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# Electron transfer between neptunium and sodium chlorite in acidic chloride media†

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Controlling aqueous 5f-element electron transfer chemistry is critical for processing efforts associated with actinide technologies. Often, redox agents are added during actinide processing steps to control actinide redox chemistry and manipulate the actinide oxidation states for the separation. Sodium chlorite, NaClO<sub>2(aq)</sub>, represents one of these useful redox agents. For example, NaClO<sub>2(aq)</sub> finds widespread application in the processing of plutonium and americium. Surprisingly, however, redox reactivity between NaClO<sub>2(aq)</sub> and other actinides, like neptunium, has been largely ignored. That knowledge gap is addressed herein. We characterized some redox reactivity between NaClO<sub>2(aq)</sub> and Np<sup>4+(aq)</sup> and identified experimental conditions that held neptunium in the +4 oxidation state or converted Np<sup>4+(aq)</sup> to NpO<sub>2<sup>2+(aq)</sup> or NpO<sub>2<sup>1+(aq)</sup>. This was achieved by carefully adjusting four variables: ingoing concentrations of (1) Np<sup>4+(aq)</sup>, (2) NaClO<sub>2(aq)</sub>, (3) Cl<sup>1-(aq)</sup>, and (4) H<sup>1+(aq)</sup>. We discovered that three neptunium oxidation states (+4, +5, and +6) could be accessed using one ubiquitous redox agent, NaClO<sub>2(aq)</sub>. These results highlight the diverse electron transfer chemistry available to neptunium in aqueous solutions, provide new insight on how neptunium reacts with NaClO<sub>2(aq)</sub>, and are discussed within the context of their importance to plutonium and americium processing.</sub></sub>

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## Introduction

Controlling electron transfer chemistry accessible to the 5f-elements in aqueous media is critical for nearly every actinide-based technological application. These applications span from isotope production, environmental restoration, nuclear fuel rod fabrication, space exploration, and harvesting actinides from weapons program waste streams.<sup>1–19</sup> The latter topic was the major motivating factor for this particular research effort. For instance, at Los Alamos National Laboratory (LANL), transuranic actinides – like plutonium (Pu) and americium (Am) – are recovered from aqueous waste streams.<sup>20–23</sup> Success associated with this waste stewardship is valuable and responsible. It reduces burden on taxpayers by lowering transuranic waste costs, decreases the amount of transuranic waste generated, and recycles valuable and rare radioisotopes (e.g., <sup>238</sup>Pu, <sup>239</sup>Pu, <sup>240</sup>Pu, <sup>241</sup>Pu, <sup>242</sup>Pu, and <sup>241</sup>Am) for national security and industrial usage.

To recover plutonium and americium from aqueous waste streams, LANL implemented the experimental chloride extraction line (EXCEL) and chloride extraction and recovery (CLEAR)

aqueous processing methods.<sup>15,24–28</sup> In terms of plutonium recovery, waste entering the processing line contains plutonium in a variety of oxidation states. Hence, successful plutonium recovery requires a valence state adjustment that holds plutonium in the +4 oxidation state during subsequent chemical processing steps. This plutonium oxidation-state adjustment can be carried out using sodium chlorite; NaClO<sub>2(aq)</sub>. The robustness associated with using NaClO<sub>2(aq)</sub> as a plutonium valence adjusting agent has been documented by the successful recovery of plutonium from diverse waste streams using EXCEL for decades.<sup>16,29</sup>

Accompanying changes to the national nuclear agenda, described in the Nuclear Posture Review, are new chemical challenges that face aqueous recovery and recycling of plutonium and americium.<sup>30–32</sup> One obstacle is associated with needs to accommodate increasingly diverse waste feedstocks. Processing these feedstocks is complicated because they contain a wide range of chemical constituents. In addition, their chemical identities and quantities change with time. One relevant example is neptunium-237: <sup>237</sup>Np, half-life (*t*<sub>1/2</sub>) = 2.144(7) × 10<sup>6</sup> y.<sup>33</sup> Concerns regarding <sup>237</sup>Np contamination stem from substantial <sup>237</sup>Np ingrowth in aged plutonium/americium containing waste. Fig. 1 qualifies these concerns. It shows that the <sup>237</sup>Np contaminant is generated *via* α-decay from its <sup>241</sup>Am [*t*<sub>1/2</sub> = 432.6(6) y] parent radionuclide, which in turn is the β<sup>–</sup> decay product from <sup>241</sup>Pu [*t*<sub>1/2</sub> = 14.329(29) y].<sup>33</sup> It also highlights how “aged” waste will contain substantial

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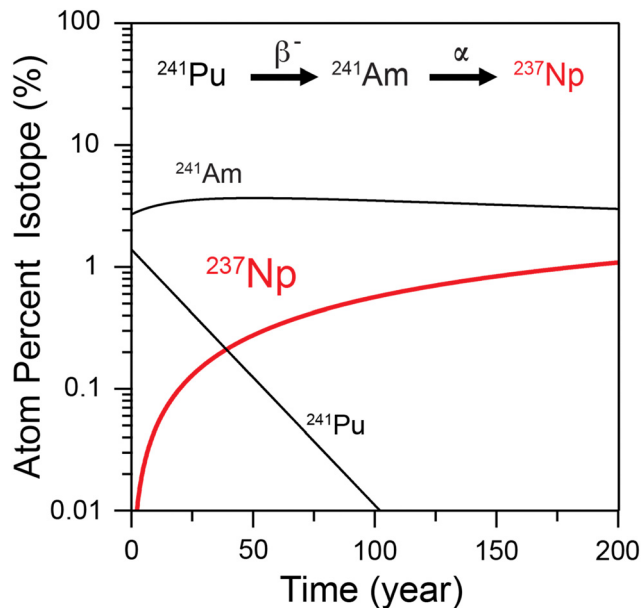



Fig. 1 Calculated isotopic decay products for selected transuranic radioisotopes within aged plutonium calculated from aged plutonium isotopic content (WGPu-C, 16–19%  $^{240}\text{Pu}$  as defined in the compendium of material composition data for radiation transport modeling).<sup>39</sup>

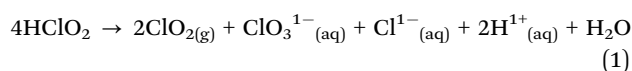
quantities of  $^{237}\text{Np}$ .<sup>34–38</sup> Unfortunately, it is unknown how  $^{237}\text{Np}_{(\text{aq})}$  will respond to the antecedently described plutonium valence adjustment step because redox chemistry between neptunium and  $\text{NaClO}_{2(\text{aq})}$  has not been studied in depth. Defining electron transfer chemistry between neptunium and  $\text{NaClO}_{2(\text{aq})}$  in aqueous solutions is relevant to plutonium/amerium processing. That information will show how  $^{237}\text{Np}$  contaminants move through the EXCEL and CLEAR processing lines and provide insight for maintaining robustness and effectiveness in plutonium/amerium waste processing efforts.

To address the aforementioned  $^{237}\text{Np}$  processing concerns, we set out to characterize some redox chemistry between  $^{237}\text{Np}_{(\text{aq})}$  and the  $\text{NaClO}_{2(\text{aq})}$ . We discovered that we could force neptunium into three different oxidation states (+4, +5, +6) using the  $\text{NaClO}_{2(\text{aq})}$  redox agent by manipulating four experimental variables. Those variables were ingoing concentrations of  $\text{Np}^{4+}_{(\text{aq})}$ ,  $\text{NaClO}_{2(\text{aq})}$ ,  $\text{Cl}^{-}_{(\text{aq})}$ , and  $\text{H}^{1+}_{(\text{aq})}$ . Although  $\alpha$ -particle radiation has been known to influence neptunium oxidation state chemistry, these attributes were not considered.<sup>40</sup> Scheme 1 and Table 1 showcase that  $\text{NaClO}_{2(\text{aq})}$  acted as a  $\text{Np}^{4+}_{(\text{aq})}$  valence holding agent under a wide range of experimental conditions. We also found experimental conditions that converted  $\text{Np}^{4+}_{(\text{aq})}$  to the  $\text{NpO}_2^{1+}_{(\text{aq})}$  mono-cation or the  $\text{NpO}_2^{2+}_{(\text{aq})}$  di-cation. Stable solutions of  $\text{NpO}_2^{1+}_{(\text{aq})}$  were obtained when the ingoing concentrations of  $\text{Np}^{4+}_{(\text{aq})}$  (0.81 mM),  $\text{NaClO}_{2(\text{aq})}$  (3.4 mM),  $\text{Cl}^{-}_{(\text{aq})}$  (0.026 M), and  $\text{H}^{1+}_{(\text{aq})}$  (0.02 M) were all relatively low. Stable solutions of  $\text{NpO}_2^{2+}_{(\text{aq})}$  were obtained when the ingoing concentration of  $\text{Np}^{4+}_{(\text{aq})}$  was high (3.1 mM to 1.5 mM),  $\text{NaClO}_{2(\text{aq})}$  was high (160 to 32 mM),  $\text{Cl}^{-}_{(\text{aq})}$  ranged 0.02 to 9.2 M, and  $\text{H}^{1+}_{(\text{aq})}$  was between 0.02 and 5.4 M. Overall, these results demonstrated that that reactivity between  $\text{Np}^{4+}_{(\text{aq})}$  and  $\text{NaClO}_{2(\text{aq})}$  was diverse,

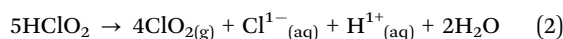
predictable, and therefore controllable. That insight advances understanding about neptunium electron transfer reactions under experimental conditions that are relevant to those described above during EXCEL and CLEAR processing of plutonium and  $^{241}\text{Am}$ .

