb(IV)}]}{dt} = k_1^* \cdot [\text{Pb(II)}]_{\text{lead carbonate}} \cdot [\text{HOCl}] + k_2^* \cdot [\text{Pb(II)}]_{\text{lead carbonate}} \cdot [\text{HOCl}] + k_3^* \cdot [\text{Pb(II)}]_{\text{lead carbonate}} \cdot [\text{HOCl}] \quad (2)$$

$$k_2^* = k_2 \cdot [\text{Pb(IV)}] \quad (3)$$

$$k_3^* = k_3 \cdot [\text{Pb(IV)}]_{\text{added}} \quad (4)$$

$$[\text{Pb(II)}]_{\text{lead carbonate}} = [\text{Pb(II)}]_0 - [\text{Pb(IV)}] \quad (5)$$

The constants that provided the best fit of the model output to the experimental results are compiled in Table 2. The simulated free chlorine concentration trends based on these reactions and apparent constants are shown together with the experimental data in Figure 3 (cerussite) and Figure S6 (hydrocerussite). The apparent constant for cerussite oxidation on PbO_2 formed *in situ* ($k_2 = 5.5 \times 10^5 \text{ M}^{-2}\cdot\text{h}^{-1}$) was 2 orders of magnitude higher than the constants for Pb(II) oxidation on the added PbO_2 ($k_3 = 6 \times 10^3 \text{ M}^{-2}\cdot\text{h}^{-1}$ and $k_4 = 6 \times 10^3 \text{ M}^{-2}\cdot\text{h}^{-1}$). The apparent constants for hydrocerussite followed the same trend. The oxidation rate constants $k_2^* - k_4^*$ were calculated by multiplying $k_2 - k_4$ by the concentration of PbO_2 present (eqs 3, 4, and S8). The rate constants used here are related to the moles of PbO_2 present, but for surface-based reactions, the controlling factor is probably the surface area of PbO_2 present. The higher specific surface area of PbO_2 formed *in situ* may be one reason for its more robust catalytic effect on Pb(II) carbonate oxidation than that of presynthesized PbO_2 . Unfortunately, we did not have a means of measuring the surface area of PbO_2 formed *in situ*. Comparing hydrocerussite and cerussite, hydrocerussite exhibited a higher autocatalytic oxidation speed than cerussite, and k_1^* and k_2 of hydrocerussite were both greater than these of cerussite. The reaction rate constants k_1^* and apparent rate constants k_3 and k_4 in Table 2 were almost 10 times higher than in the study by Liu et al.¹⁶ because the surface areas of cerussite and hydrocerussite (3.38 and 10.70 m^2/g) employed in this study were almost 10 times higher than Liu et al.¹⁶ (0.42 and 0.81 m^2/g).

The effect of pH on cerussite oxidation was examined (Figure 4). The duration of the lag stage was not affected by increasing pH for cerussite. Free chlorine consumption speeds were almost the same at pH 6.5, 7.5, and 8.5 for cerussite. For hydrocerussite oxidation, the free chlorine consumption speed decreased with increasing pH (Figure S7). Similar results were observed by Zhang and Lin.⁴² The influence of pH on Pb(II) carbonate oxidation by free chlorine could be related to two phenomena. First, the pH determines the ratio of HOCl/

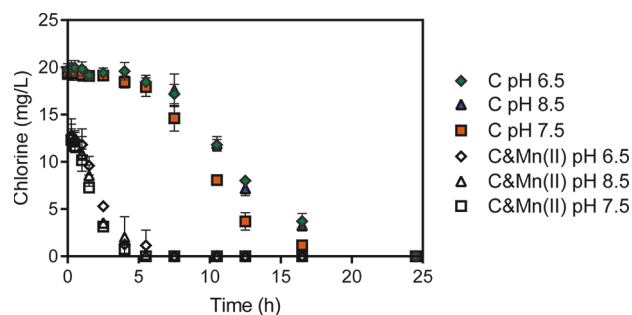


Figure 4. Effect of pH on cerussite oxidation. Condition: initial free chlorine = 0.31 mM (22 mg/L) as Cl_2 , initial Pb = 2 mM, carbonate = 0.2 mM as C. Experimental results were shown as points.