Before discussing reactivity between  $\text{Np}^{4+}_{(\text{aq})}$  and  $\text{NaClO}_{2(\text{aq})}$ , we found it instructive to comment – at a high level – on the aqueous reactivity of  $\text{NaClO}_{2(\text{aq})}$  in  $\text{Cl}^{-}_{(\text{aq})}$  containing solutions. The chemistry of  $\text{NaClO}_{2(\text{aq})}$  has been the subject of numerous experimental studies described previously.<sup>41–47</sup> The  $\text{NaClO}_2$  salt has been structurally characterized.<sup>48–50</sup> The  $\text{ClO}_2^{1-}$  anion contains chlorine in the +3 oxidation state, and (based on the standard half-cell potentials) is the strongest oxidizer in the chlorine oxyanion family.<sup>45</sup> As testament, many salts containing the  $\text{ClO}_2^{1-}$  anion are reported to decompose explosively when exposed to heat and shock, e.g. salts of  $\text{Ag}^{1+}$ ,  $\text{Hg}^{1+}$ ,  $\text{Tl}^{1+}$ ,  $\text{Pb}^{2+}$ ,  $\text{Cu}^{2+}$ , and  $\text{NH}_4^{1+}$ .<sup>51</sup> Hence, inorganic  $\text{ClO}_2^{1-}$  salts are often regarded as being unstable and highly reactive. In aqueous solutions, the conjugate acid of  $\text{ClO}_2^{1-}$  (chlorous acid;  $\text{HClO}_2$ ) is a weak acid ( $\text{p}K_a \sim 1.94$ ),<sup>52</sup> one of the most reactive (least stable) oxoacids of chlorine,<sup>53</sup> and has only been observed in aqueous solution at low concentrations.<sup>54</sup> In aqueous chloride solutions, the chemistry of the  $\text{ClO}_2^{1-}_{(\text{aq})}/\text{HClO}_{2(\text{aq})}$  pair is dominated by disproportionation reactions whose product ratios are heavily influenced by the  $\text{H}^{1+}_{(\text{aq})}$  and  $\text{Cl}^{-}_{(\text{aq})}$  concentrations associated with the aqueous matrix.<sup>47,55–57</sup> For example, in aqueous matrixes and in the absence of  $\text{Cl}^{-}_{(\text{aq})}$ , the  $\text{ClO}_2^{1-}_{(\text{aq})}$  anion is reported to disproportionate and form gaseous chlorine dioxide [ $\text{Cl}^{\text{IV}}\text{O}_{2(\text{g})}$ ], chlorate [ $\text{Cl}^{\text{V}}\text{O}_3^{1-}_{(\text{aq})}$ ], and chloride [ $\text{Cl}^{-}_{(\text{aq})}$ ] (eqn (1)). In contrast, when  $\text{Cl}^{-}_{(\text{aq})}$  concentrations are high (see ref. 58 for details), the disproportionation reaction favors formation of the  $\text{ClO}_{2(\text{g})}$  and  $\text{Cl}^{-}_{(\text{aq})}$  pair (eqn (2)) and the formation of  $\text{ClO}_3^{1-}_{(\text{aq})}$  is suppressed. According to Kieffer and Gordon, the rate of  $\text{ClO}_2^{1-}_{(\text{aq})}$  disproportionation also varies with  $\text{Cl}^{-}_{(\text{aq})}$  content. For example the disproportionation reaction rate increases substantially when moving from low (no ingoing)  $\text{Cl}^{-}_{(\text{aq})}$  concentration ( $t_{1/2} = 389 \pm 3$  min without added  $\text{Cl}^{-}_{(\text{aq})}$ ) to higher (100 mM)  $\text{Cl}^{-}_{(\text{aq})}$  concentrations ( $t_{1/2} = 6.85 \pm 0.05$  min).<sup>56,57</sup> These disproportionation rates are not exact, and likely represent a combination of multiple chemical events and intertwined reactions.<sup>58</sup>

$\text{ClO}_2^{1-}_{(\text{aq})}$  disproportionation in low  $\text{Cl}^{-}_{(\text{aq})}$  conditions



$\text{ClO}_2^{1-}_{(\text{aq})}$  disproportionation in high  $\text{Cl}^{-}_{(\text{aq})}$  conditions

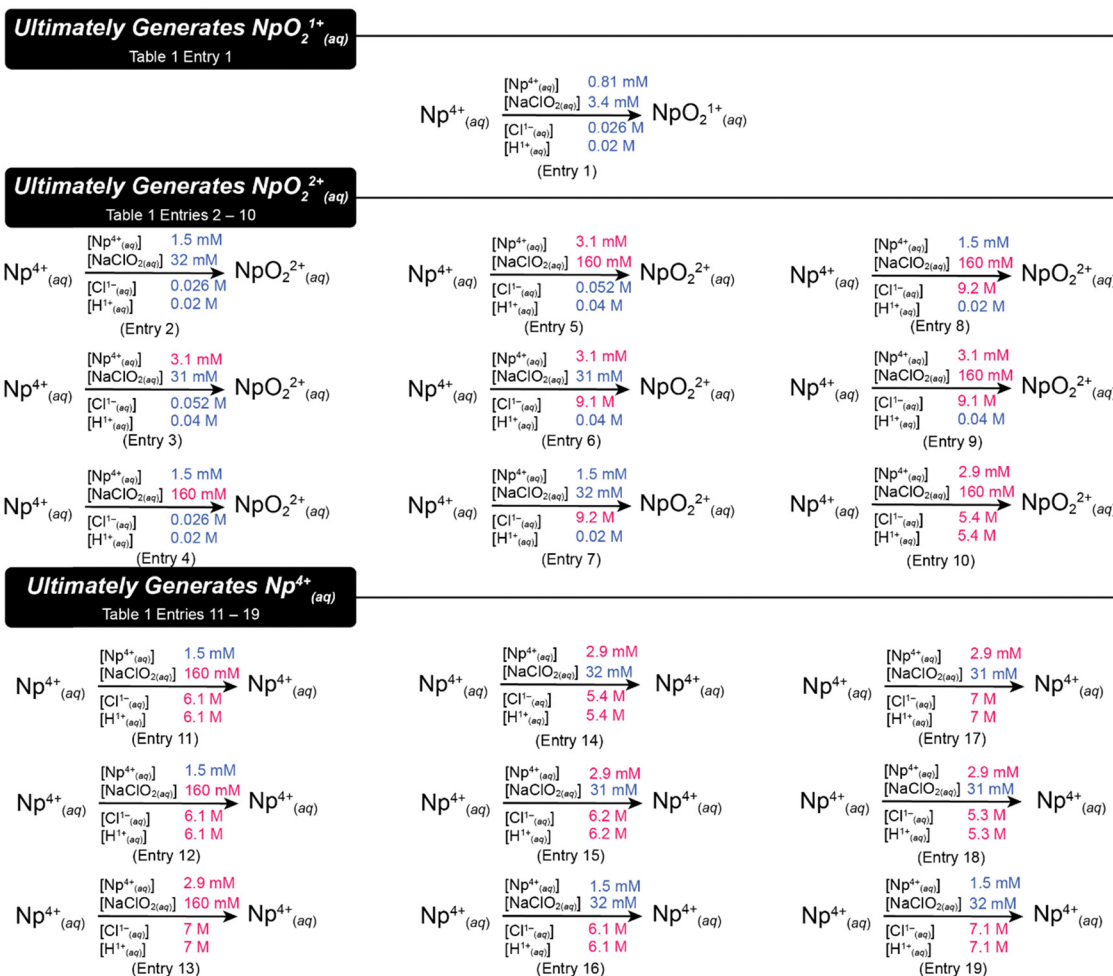


## Results and discussion

### Generating oxidation state pure solutions of $\text{NpO}_2^{1+}_{(\text{aq})}$ and $\text{NpO}_2^{2+}_{(\text{aq})}$ with $\text{NaClO}_{2(\text{aq})}$

We discovered experimental conditions that selectively oxidized  $\text{Np}^{4+}_{(\text{aq})}$  to either  $\text{NpO}_2^{1+}_{(\text{aq})}$  or  $\text{NpO}_2^{2+}_{(\text{aq})}$  in aqueous acidic solutions of  $\text{HCl}_{(\text{aq})}$  (0.02 to 5.4 M; entries 1 to 10 in Table 1)





**Scheme 1** General  $\text{Np}^{4+}_{(aq)}$  reaction products under various experimental conditions. The “entry” showed in parentheses references the experimental conditions documented in Table 1. Colors associated with concentrations are depicted as relatively high or low.

without the exclusion of air. To carry out these studies, we generated an oxidation state and chemically pure aqueous stock solution of  $\text{Np}^{4+}_{(aq)}$  (66 mM) using a previously described method.<sup>59</sup> Aliquots from this  $\text{Np}^{4+}_{(aq)}$  stock solution were combined with the  $\text{NaClO}_{2(aq)}$  oxidant at fixed  $\text{Cl}^-_{(aq)}$  concentrations. Transformations that converted  $\text{Np}^{4+}_{(aq)}$  to either  $\text{NpO}_2^{1+}_{(aq)}$  and/or  $\text{NpO}_2^{2+}_{(aq)}$  were subsequently monitored using ultraviolet-visible (UV-vis) absorption spectroscopy. Scheme 1 and Table 1 provide a high-level summary of how the matrix compositions impacted product formation.