OCl^- , and for many species, oxidation rates are faster with HOCl than with OCl^- .⁴⁶ Second, the pH determines the species of dissolved Pb(II), which also affects the oxidation rates according to Figure 3. Prior research demonstrated that high DIC concentration could inhibit Pb(II) carbonate oxidation by free chlorine via formation of dissolved Pb(II) carbonate complexes, which are much less reactive compared with lead hydroxo complexes.^{8,16,44}

The Catalytic Effect of Mn Oxide on Lead Carbonate Oxidation. The addition of Mn(II) significantly enhanced the rate of cerussite oxidation (Figure 5). With free chlorine Mn(II), oxidation was much faster than Pb(II) oxidation, and it produced $\delta\text{-MnO}_2$ (Figure S2). The free chlorine

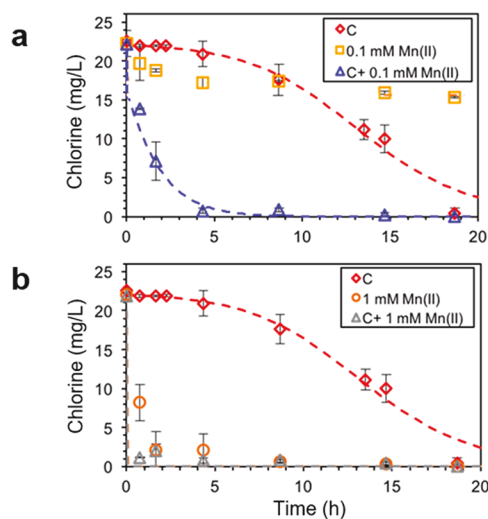


Figure 5. Oxidation of cerussite (C) by chlorine with (a) 0.1 and (b) 1 mM Mn(II) concentrations. Conditions: pH = 7.5 ± 0.2 , initial free chlorine = 0.31 mM (22 mg/L as Cl_2), initial Pb = 2 mM, carbonate = 0.2 mM as C. Experimental results are shown as points, and modeling results are shown as dashed lines. The average values from duplicate experiments are shown, and the error bars represent the standard deviation from the duplicate experiments.

concentration was in excess of the amount required to oxidize 0.1 mM Mn(II), so after Mn(II) was completely oxidized, there was still sufficient free chlorine to oxidize Pb(II) carbonate. The rate of Pb(II) carbonate oxidation was accelerated as the rate of free chlorine consumption was much higher than with the lead carbonate oxidation in the absence of Mn(II). Hydrocerussite oxidation with the presence of Mn(II) showed the same trends as cerussite (Figure S9).

A kinetic model for free chlorine concentration during Pb(II) oxidation in systems to which Mn(II) had been added was developed (eqs 6 and 7). Because the rate of Mn(II) oxidation was at least 3 orders of magnitude higher than the rates of other reactions used in the model, we assumed that all of the Mn(II) was instantaneously oxidized to δ -MnO₂; efforts to account for the kinetics of Mn(II) oxidation made the model more complicated but did not improve the ability to simulate Pb(II) oxidation. This assumption was confirmed by the similar catalytic effects on Pb(II) carbonate oxidation of δ -MnO₂ solids formed *in situ* and added (Figure 5, S8, and S9). In developing the model of Pb(II) oxidation in the presence of Mn oxide, the constants for noncatalytic Pb(II) oxidation and for oxidation catalyzed by PbO₂ were set at the values determined from the previously discussed experiments in which no Mn had been added. The k_5 value was determined by fitting the model output to experimental data of chlorine consumption in the presence of 0.1 mM Mn(II). The value obtained then can be successfully applied to the system with 1 mM Mn(II) (Figure 5). The definition and units of each constant are listed in Table 1. The actual second order oxidation rate constants were determined using apparent constants as noted in eq 7.

$$\frac{d[\text{Pb(IV)}]}{dt} = k_1^* [\text{Pb(II)}]_{\text{lead carbonate}} [\text{HOCl}] + k_2^* [\text{Pb(IV)}] \cdot [\text{Pb(II)}]_{\text{lead carbonate}} [\text{HOCl}] + k_5^* [\text{Pb(II)}]_{\text{lead carbonate}} [\text{HOCl}] \quad (6)$$

$$k_5^* = k_5 \cdot [\delta - \text{MnO}_2] \quad (7)$$

Based on our modeling results (Table 2), the apparent constants for Pb(II) oxidation with δ -MnO₂ were 1 order of magnitude higher than those of PbO₂ formed *in situ*. Considering the amount of δ -MnO₂ in the system, the actual catalytic rate constant with δ -MnO₂ could be at least 3 orders of magnitude higher than that of PbO₂ formed *in situ* and PbO₂ added. Apart from the catalytic effect of Mn(II) on Pb(II) carbonate oxidation, we also observed faster chlorine consumption rates for Mn(II) when Pb(II) carbonate was present than when Mn(II) was present on its own, which may indicate that the surface of Pb(II) carbonate helped catalyze the oxidation of Mn(II) by chlorine.

Products of Pb(II) Carbonate Oxidation. The reaction of cerussite and hydrocerussite with free chlorine produced PbO₂. Scrutinyte was formed from lead carbonate oxidation as determined by XRD (Figure 6). The effect of Mn(II) on lead carbonate oxidation by free chlorine was further confirmed by XRD patterns. The structural changes of Pb(II) carbonate solids and the formation of PbO₂ were also observed by SEM results (Figure S10). With 1 mM Mn(II), there was no change in the cerussite XRD pattern because all the free chlorine was consumed by Mn(II) oxidation. For cerussite oxidation by free chlorine, the XRD patterns for 5 h samples were almost the same as those of the initial cerussite, which was consistent with batch experiment results that the reactions were in a lag stage

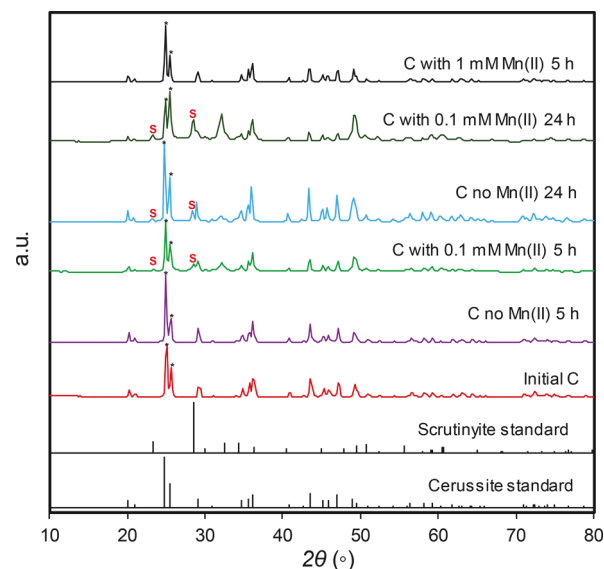


Figure 6. X-ray diffraction patterns of solids from cerussite (C) oxidation experiments. Dominant peaks associated with scrutinyte and cerussite were indicated, and reference powder diffraction file patterns are included for these phases (04-008-7674 and 01-073-4362). The main peaks of scrutinyte are indicated as S. The main peaks of cerussite are indicated as *.

for the first 5 h. After 24 h reaction with free chlorine, peaks for scrutinyte at 2θ of 23° and 28° were observed for the reacted cerussite. Liu et al.¹⁶ examined cerussite and hydrocerussite oxidation by 50 mg/L free chlorine and observed the signature peaks for scrutinyte at a longer oxidation time, which was probably because of the higher DIC concentration (0.001 M) used in their experiments.

With the presence of Mn(II), the same peaks for scrutinyte were observed after cerussite reaction with free chlorine for both 5 and 24 h. These XRD results were consistent with the observed free chlorine consumption profiles from batch experiments. As shown in Figure 6, scrutinyte was the only product of cerussite oxidation by free chlorine. The same overall trend was observed for hydrocerussite oxidation (Figure S11); with the addition of 0.1 mM Mn(II), scrutinyte was observed in the solid sample for 5 and 24 h reaction with free chlorine. The formation of pure scrutinyte from hydrocerussite oxidation was also observed in the studies by Wang et al.⁸ and Liu et al.¹⁶ When starting with cerussite and no DIC, Wang et al.⁸ observed cerussite transformed to hydrocerussite after 1 day, and a mixture of scrutinyte and plattnerite formed after 28 days. Nevertheless, in both the work of Liu et al.¹⁶ and our research, plattnerite was not observed after cerussite oxidation. The difference in these studies may be due to the difference in experimental pH and carbonate concentration. Higher pH and higher carbonate concentration can lead to higher generation of scrutinyte over plattnerite in PbO₂ formation. We also note that there are two possible processes for the oxidation of lead carbonate solids to PbO₂, dissolution-oxidation-precipitation (DOR), and direct solid-state oxidation.^{8,16,47} In this study, DOR is considered to be the main PbO₂ formation pathways. Although there might be direct solid-state oxidation of a Pb(II) layer at the surface of lead carbonate solids, this process will yield a protective layer of PbO₂ on lead carbonate solid surface and inhibit further oxidation. Nevertheless, thorough oxidation of Pb(II) carbo-

nate solids to PbO_2 was observed by previous studies,^{8,48} which indicates that DOR is more significant.