We found that exclusive formation of  $\text{NpO}_2^{1+}_{(aq)}$  occurred when ingoing concentrations for the  $^{237}\text{Np}_{(aq)}$  analyte (0.81 mM),  $\text{NaClO}_{2(aq)}$  redox agent (3.4 mM),  $\text{Cl}^-_{(aq)}$  complexing agent (0.026 M), and  $\text{H}^+_{(aq)}$  (0.02 M) were all low, relative to the other experimental conditions we examined (entry 1 in Table 1). As documented in Fig. 2, addition of  $\text{NaClO}_{2(aq)}$  to a solution of  $\text{Np}^{4+}_{(aq)}$  caused the UV-vis absorption peaks from  $\text{Np}^{4+}_{(aq)}$  at  $10\,500\text{ cm}^{-1}$  (960 nm) and  $8\,700\text{ cm}^{-1}$  (1150 nm) to vanish within 1 day (see Fig. 3). Concomitantly, new peaks from  $\text{NpO}_2^{1+}_{(aq)}$  at  $10\,200\text{ cm}^{-1}$  (980 nm) and  $9\,100\text{ cm}^{-1}$  (1100 nm) emerged. Fig. 2 and 3 additionally documented that this

$\text{NpO}_2^{1+}_{(aq)}$  product was stable during a 5-day monitoring process, as no other neptunium oxidation states were detected by UV-vis spectroscopy. To obtain additional insight into the transformation of  $\text{Np}^{4+}_{(aq)}$  to  $\text{NpO}_2^{1+}_{(aq)}$ , we characterized the rate at which absorption peaks from  $\text{Np}^{4+}_{(aq)}$  decreased in intensity and peaks associated with  $\text{NpO}_2^{1+}_{(aq)}$  were prevalent for 5 days (Fig. 3). The reaction rate laws were calculated, and subsequent analyses suggested that oxidation of  $\text{Np}^{4+}_{(aq)}$  to  $\text{NpO}_2^{1+}_{(aq)}$  proceeded by fractional order dependence on both the  $\text{Np}^{4+}_{(aq)}$  and  $\text{ClO}_2^{1-}_{(aq)}$  reagents,  $0.300 \pm 0.008$  and  $0.110 \pm 0.002$ , respectively. Fractional order reactions can be indicative of chain reactions propagated by radical species present in solution and many radical reactions pathways could be responsible for the  $\text{NaClO}_{2(aq)}$  initiated conversion of  $\text{Np}^{4+}_{(aq)}$  to  $\text{NpO}_2^{1+}_{(aq)}$ .<sup>41,60–63</sup>

Subtle changes in the aqueous matrix identity substantially impacted the  $\text{Np}^{4+}_{(aq)}$  oxidation process. As a representative example, data displayed in Fig. 4 documented how oxidizing  $\text{Np}^{4+}_{(aq)}$  to  $\text{NpO}_2^{1+}_{(aq)}$  was dependent on the ingoing  $\text{NaClO}_{2(aq)}$  quantities. To generate these data, we sequentially added aliquots of  $\text{NaClO}_{2(aq)}$  (0.072  $\mu\text{mol}$ ; 10  $\mu\text{L}$  of a 7.2 mM stock solution) to an



**Table 1** Experimental conditions associated with all reactions between  $\text{Np}^{4+}_{(\text{aq})}$  and  $\text{NaClO}_{2(\text{aq})}$  described in this paper. Ingoing concentrations of  $\text{Np}^{4+}_{(\text{aq})}$ ,  $\text{NaClO}_{2(\text{aq})}$ ,  $\text{H}^{1+}_{(\text{aq})}$ , and  $\text{Cl}^{1-}_{(\text{aq})}$  are recorded. Also shown is the dominant neptunium species that forms at two different time points (after 1 vs. 24 h). Note, ingoing  $\text{Cl}^{1-}_{(\text{aq})}$  concentration includes three sources: from  $\text{HCl}_{(\text{aq})}$ , from  $\text{LiCl}$ , and from the ingoing  $\text{Np}^{4+}_{(\text{aq})}$  stock solution. Values in the "Ingoing  $\text{Cl}^{1-}$  Total (M)" concentration column represent total ingoing  $\text{Cl}^{1-}$  concentration (top number), the contributions from  $\text{HCl}_{(\text{aq})}$  and  $\text{LiCl}_{(\text{aq})}$  (bottom numbers in parentheses), and contributions from the  $\text{Np}^{4+}_{(\text{aq})}$  stock solution

Entry	Dominant species ( $t = 1$ h)	Dominant species ( $t = 24$ h)	Initial $\text{Np}^{4+}_{(\text{aq})}$ (mM)	Initial $\text{NaClO}_{2(\text{aq})}$ (mM)	Initial $\text{H}^{1+}_{(\text{aq})}$ (M)	Ingoing $\text{Cl}^{1-}$ total <sup>a</sup> (M)
						( $\text{Cl}^{1-}$ from $\text{HCl}$ ) + ( $\text{Cl}^{1-}$ from $\text{LiCl}$ ) + ( $\text{Cl}^{1-}$ from $\text{NpCl}_4$ )
1	$\text{NpO}_2^{1+}_{(\text{aq})}$	$\text{NpO}_2^{1+}_{(\text{aq})}$	0.81	3.4	0.02	0.023 (0.02) + (0) + 0.003
2	$\text{NpO}_2^{1+}_{(\text{aq})}$	$\text{NpO}_2^{2+}_{(\text{aq})}$	1.5	32	0.02	0.026 (0.02) + (0) + 0.006
3	$\text{NpO}_2^{1+}_{(\text{aq})}$	$\text{NpO}_2^{2+}_{(\text{aq})}$	3.1	31	0.04	0.052 (0.04) + (0) + 0.012
4	$\text{NpO}_2^{1+}_{(\text{aq})}$	$\text{NpO}_2^{2+}_{(\text{aq})}$	1.5	160	0.02	0.026 (0.02) + (0) + 0.006
5	$\text{NpO}_2^{1+}_{(\text{aq})}$	$\text{NpO}_2^{2+}_{(\text{aq})}$	3.1	160	0.04	0.052 (0.04) + (0) + 0.012
6	$\text{Np}^{4+}_{(\text{aq})}$	$\text{NpO}_2^{2+}_{(\text{aq})}$	3.1	31	0.04	9.1 (0.04) + (9.0) + 0.012
7	$\text{Np}^{4+}_{(\text{aq})}$	$\text{NpO}_2^{2+}_{(\text{aq})}$	1.5	32	0.02	9.2 (0.02) + (9.2) + 0.006
8	$\text{Np}^{4+}_{(\text{aq})}$	$\text{NpO}_2^{2+}_{(\text{aq})}$	1.5	160	0.02	9.2 (0.02) + (9.2) + 0.006
9	$\text{Np}^{4+}_{(\text{aq})}$	$\text{NpO}_2^{2+}_{(\text{aq})}$	3.1	160	0.04	9.1 (0.04) + (9.0) + 0.012
10	$\text{Np}^{4+}_{(\text{aq})}$	$\text{NpO}_2^{2+}_{(\text{aq})}$	2.9	160	5.4	5.4 (5.4) + (0) + 0.012
11	$\text{Np}^{4+}_{(\text{aq})}$	$\text{Np}^{4+}_{(\text{aq})}$	1.5	160	6.1	6.1 (6.1) + (0) + 0.006
12	$\text{Np}^{4+}_{(\text{aq})}$	$\text{Np}^{4+}_{(\text{aq})}$	1.5	160	7.1 <sup>c</sup>	7.1 (7.1) + (0) + 0.006
13	$\text{Np}^{4+}_{(\text{aq})}$	$\text{Np}^{4+}_{(\text{aq})}$	2.9	160	7	7.0 (7.0) + (0) + 0.012
14	$\text{Np}^{4+}_{(\text{aq})}$	$\text{Np}^{4+}_{(\text{aq})}$	2.9	32	5.4	5.4 (5.4) + (0) + 0.012
15	$\text{Np}^{4+}_{(\text{aq})}$	$\text{Np}^{4+}_{(\text{aq})}$	2.9	31	6.2	6.2 (6.2) + (0) + 0.012
16	$\text{Np}^{4+}_{(\text{aq})}$	$\text{Np}^{4+}_{(\text{aq})}$	1.5	32	6.1	6.1 (6.1) + (0) + 0.006
17	$\text{Np}^{4+}_{(\text{aq})}$	$\text{Np}^{4+}_{(\text{aq})}$	2.9	31	7.0	7.0 (7.0) + (0) + 0.012
18	$\text{Np}^{4+}_{(\text{aq})}$	$\text{Np}^{4+}_{(\text{aq})}$	2.9	31	5.3	5.4 (5.4) + (0) + 0.012
19	$\text{Np}^{4+}_{(\text{aq})}$	$\text{Np}^{4+}_{(\text{aq})}$	1.5	32	7.1	7.1 (7.1) + (0) + 0.006

<sup>a</sup> Calculations show ingoing  $\text{Cl}^{1-}_{(\text{aq})}$  content from  $\text{HCl}_{(\text{aq})}$ ,  $\text{LiCl}_{(\text{aq})}$ , and  $\text{NpCl}_{4(\text{aq})}$  and do not include  $\text{NaClO}_{2(\text{aq})}$  nor  $\text{NaClO}_{2(\text{aq})}$  disproportionation products.

aqueous solution that was dilute in  $\text{Np}^{4+}_{(\text{aq})}$  (1.6 mM, 1  $\mu\text{mol}$ , 600  $\mu\text{L}$ ) and dilute in  $\text{HCl}_{(\text{aq})}$  (0.02 M). Analyses by UV-vis spectroscopy showed that the  $\text{Np}^{4+}_{(\text{aq})}$  reagent reacted immediately with  $\text{NaClO}_{2(\text{aq})}$ . Then, the  $\text{Np}^{4+}_{(\text{aq})}$  and  $\text{NpO}_2^{1+}_{(\text{aq})}$  product-to-reagent ratio stabilized within 60 s. These experiments showed that reacting a small amount [more than two equivalents vs.  $\text{Np}^{4+}_{(\text{aq})}$ ] of  $\text{NaClO}_{2(\text{aq})}$  (2.2  $\mu\text{mol}$ ; 330  $\mu\text{L}$  of a 7.2 mM solution) with one equivalent of  $\text{Np}^{4+}_{(\text{aq})}$  (1  $\mu\text{mol}$ ) generated a one-to-one mixture of  $\text{Np}^{4+}_{(\text{aq})}$  and  $\text{NpO}_2^{1+}_{(\text{aq})}$ . Increasing the quantity of  $\text{NaClO}_{2(\text{aq})}$  increased the amount of  $\text{NpO}_2^{1+}_{(\text{aq})}$  generated until approximately 4.5 equivalents of  $\text{NaClO}_{2(\text{aq})}$  (4.5  $\mu\text{mol}$ ; 630  $\mu\text{L}$  of 7.2 mM solution) vs. one equivalent of  $\text{Np}^{4+}_{(\text{aq})}$  (1  $\mu\text{mol}$ ) had been added. In this situation, the  $\text{Np}^{4+}_{(\text{aq})}$  reagent was completely consumed and  $\text{NpO}_2^{1+}_{(\text{aq})}$  formed. The  $\text{NpO}_2^{1+}_{(\text{aq})}$  species was stable under ambient conditions during a 5-day monitoring process.

The  $\text{Np}^{4+}_{(\text{aq})}$  cation also oxidized to  $\text{NpO}_2^{1+}_{(\text{aq})}$  when the  $\text{NaClO}_{2(\text{aq})}$  concentration was increased from 3.4 mM to 32 mM; entry 2 in Table 1. However, under these conditions the  $\text{NpO}_2^{1+}_{(\text{aq})}$  mono-cation was not stable with time and further oxidation to  $\text{NpO}_2^{2+}_{(\text{aq})}$  occurred (Fig. 5). Absorption peaks from  $\text{NpO}_2^{1+}_{(\text{aq})}$  at 10 400  $\text{cm}^{-1}$  (980 nm) and 8700  $\text{cm}^{-1}$  (1150 nm) were replaced within one hour by features characteristic of  $\text{NpO}_2^{2+}_{(\text{aq})}$ ; 7900  $\text{cm}^{-1}$  (1270 nm). Monitoring this product by UV-vis spectroscopy revealed that  $\text{NpO}_2^{2+}_{(\text{aq})}$  was stable for at least 5 days, when monitoring ceased. Comparing this data with that presented above highlighted how formation of  $\text{NpO}_2^{2+}_{(\text{aq})}$  vs.  $\text{NpO}_2^{1+}_{(\text{aq})}$  could be controlled based on ingoing  $\text{NaClO}_{2(\text{aq})}$  concentrations. The  $\text{NpO}_2^{1+}_{(\text{aq})}$  mono-cation was generated when the ingoing concentration of  $\text{NaClO}_{2(\text{aq})}$  was low (3.4 mM; entry 1, Table 1). In contrast, the  $\text{NpO}_2^{2+}_{(\text{aq})}$



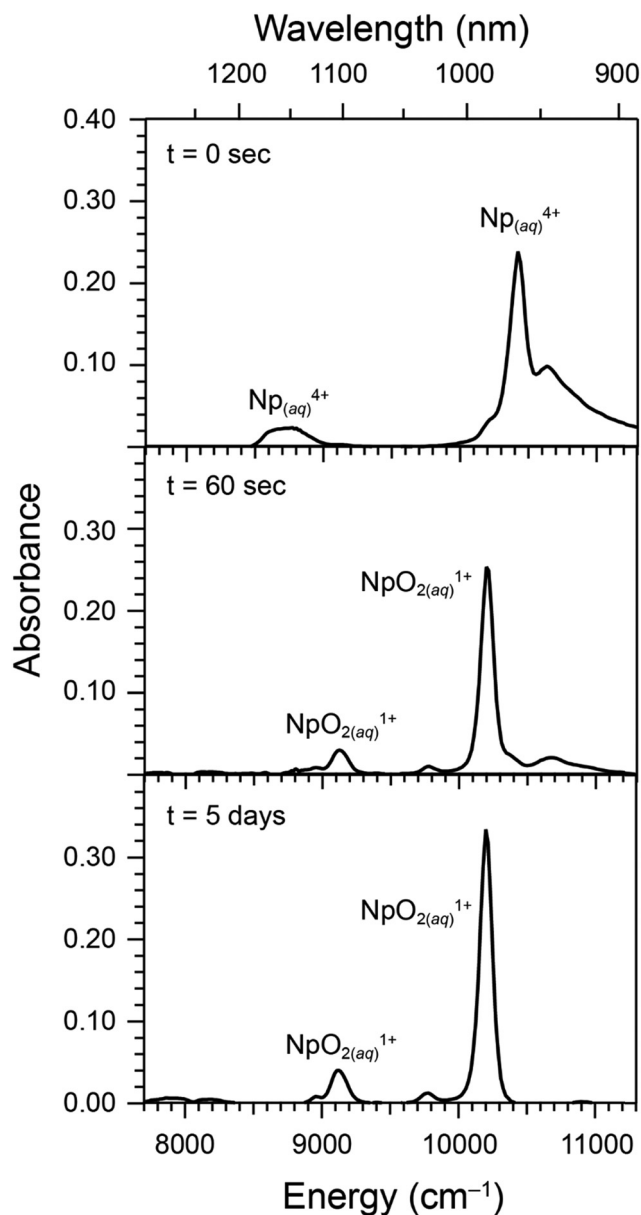


Fig. 2 UV-vis absorption spectra documenting oxidation of  $\text{Np}^{4+}_{(\text{aq})}$  to  $\text{NpO}_2^{1+}_{(\text{aq})}$  as a function of time (top, initial; middle, after 60 s; bottom, after 5 days). Entry #1 in Table 1 describe the reaction conditions. Ingoing concentrations were low for  $\text{Np}^{4+}_{(\text{aq})}$  (0.81 mM), low for  $\text{NaClO}_{2(\text{aq})}$  (3.4 mM), low for  $\text{H}^{1+}_{(\text{aq})}$  (0.02 M), and low for  $\text{Cl}^{1-}_{(\text{aq})}$  (0.026 M). Hence, the ingoing  $\text{HCl}_{(\text{aq})}$  concentration was 0.02 M and there no added  $\text{LiCl}_{(\text{aq})}$ .

di-cation was generated with the ingoing concentration of  $\text{NaClO}_2$  was high (32 mM; entry 2, Table 1). There is a disclaimer associated with this conclusion. Controlling oxidation of  $\text{Np}^{4+}_{(\text{aq})}$  to either  $\text{NpO}_2^{1+}_{(\text{aq})}$  or  $\text{NpO}_2^{2+}_{(\text{aq})}$  required the ingoing concentrations of  $\text{Np}^{4+}_{(\text{aq})}$  and  $\text{NaClO}_{2(\text{aq})}$  as well as ingoing concentrations of  $\text{Cl}^{1-}_{(\text{aq})}$  and  $\text{H}^{1+}_{(\text{aq})}$  to be carefully managed; see experiments captured in entries 2 to 19 in Table 1. For example, adding  $\text{NaClO}_{2(\text{aq})}$  (32 mM or 31 mM to 160 mM entries 2 to 5) to  $\text{Np}^{4+}_{(\text{aq})}$  (1.5 mM to 3.1 mM) with low  $\text{Cl}^{1-}_{(\text{aq})}$  and  $\text{H}^{1+}_{(\text{aq})}$  concentrations (0.02 to 0.04 M) generated  $\text{NpO}_2^{1+}_{(\text{aq})}$  at the 1 h timestamp and that  $\text{NpO}_2^{1+}_{(\text{aq})}$  converted

to  $\text{NpO}_2^{2+}_{(\text{aq})}$  within 24 h. Increasing the  $\text{Cl}^{1-}_{(\text{aq})}$  content changed this outcome. By using the  $\text{LiCl}_{(\text{aq})}$  to increase  $\text{Cl}^{1-}_{(\text{aq})}$  concentrations to  $>9$  M (entries 6 to 9) and holding the  $\text{H}^{1+}_{(\text{aq})}$  concentration low (0.02 M to 0.04 M) stabilized the  $\text{Np}^{4+}_{(\text{aq})}$  regent as the dominant specie present in solution at the 1 h timestamp. Then, the  $\text{Np}^{4+}_{(\text{aq})}$  cation oxidized to  $\text{NpO}_2^{2+}_{(\text{aq})}$  within 24 h. Increasing both  $\text{H}^{1+}_{(\text{aq})}$  and  $\text{Cl}^{1-}_{(\text{aq})}$  content to  $>5.4$  M shut down the oxidation of  $\text{Np}^{4+}_{(\text{aq})}$  to  $\text{NpO}_2^{2+}_{(\text{aq})}$  and  $\text{Np}^{4+}_{(\text{aq})}$  was stable throughout the 24 h monitoring duration.

We characterized the reaction rates for entry 2 in Table 1 that were associated with the transformation of (1st)  $\text{Np}^{4+}_{(\text{aq})}$  to  $\text{NpO}_2^{1+}_{(\text{aq})}$  and (2nd)  $\text{NpO}_2^{1+}_{(\text{aq})}$  to  $\text{NpO}_2^{2+}_{(\text{aq})}$  (Fig. 6). The oxidation of  $\text{Np}^{4+}_{(\text{aq})}$  to  $\text{NpO}_2^{1+}_{(\text{aq})}$  proceeded with fractional order dependence on both the  $\text{Np}^{4+}_{(\text{aq})}$  ( $0.300 \pm 0.008$ ) and  $\text{ClO}_2^{1-}_{(\text{aq})}$  ( $0.110 \pm 0.002$ ) reagents. The reaction rate law derived from the oxidation of  $\text{NpO}_2^{1+}_{(\text{aq})}$  to  $\text{NpO}_2^{2+}_{(\text{aq})}$  was approximately first order in  $\text{NpO}_2^{1+}_{(\text{aq})}$  ( $0.98 \pm 0.02$ ) and higher order ( $1.7 \pm 0.01$ ) with regards to  $\text{NaClO}_{2(\text{aq})}$ . No evidence was obtained that suggested the  $\text{NpO}_2^{2+}_{(\text{aq})}$  di-cation formed *via* disproportionation of the  $\text{NpO}_2^{1+}_{(\text{aq})}$  mono-cation to  $\text{Np}^{4+}_{(\text{aq})}$  and  $\text{NpO}_2^{2+}_{(\text{aq})}$ . For instance, regeneration of  $\text{Np}^{4+}_{(\text{aq})}$  – or a sustained presence of  $\text{Np}^{4+}_{(\text{aq})}$  – after initial formation of  $\text{NpO}_2^{1+}_{(\text{aq})}$  was not observed. Instead, our data suggested that  $\text{NpO}_2^{1+}_{(\text{aq})}$  converted directly to  $\text{NpO}_2^{2+}_{(\text{aq})}$ . Admittedly, this data does not completely rule out the possibility of a low-concentration  $\text{Np}^{4+}_{(\text{aq})}$  transient intermediate nor the possibility of disproportionation reaction pathways. However, with the data in hand at this time, it seemed possible that  $\text{NpO}_2^{1+}_{(\text{aq})}$  was converted directly  $\text{NpO}_2^{2+}_{(\text{aq})}$  by the redox processes accessed with  $\text{NaClO}_{2(\text{aq})}$  and/or  $\text{NaClO}_{2(\text{aq})}$  decomposition productions, see eqn (1) and (2).