Flow-through Experiment. In the batch experiments, a high free chlorine concentration (0.28 ± 0.03 mM as Cl_2) was applied initially and then depleted as it oxidized Pb(II) and Mn(II) species. In contrast to the batch experiments, flow-through experiments were conducted with a more realistic free chlorine concentration (0.034 mM or 2.4 mg/L as Cl_2) that was continuously supplied. The oxidation of cerussite was enhanced with $\delta\text{-MnO}_2$ in the flow-through experiment (Figure 7). After a period of relatively high effluent

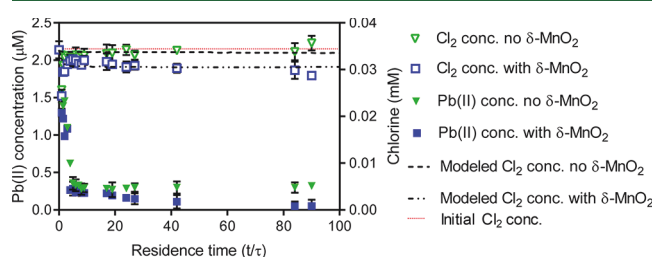


Figure 7. Effluent lead and chlorine concentrations from flow-through experiments with cerussite in the absence and presence of $\delta\text{-MnO}_2$. Experiments were conducted at a hydraulic residence time of 20 min. Samples were not taken during the middle of the experiments (50–80 τ). Modeling results are shown as dashed lines. Conditions: $\text{pH} = 7.5 \pm 0.2$, initial free chlorine = 0.034 mM (2.4 mg/L) as Cl_2 , initial $\text{Pb} = 2$ mM, carbonate = 0.2 mM as C.

concentrations (~ 1.5 μM) of Pb(II) as the CSTRs started up, the concentrations of Pb(II) dropped to much lower values (0.05–0.32 μM). The effluent concentrations of Pb(II) were lower with $\delta\text{-MnO}_2$ (0.05 μM) than without (0.32 μM). For the system without $\delta\text{-MnO}_2$, the concentrations of Pb(II) stabilized at a steady-state concentration. A steady-state was not necessarily expected since the rate of Pb(II) consumption would be expected to increase as more PbO_2 accumulated in the reactor; however, the rate of PbO_2 accumulation may have been slow enough for the system to be at steady-state. For the system with $\delta\text{-MnO}_2$, a steady state was not reached, and Pb(II) concentrations decreased over the duration of the experiment, which indicates that the amount of PbO_2 in the system was probably increasing with time. The faster oxidation of cerussite with $\delta\text{-MnO}_2$ was also confirmed by higher chlorine consumption in the effluent. When similar experiments were conducted with hydrocerussite (Figure S12), we observed the similar trend that $\delta\text{-MnO}_2$ accelerates hydrocerussite oxidation.

For model development, we considered the conditions with and without $\delta\text{-MnO}_2$ (eqs 8 and 9). Q and V are the flow rate and volume, respectively. The amount of Pb(II) (≤ 1.2 μmol) in the total outlet was neglected in the model development because it was negligible when compared with the amount of Pb(II) and Pb(IV) solids (≥ 38.8 μmol) left in the CSTRs after the experiment. For cerussite oxidation without $\delta\text{-MnO}_2$, the $\delta\text{-MnO}_2$ concentration was set to zero in eqs 8 and 9 which eliminated particular terms in the equations. The constants were defined in Table 1, and rate constants (k_1^* , k_2^* , and k_5^*) were obtained by using the same rate constants obtained from batch experiments (Table 2). The application of parameters from batch experiments to flow-through experiment was based on the hypothesis that the behavior of solids in batch experiment was consistent with the behavior of solids in the

flow-through experiments, which was confirmed by the overall agreement between experimental and modeled chlorine concentrations in the effluent. Chlorine concentration profiles in the effluent derived from modeling work are shown in Figure 7 and Figure S11 as dashed lines. The simulations provided general agreement with the flow-through experimental data that the presence of $\delta\text{-MnO}_2$ accelerates chlorine consumption, which indicates a higher rate of Pb(II) oxidation.