There were other experimental conditions that generated a  $\text{NpO}_2^{1+}_{(\text{aq})}$  intermediate (within an hour) that in turn oxidized  $\text{Np}^{4+}_{(\text{aq})}$  to  $\text{NpO}_2^{2+}_{(\text{aq})}$  (entries 3 to 10 from Table 1). Each of these scenarios had three commonalities: the ingoing  $\text{H}^{1+}_{(\text{aq})}$  concentrations ranged 0.02 to 5.4 M, the ingoing  $\text{Cl}^{1-}_{(\text{aq})}$  concentration ranged from 0.026 to 9.1 M, the ingoing  $\text{Np}^{4+}_{(\text{aq})}$  concentrations were larger than 0.81 mM (entries 2 through 5 in Table 1), and the ingoing  $\text{NaClO}_{2(\text{aq})}$  concentrations were high ( $>31$  mM). Modifying these variables changed the reaction outcomes. For example, increasing  $\text{Cl}^{1-}_{(\text{aq})}$  content to  $>9$  M – using  $\text{LiCl}_{(\text{aq})}$  – masked observation of a  $\text{NpO}_2^{1+}_{(\text{aq})}$  intermediate (entries 6 to 10 in Table 1; Fig. 7 and 8). In these situations, we only observed disappearance of  $\text{Np}^{4+}_{(\text{aq})}$  and formation of  $\text{NpO}_2^{2+}_{(\text{aq})}$  (entries 6 to 9). Monitoring this reaction as a function of time revealed that the loss of  $\text{Np}^{4+}_{(\text{aq})}$  and formation of  $\text{NpO}_2^{2+}_{(\text{aq})}$  proceeded at fourth order ( $4.11 \pm 0.09$ ) dependence with respect to  $\text{Np}^{4+}_{(\text{aq})}$  and fractional ( $0.50 \pm 0.07$ ) order with respect to  $\text{NaClO}_{2(\text{aq})}$  (Fig. 8). Although fourth order rate laws have been measured before,<sup>64–67</sup> we acknowledge their unusuality and rarity. In our case, we speculated that the measured fourth order rate law resulted from numerous intertwined reactions that were occurring simultaneously. One obvious possibility included  $\text{NaClO}_{2(\text{aq})}$  decomposition products reacting directly with  $\text{Np}^{4+}_{(\text{aq})}$  to generate  $\text{NpO}_2^{2+}_{(\text{aq})}$  and maybe a transient  $\text{NpO}_2^{1+}_{(\text{aq})}$  intermediate. It is also



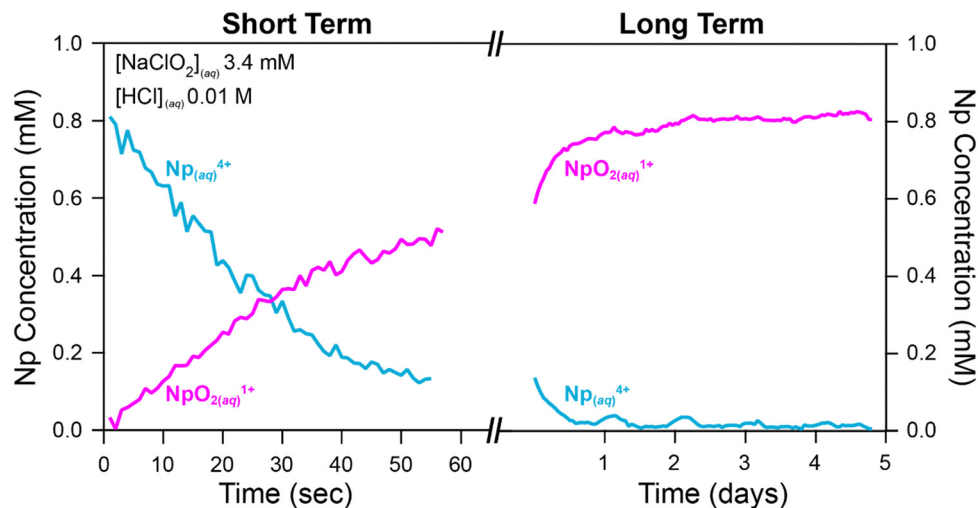


Fig. 3 Peak intensity from UV-vis spectra from  $\text{Np}^{4+}_{(\text{aq})}$  (teal trace from the absorption peak at  $10\,400\text{ cm}^{-1}$ ,  $960\text{ nm}$ ) and  $\text{NpO}_2^{1+}_{(\text{aq})}$  (magenta trace from the absorption peak at  $9100\text{ cm}^{-1}$ ,  $1100\text{ nm}$ ) plotted as a function of two different time intervals (left, 0 to 1 min; right, 0.1 to 5 days). Entry #1 in Table 1 describe the reaction conditions. Ingoing concentrations were low for  $\text{Np}^{4+}_{(\text{aq})}$  (0.81 mM), low for  $\text{NaClO}_{2(\text{aq})}$  (3.4 mM), low for  $\text{H}^{1+}_{(\text{aq})}$  (0.02 M), and low for  $\text{Cl}^{1-}_{(\text{aq})}$  (0.02 M). Hence, the ingoing  $\text{HCl}_{(\text{aq})}$  concentration was 0.02 M and there was no added LiCl.

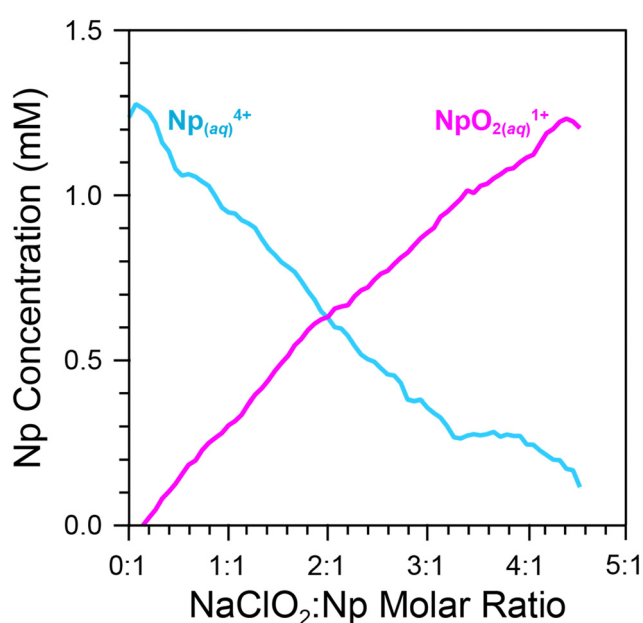


Fig. 4 A plot showing how oxidation of  $\text{Np}^{4+}_{(\text{aq})}$  (teal trace from the absorption peak at  $10\,400\text{ cm}^{-1}$ ,  $960\text{ nm}$ ) to  $\text{NpO}_2^{1+}_{(\text{aq})}$  (magenta trace from the absorption peak at  $10\,200\text{ cm}^{-1}$ ,  $980\text{ nm}$ ) depended on the amount of added  $\text{NaClO}_{2(\text{aq})}$ . Ingoing concentrations were low for  $\text{Np}^{4+}_{(\text{aq})}$  (1.6 mM), low for  $\text{H}^{1+}_{(\text{aq})}$  (0.02 M), and low for  $\text{Cl}^{1-}_{(\text{aq})}$  (0.026 M). Hence, the ingoing  $\text{HCl}_{(\text{aq})}$  concentration was 0.02 M and there was no added LiCl. The  $\text{NaClO}_{2(\text{aq})}$  started at zero and increased with addition of  $10\text{ }\mu\text{L}$  aliquots of a  $\text{NaClO}_{2(\text{aq})}$  (7.2 mM) solution. Concentrations were corrected for dilutions.

possible that the  $\text{NaClO}_{2(\text{aq})}$  could react with  $\text{Np}^{4+}_{(\text{aq})}$  directly. If any  $\text{NpO}_2^{1+}_{(\text{aq})}$  formed, that mono-cation could disproportionate to generate  $\text{NpO}_2^{2+}_{(\text{aq})}$  and regenerate  $\text{Np}^{4+}_{(\text{aq})}$ . Note, we did not detect evidence for formation of a  $\text{Np}^{4+}_{(\text{aq})}$  intermediate by UV-vis. Hence, if  $\text{Np}^{4+}_{(\text{aq})}$  formed, it was generated in relatively small quantities.

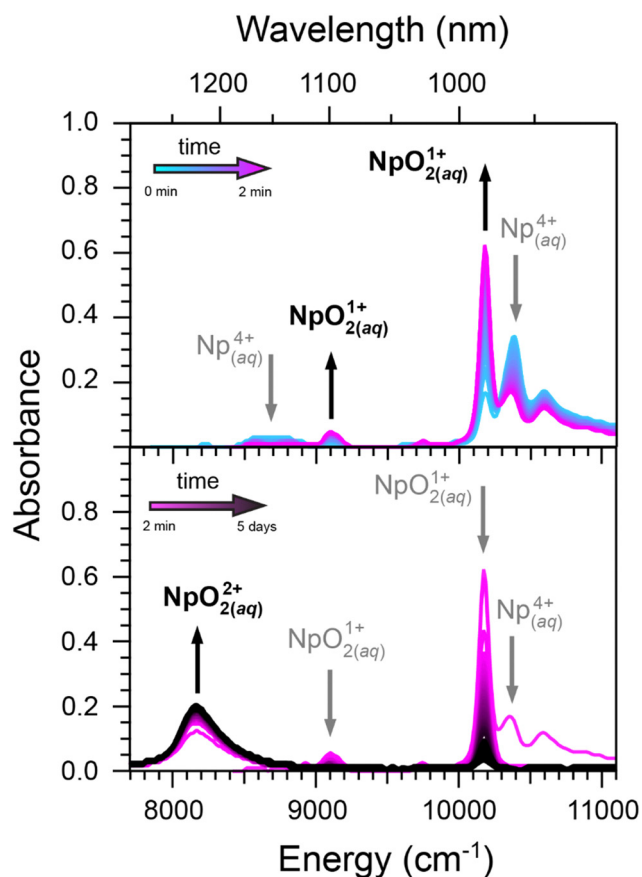


Fig. 5 Top: UV-vis absorption spectra documenting oxidation of  $\text{Np}^{4+}_{(\text{aq})}$  (teal trace) to  $\text{NpO}_2^{1+}_{(\text{aq})}$  (magenta trace) as a function of time (2 min). Bottom: Subsequent oxidation of  $\text{NpO}_2^{1+}_{(\text{aq})}$  (magenta trace) to  $\text{NpO}_2^{2+}_{(\text{aq})}$  (black trace). Entry #2 in Table 1 describe the reaction conditions. Ingoing concentrations were low for  $\text{Np}^{4+}_{(\text{aq})}$  (1.5 mM), low for  $\text{NaClO}_{2(\text{aq})}$  (32 mM), low for  $\text{H}^{1+}_{(\text{aq})}$  (0.02 M), and low for  $\text{Cl}^{1-}_{(\text{aq})}$  (0.02 M). Hence, the ingoing  $\text{HCl}_{(\text{aq})}$  concentration was 0.02 M and there was no added LiCl.



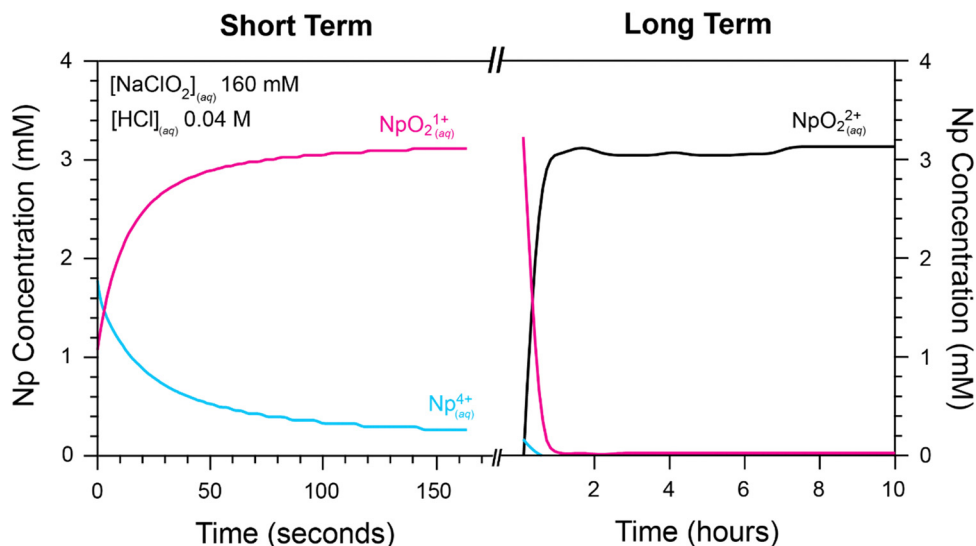


Fig. 6 Peak intensity from UV-vis spectra from  $\text{Np}^{4+}_{(\text{aq})}$  (teal trace at  $10\,400\text{ cm}^{-1}$ ,  $960\text{ nm}$ ),  $\text{NpO}_2^{1+}_{(\text{aq})}$  (magenta trace at  $9100\text{ cm}^{-1}$ ,  $1100\text{ nm}$ ), and  $\text{NpO}_2^{2+}_{(\text{aq})}$  (black trace at  $8200\text{ cm}^{-1}$ ,  $1250\text{ nm}$ ) plotted as a function of two different time intervals (left, 0 to 2 min; right, approximately 2 min to 10 days). Entry #5 in Table 1 describe the reaction conditions. Ingoing concentrations were high for  $\text{Np}^{4+}_{(\text{aq})}$  (3.1 mM), high for  $\text{NaClO}_{2(\text{aq})}$  (160 mM), low for  $\text{H}^{1+}_{(\text{aq})}$  (0.04 M), and low for  $\text{Cl}^{1-}_{(\text{aq})}$  (0.052 M). Hence, the ingoing  $\text{HCl}_{(\text{aq})}$  concentration was 0.04 M and there was no added LiCl.

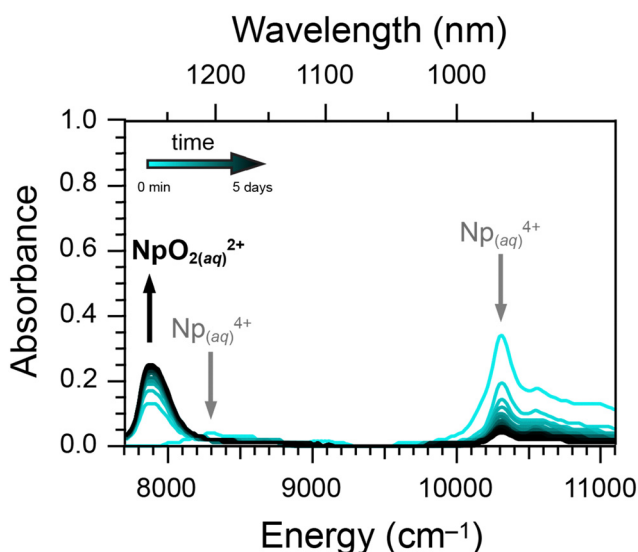


Fig. 7 UV-vis absorption spectra documenting oxidation of  $\text{Np}^{4+}_{(\text{aq})}$  (teal trace) to  $\text{NpO}_2^{2+}_{(\text{aq})}$  (black trace). Entry #7 in Table 1 describe the reaction conditions. Ingoing concentrations were 1.5 mM for  $\text{Np}^{4+}_{(\text{aq})}$ , 32 mM for  $\text{NaClO}_{2(\text{aq})}$ , 0.02 M for  $\text{H}^{1+}_{(\text{aq})}$ , and 9.2 M for  $\text{Cl}^{1-}_{(\text{aq})}$ . Hence the ingoing  $\text{HCl}_{(\text{aq})}$  concentration was 0.02 M and for  $\text{LiCl}_{(\text{aq})}$  was 9.2 M.

Regardless, we are intrigued by this reaction and our current efforts are focused on better characterizing how  $\text{NaClO}_{2(\text{aq})}$  incites  $\text{Np}^{4+}_{(\text{aq})}$  transformation to  $\text{NpO}_2^{2+}_{(\text{aq})}$  in high  $\text{Cl}^{1-}_{(\text{aq})}$  solutions.

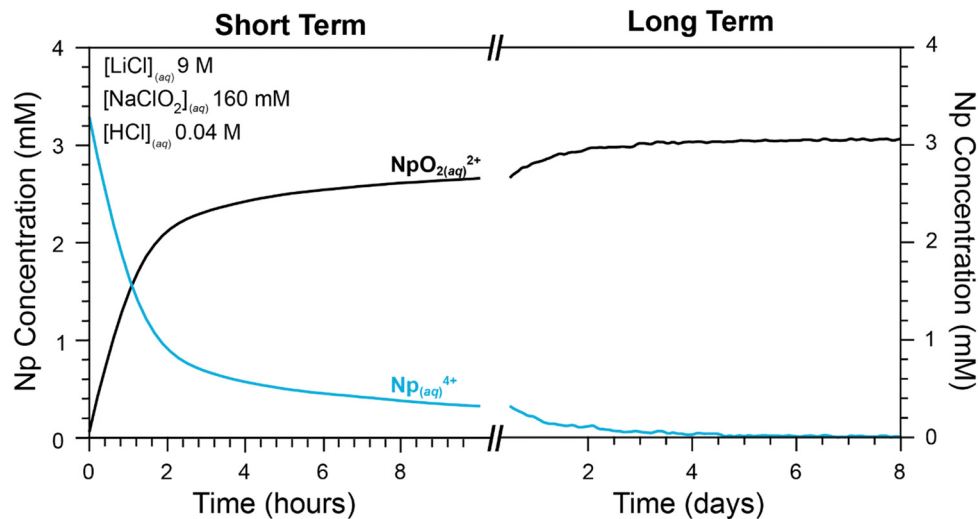
As shown in Table 1 (entries 11 to 19), oxidation of  $\text{Np}^{4+}_{(\text{aq})}$  to  $\text{NpO}_2^{2+}_{(\text{aq})}$  was halted when the  $\text{H}^{1+}_{(\text{aq})}$  and  $\text{Cl}^{1-}_{(\text{aq})}$  concentrations were  $\geq 5.4\text{ M}$ , when the ingoing  $\text{Np}^{4+}_{(\text{aq})}$  concentration ranged 1.5 to 2.9 mM, and the  $\text{NaClO}_{2(\text{aq})}$  concentration was high (31 to 160 mM). It is particularly important to highlight these observations because of their relevance to EXCEL processing

methods described in the introduction. Notice, that the  $\text{NaClO}_{2(\text{aq})}$  redox agent held neptunium in the +4 oxidation state under conditions that mimicked plutonium processing by EXCEL: when the  $\text{H}^{1+}_{(\text{aq})}$  and  $\text{Cl}^{1-}_{(\text{aq})}$  concentrations were  $> 5.4\text{ M}$  and the  $\text{NaClO}_{2(\text{aq})}$  concentrations varied from 31 mM to 160 mM, and  $\text{Np}^{4+}_{(\text{aq})}$  concentrations varied from 1.5 mM to 2.9 mM (entries 11 to 19 in Table 1). This data suggested that – all things being equal – neptunium should follow plutonium through within the EXCEL processing steps.