$$\frac{d[\text{HOCl}]}{dt} \cdot V = [\text{HOCl}]_{\text{inlet}} \cdot Q - \{k_1^* \cdot [\text{Pb(II)}]_{\text{lead carbonate}} \cdot [\text{HOCl}] - k_2^* \cdot [\text{Pb(II)}]_{\text{lead carbonate}} \cdot [\text{HOCl}] - k_5^* \cdot [\text{Pb(II)}]_{\text{lead carbonate}} \cdot [\text{HOCl}]\} \cdot V - [\text{HOCl}] \cdot Q \quad (8)$$

$$\frac{d[\text{Pb(IV)}]}{dt} = k_1^* \cdot [\text{Pb(II)}]_{\text{lead carbonate}} \cdot [\text{HOCl}] + k_2^* \cdot [\text{Pb(II)}]_{\text{lead carbonate}} \cdot [\text{HOCl}] + k_5^* \cdot [\text{Pb(II)}]_{\text{lead carbonate}} \cdot [\text{HOCl}] \quad (9)$$

Environmental Implications. PbO_2 solids can maintain low dissolved Pb(II) concentrations, but they are not always observed on LSLs that deliver drinking water with free chlorine as the disinfectant. Water chemistry can strongly influence the oxidation of Pb(II) carbonate in the actual distribution system. Under a more realistic and continuously supplied free chlorine concentration (0.034 mM or 2.4 mg/L as Cl_2), $\delta\text{-MnO}_2$ also enhanced Pb(II) carbonate oxidation.

The effect of Mn on Pb(II) carbonate oxidation can be successfully explained by the $\delta\text{-MnO}_2$ that formed *in situ* acting as a heterogeneous catalyst for Pb(II) oxidation. Modeling work showed that the catalytic effect of $\delta\text{-MnO}_2$ formed *in situ* is almost 3 orders of magnitude faster than that of an equal amount of PbO_2 formed *in situ*. Although results in this study were obtained from batch and flow-through experiments, it can be extrapolated to LSLs. In a pipe reactor study by Bae et al.,⁴⁹ an accidental dosage of Mn(II) was found to skyrocket the free chlorine consumption speed and decrease the Pb(II) concentration from ~ 50 $\mu\text{g/L}$ to less than 10 $\mu\text{g/L}$ in the effluent of lead pipes. Our experimental and modeling results can help explain the phenomenon of Pb(II) decreasing in the effluent of lead pipes by PbO_2 formation catalyzed by *in situ* formation of MnO_2 . We anticipate that the study of water chemistry in bench-scale experiments can be employed to study the Pb corrosion in drinking water distribution systems with different levels of Mn in the water. The same findings may potentially be extended to other metal oxide deposits on the inner surface of LSLs, such as those of iron and copper. Nevertheless, the catalytic effect of these metal oxides may not be as robust as $\delta\text{-MnO}_2$, which has a relative high surface area, fast Pb(II) adsorption kinetics, and high Pb(II) adsorption capacity.⁵⁰ Furthermore, our findings likely have broader implications for the role of deposits (such as Mn, Fe, and Cu oxide) in drinking water distribution systems. Previous studies have explored the role of these deposits as sinks for trace inorganic contaminants (arsenic, selenium, vanadium, etc.) via the sorption process.^{11,14,51,52} Here, our findings reveal another possible role of these deposits in mediating redox transformations of trace inorganic contaminants such as Pb(II) .

■ ASSOCIATED CONTENT

Supporting Information

The Supporting Information is available free of charge on the ACS Publications website at DOI: 10.1021/acs.est.8b07356.

Table S1, initial model development for Pb(II) carbonate oxidation with added PbO₂; Figures S1–S12, standard reduction potentials of chemicals in experiment, XRD patterns for synthesized scrutinyite and solids obtained from Mn(II) oxidation, flow-through experiment setup, chlorine concentration with PbO₂, free chlorine concentration profiles for hydrocerussite oxidation with and without PbO₂, effect of pH on hydrocerussite oxidation by chlorine, oxidation of Pb(II) carbonate by chlorine with and without 0.1 mM δ-MnO₂ solids, oxidation of hydrocerussite by chlorine with different Mn(II) concentrations, SEM images of Pb(II) carbonate before and after reactions, XRD patterns for of solids from hydrocerussite oxidation experiments, and results of flow-through experiments with hydrocerussite (PDF)

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