#### Molar extinction coefficients

Table 2 compares molar extinction coefficients ( $\epsilon$ ) for neptunium absorption peaks calculated for experimental conditions described in Table 1 when Np was in a single oxidation state. These data were important for quantifying  $\text{Np}^{4+}_{(\text{aq})}$ ,  $\text{NpO}_2^{1+}_{(\text{aq})}$ , and  $\text{NpO}_2^{2+}_{(\text{aq})}$  abundance in the reactions studied herein and provided insight into the solution phase behavior of  $\text{Np}^{4+}_{(\text{aq})}$  and  $\text{NpO}_2^{2+}_{(\text{aq})}$ . In terms of the latter topic, we observed that  $\epsilon$  for the  $\text{Np}^{4+}_{(\text{aq})}$  absorption feature at  $10\,400\text{ cm}^{-1}$  ( $960\text{ nm}$ ) decreased with increasing  $\text{Cl}^{1-}_{(\text{aq})}$  concentration. In addition, increasing  $\text{Cl}^{1-}_{(\text{aq})}$  shifted the  $\text{Np}^{4+}_{(\text{aq})}$  absorption peaks lower in energy (Fig. 9). These absorption changes are often associated with increased  $\text{Cl}^{1-}_{(\text{aq})}$  complexation of neptunium.<sup>36,37,68</sup> It was interesting to note that  $\epsilon$  for the  $\text{Np}^{4+}_{(\text{aq})}$  peak at  $8700\text{ cm}^{-1}$  ( $1150\text{ nm}$ ) peak was shifted in energy but the  $\epsilon$  value was unaffected by changes in  $\text{Cl}^{1-}_{(\text{aq})}$  content (see Fig. 9). For  $\text{NpO}_2^{2+}_{(\text{aq})}$ , increasing the  $\text{Cl}^{1-}_{(\text{aq})}$  caused the absorption feature at  $8200\text{ cm}^{-1}$  ( $1220\text{ nm}$ ) to decrease in energy and shift higher in energy, which is also often attributed to increased  $\text{Cl}^{1-}_{(\text{aq})}$  complexation.<sup>69</sup> Another intriguing observation was that  $\epsilon$  for the  $10\,400\text{ cm}^{-1}$  absorbance peak from  $\text{Np}^{4+}_{(\text{aq})}$  was more impacted by  $\text{Cl}^{1-}_{(\text{aq})}$  concentrations than those from  $\text{NpO}_2^{2+}_{(\text{aq})}$  at  $8200\text{ cm}^{-1}$ . This stronger depended of  $\epsilon$  on  $\text{Cl}^{1-}_{(\text{aq})}$  concentration tracked with the Lewis acidity for the  $\text{Np}^{4+}_{(\text{aq})}$  vs.





**Fig. 8** Peak intensity from UV-vis spectra from  $\text{Np}^{4+}_{(\text{aq})}$  (teal trace at  $10\,400\text{ cm}^{-1}$ ,  $960\text{ nm}$ ) and  $\text{NpO}_2^{2+}_{(\text{aq})}$  (black trace at  $7900\text{ cm}^{-1}$ ,  $1270\text{ nm}$ ) plotted as a function of two different time intervals (left, 0 to 2 min; right, 2 min to 10 days). Entry #9 in Table 1 describe the reaction conditions. Ingoing concentrations were high for  $\text{Np}^{4+}_{(\text{aq})}$  (3.1 mM), high for  $\text{NaClO}_{2(\text{aq})}$  (160 mM), low for  $\text{H}^{1+}_{(\text{aq})}$  (0.04 M), and high for  $\text{Cl}^{1-}_{(\text{aq})}$  (9.1 M). Hence, the ingoing  $\text{HCl}_{(\text{aq})}$  concentration was 0.04 M and for  $\text{LiCl}_{(\text{aq})}$  was 9.1 M.

**Table 2** Molar extinction coefficients ( $\epsilon$ ) obtained for neptunium species present in aqueous solutions under the experimental conditions specified below

Oxidation state	Energy ( $\text{cm}^{-1}$ )	Wavelength (nm)	Table 1 entry #	$\epsilon$ ( $\text{M}^{-1}\text{ cm}^{-1}$ )
$\text{Np}^{4+}_{(\text{aq})}$	10 400	960	1	294
$\text{Np}^{4+}_{(\text{aq})}$	10 400	960	3	304
$\text{Np}^{4+}_{(\text{aq})}$	10 400	960	2	341
$\text{Np}^{4+}_{(\text{aq})}$	10 400	960	7	220
$\text{Np}^{4+}_{(\text{aq})}$	10 400	960	6	171
$\text{Np}^{4+}_{(\text{aq})}$	10 400	960	N/A	$162^a$
$\text{Np}^{4+}_{(\text{aq})}$	8700	1150	2	19
$\text{Np}^{4+}_{(\text{aq})}$	8700	1150	3	20
$\text{Np}^{4+}_{(\text{aq})}$	8700	1150	7	26
$\text{Np}^{4+}_{(\text{aq})}$	8700	1150	6	20
$\text{NpO}_2^{1+}_{(\text{aq})}$	10 200	980	1	334
$\text{NpO}_2^{1+}_{(\text{aq})}$	10 200	980	3	660
$\text{NpO}_2^{1+}_{(\text{aq})}$	10 200	980	4	790
$\text{NpO}_2^{1+}_{(\text{aq})}$	10 200	980	5	446
$\text{NpO}_2^{1+}_{(\text{aq})}$	10 200	980	N/A	$375^a$
$\text{NpO}_2^{1+}_{(\text{aq})}$	9100	1100	1	41
$\text{NpO}_2^{1+}_{(\text{aq})}$	9100	1100	3	53
$\text{NpO}_2^{1+}_{(\text{aq})}$	9100	1100	4	60
$\text{NpO}_2^{1+}_{(\text{aq})}$	9100	1100	5	50
$\text{NpO}_2^{2+}_{(\text{aq})}$	8200	1220	4	127
$\text{NpO}_2^{2+}_{(\text{aq})}$	8200	1220	5	115
$\text{NpO}_2^{2+}_{(\text{aq})}$	8200	1220	N/A	$43^a$
$\text{NpO}_2^{2+}_{(\text{aq})}$	7900	1270	8	190
$\text{NpO}_2^{2+}_{(\text{aq})}$	7900	1270	7	190
$\text{NpO}_2^{2+}_{(\text{aq})}$	7900	1270	9	174

<sup>a</sup> In 2 M  $\text{HClO}_{4(\text{aq})}$  from ref. 37.

$\text{NpO}_2^{2+}_{(\text{aq})}$  cations as well as the variances in  $\text{Cl}^{1-}_{(\text{aq})}$  complexation constants.<sup>37,70</sup>

## Outlook

Herein, we observed that  $\text{NaClO}_{2(\text{aq})}$  enabled  $\text{Np}^{4+}_{(\text{aq})}$  to access many redox events in acidic aqueous solutions (Scheme 1).

Moreover, oxidation product identities could be controlled as a function of time by manipulating four variables; ingoing concentrations of (1)  $\text{Np}^{4+}_{(\text{aq})}$ , (2)  $\text{NaClO}_{2(\text{aq})}$ , (3)  $\text{Cl}^{1-}_{(\text{aq})}$ , and (4)  $\text{H}^{1+}_{(\text{aq})}$ . For example,  $\text{NpO}_2^{1+}_{(\text{aq})}$  exclusively formed when  $\text{Np}^{4+}_{(\text{aq})}$ ,  $\text{NaClO}_{2(\text{aq})}$ ,  $\text{Cl}^{1-}_{(\text{aq})}$ , and  $\text{H}^{1+}_{(\text{aq})}$  concentrations were all relatively low. Increasing the  $\text{Cl}^{1-}_{(\text{aq})}$  concentration and/or increasing the  $\text{NaClO}_{2(\text{aq})}$  concentration generated  $\text{NpO}_2^{2+}_{(\text{aq})}$ , provided that the  $\text{H}^{1+}_{(\text{aq})}$  concentration was also low. Finally,  $\text{Np}^{4+}_{(\text{aq})}$  was stabilized when the  $\text{H}^{1+}_{(\text{aq})}$  and  $\text{Cl}^{1-}_{(\text{aq})}$  concentrations were high ( $> 5.4\text{ M}$ ), regardless of the ingoing  $\text{NaClO}_{2(\text{aq})}$  and  $\text{Np}^{4+}_{(\text{aq})}$  concentrations.

Regarding applications for plutonium processing methods described in the introduction, we make the following prediction. Our data suggested that the neptunium oxidation state during plutonium processing will be either maintained at +4 or a mixture of +4 and +6 if plutonium separations are carried out at high  $\text{NaClO}_{2(\text{aq})}$ , high  $\text{Cl}^{1-}_{(\text{aq})}$ , high  $\text{H}^{1+}_{(\text{aq})}$ , and low  $\text{Np}^{4+}_{(\text{aq})}$  concentrations. If we are correct, the  $^{237}\text{Np}_{(\text{aq})}$  contaminant will follow  $\text{Pu}^{4+}_{(\text{aq})}$  through processing lines. However, we acknowledge that there are substantial differences between the fundamental studies carried out here *vs.* real-world processing environments (*e.g.* radiation effects, scaling effects from other impurities, *etc.*). To improve relevancy, future work will center on understanding how other constituents present in various plutonium and americium waste streams impact redox chemistry between  $^{237}\text{Np}_{(\text{aq})}$  and  $\text{NaClO}_{2(\text{aq})}$ .

The data in Table 1 and Scheme 1 also demonstrated that the neptunium,  $\text{NaClO}_{2(\text{aq})}$ ,  $\text{H}^{1+}_{(\text{aq})}$ , and  $\text{Cl}^{1-}_{(\text{aq})}$  concentration variables were intertwined and must be delicately managed to control the  $\text{Np}^{4+}_{(\text{aq})}$  oxidation to  $\text{NpO}_2^{1+}_{(\text{aq})}$  *vs.*  $\text{NpO}_2^{2+}_{(\text{aq})}$  by way of  $\text{NaClO}_{2(\text{aq})}$ . These studies provided a more sophisticated understanding of aqueous actinide electron transfer chemistry and offer opportunity to better predict and control actinide redox processes. Future efforts will center on developing a





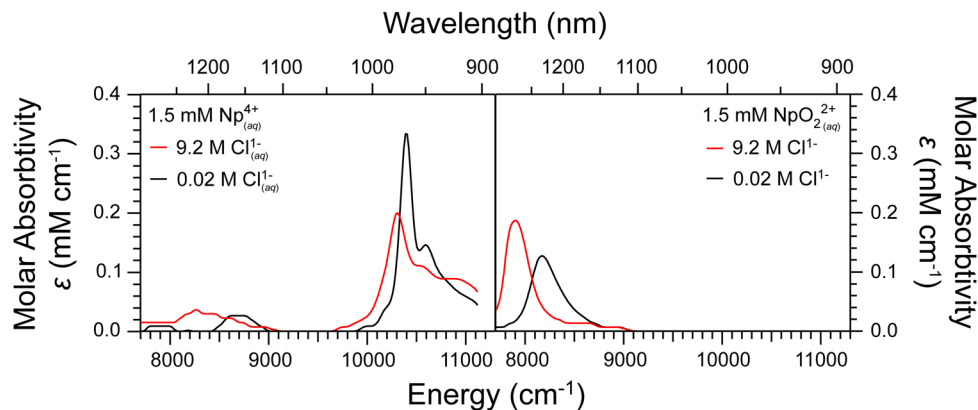


Fig. 9 UV-vis absorption spectra documenting oxidation of the  $\text{Np}^{4+}_{(\text{aq})}$  reagent (left) to the  $\text{NpO}_2^{2+}_{(\text{aq})}$  product (right) from entries #4 (black trace) and #8 (red trace). This figure highlights how moving from low concentration  $\text{Cl}^{-}$  (0.026 M; entry #4, black trace) to high concentration  $\text{Cl}^{-}$  (9.2 M, entry #8, red trace) impacts the absorption features. Note, other than  $\text{Cl}^{-}$  concentration, the other experimental variables were held constant between these two experiments. Ingoing concentrations associated with the entry #4 (black trace) experiment were 1.5 mM for  $\text{Np}^{4+}_{(\text{aq})}$ , 160 mM for  $\text{NaClO}_{2(\text{aq})}$ , 0.02 M for  $\text{H}^{1+}_{(\text{aq})}$ , and 0.026 M for  $\text{Cl}^{-}_{(\text{aq})}$ . In this case, the ingoing  $\text{HCl}_{(\text{aq})}$  concentration was 0.02 M and there was no  $\text{LiCl}$  added. Ingoing concentrations associated with the entry #8 (red trace) experiment were 1.5 mM for  $\text{Np}^{4+}_{(\text{aq})}$ , 160 mM for  $\text{NaClO}_{2(\text{aq})}$ , 0.02 M for  $\text{H}^{1+}_{(\text{aq})}$ , and 9.2 M for  $\text{Cl}^{-}_{(\text{aq})}$ .

better understanding on what reaction pathways are responsible for the oxidation of  $\text{Np}^{4+}_{(\text{aq})}$  by  $\text{NaClO}_{2(\text{aq})}$ . On an applied level, these types of studies – alongside the data reported herein – will impact waste stewardship programs, production of pure isotopes, and touch on many processing campaigns associated with actinide reliant technologies.

## Methods

### General considerations

**CAUTION!** Neptunium-237 [ $^{237}\text{Np}$ ,  $t_{1/2} = 2.144(6) \times 10^6 \text{ y}$ ]<sup>33</sup> and its daughter products constitute serious health threats because of radioactive decay. Hence, all experiments that involved manipulation of these radionuclides were conducted in a radiological buffer area that contained HEPA filtered hoods, continuous air monitors, negative pressure gloveboxes, and monitoring equipment appropriate for  $\alpha$ -,  $\beta$ -, and  $\gamma$ -particle detection. Entrance into the laboratory space was controlled with a hand and foot monitoring instrument for  $\alpha$ -,  $\beta$ -, and  $\gamma$ -emitting isotopes and a full body personal contamination monitoring station.

Aqueous hydrochloric acid [ $\text{HCl}_{(\text{aq})}$ , Fisher Scientific, Optima<sup>®</sup> grade] and sodium chlorite ( $\text{NaClO}_2$ , 80%, Sigma Aldrich) was obtained commercially and used as received. Water ( $\text{H}_2\text{O}$ ) was deionized, passed through a Barnstead water purification system to achieve resistivity of 18  $\text{M}\Omega \text{ cm}$ , and purified further *via* distillation using a Teflon distillation apparatus. Chemically pure and oxidation state pure stock solutions of  $\text{Np}^{4+}_{(\text{aq})}$  were obtained from  $^{237}\text{Np}$  samples that had been recovered from previous experimental campaigns, as described previously.<sup>59</sup> In general, these  $^{237}\text{Np}$  samples were combined and processed using a series of precipitations, valence adjustments, and ion exchange chromatography. The end result was a chemically pure, emerald green stock solution that contained  $\text{Np}^{4+}_{(\text{aq})}$  (66 mM) in  $\text{HCl}_{(\text{aq})}$  (1 M).<sup>59</sup>

### UV-Vis spectroscopy

All UV-vis spectroscopy measurements were conducted using a Stellar Net EXTENDED RANGE NIR Spectrometer and/or Ocean Insight NIR Quest Spectrometer. To mitigate hazards associated with making optical measurements on radioactive samples, the cuvette holder was housed within a HEPA filtered chemical fume hood. This holder was connected to the UV-vis spectrometer using fiber optics and the neptunium samples were contained within the fume hood in screw top quartz cuvettes (Starna Scientific). Neptunium concentrations were determined by monitoring the following absorption peaks: 10 400  $\text{cm}^{-1}$  (960 nm) and/or 8700  $\text{cm}^{-1}$  (1150 nm) for  $\text{Np}^{4+}_{(\text{aq})}$ , 10 200  $\text{cm}^{-1}$  (980 nm) and/or 9100  $\text{cm}^{-1}$  (1100 nm) for  $\text{NpO}_2^{1+}_{(\text{aq})}$ , 8200  $\text{cm}^{-1}$  (1220 nm) and/or 7900  $\text{cm}^{-1}$  (1270 nm) for  $\text{NpO}_2^{2+}_{(\text{aq})}$ .

### Data acquisition and analysis

Before handling the Np stock, a reference UV-vis spectra was initially obtained from pure  $\text{HCl}_{(\text{aq})}$  (0, 6, 7, or 8 M) or  $\text{LiCl}_{(\text{aq})}$  (10 M) solutions prior to assaying  $\text{Np}^{4+}_{(\text{aq})}$  and  $\text{NaClO}_{2(\text{aq})}$  containing solutions. Data from  $\text{Np}^{4+}_{(\text{aq})}$  and  $\text{NaClO}_{2(\text{aq})}$  reactions were background subtracted by setting the intensity at 9520  $\text{cm}^{-1}$  (1050 nm) to zero. Reaction rate constants were determined using the systematic numerical procedure by which the concentration of one variable was varied (*e.g.* ingoing  $\text{Np}^{4+}_{(\text{aq})}$ ) whilst the other three variables were held constant [in this scenario, ingoing  $\text{NaClO}_{2(\text{aq})}$ ,  $\text{H}^{1+}_{(\text{aq})}$ , and  $\text{Cl}^{-}_{(\text{aq})}$ ].<sup>71,72</sup> The linear initial reaction rate was taken to be the rate by which  $\text{Np}^{4+}_{(\text{aq})}$  reacted (after mixing) by using linear regression. Errors for the analysis of reactions rates were calculated through propagation of error in measurements of ingoing  $\text{NaClO}_{2(\text{aq})}$  mass, pipetting, and volumetric flasks. An example calculation may be found in the ESI.† Molar extinction coefficients were calculated using the Beer-Lambert Law.<sup>73</sup>



### Using dilute aqueous sodium chlorite solutions, $\text{NaClO}_{2(\text{aq})}$ , to generate oxidation state pure solutions of neptunyl(v), $\text{NpO}_2^{1+}(\text{aq})$

In a fume hood and with no attempt to exclude air and moisture, an aliquot (15  $\mu\text{L}$ ; 0.2 mg, 1  $\mu\text{mol}$  of  $\text{Np}^{4+}$ ) of the aforementioned  $\text{Np}^{4+}(\text{aq})$  stock solution (66 mM  $\text{Np}^{4+}$ ) in  $\text{HCl}_{(\text{aq})}$  (1 M) was added to a screw-top quartz cuvette charged with  $\text{H}_2\text{O}$  (600  $\mu\text{L}$ ). The  $\text{Np}^{4+}(\text{aq})$  concentration for this new 615  $\mu\text{L}$  solution was now 1.6 mM. Meanwhile, in a separate beaker, a dilute solution of  $\text{NaClO}_{2(\text{aq})}$  (6.9 mM) was prepared by dissolving  $\text{NaClO}_{2(\text{s})}$  (0.0389 g, 80%, 0.344 mmol) in  $\text{H}_2\text{O}$  (50 mL). An aliquot (600  $\mu\text{L}$ , 4  $\mu\text{mol}$ ) from this  $\text{NaClO}_{2(\text{aq})}$  solution was added all at once to the  $\text{Np}^{4+}(\text{aq})$  containing cuvette. **CAUTION!** The combination of  $\text{HCl}_{(\text{aq})}$  and  $\text{NaClO}_{2(\text{aq})}$  is vigorous, can bubble, and generates toxic gases (like  $\text{ClO}_2$ ). To mitigate these hazards, we implemented the following engineering and administrative controls. First, the aliquot of the  $\text{NaClO}_{2(\text{aq})}$  was added slowly (over the course of 5 seconds to the  $\text{Np}^{4+}(\text{aq})$  containing cuvette). Second, the manipulation was carried out in a chemical fume hood, behind glass shielding, to guard the worker from evolved gases. Third, secondary containment was used to avoid spreading  $^{237}\text{Np}$ -contamination, which can result from potential splattering. Finally, the screw top cuvette was only loosely capped, to avoid pressurization. The reagents were mixed by pumping the solution (from bottom to top) with a transfer pipette at least 4 times. This mixing action marked zero time for the reaction. It is relevant to note that the concentrations at the zero time for this solution (1.2 mL) were 0.81 mM for  $\text{Np}_{(\text{aq})}$ , 3.4 mM for  $\text{NaClO}_{2(\text{aq})}$ , and 0.01 M for  $\text{HCl}_{(\text{aq})}$ . The cuvette was inserted into the UV-vis holder, the apparatus covered (to exclude ambient light), and the spectra were collected repeatedly over the course of a 5-day monitoring process.

### Using concentrated aqueous sodium chlorite solutions, $\text{NaClO}_{2(\text{aq})}$ , to generate oxidation state pure solutions of neptunyl(vi), $\text{NpO}_2^{2+}(\text{aq})$

In a fume hood and with no attempt to exclude air and moisture, oxidation state pure solutions of  $\text{NpO}_2^{2+}(\text{aq})$  were prepared using an adaptation of the procedure described in the preceding section titled, "Using Dilute Aqueous Sodium Chlorite Solutions,  $\text{NaClO}_{2(\text{aq})}$ , to Generate Oxidation State Pure Solutions of Neptunyl(v)." The major difference between these procedures was associated with the amount of added  $\text{NaClO}_{2(\text{aq})}$  used to oxidize  $\text{Np}^{4+}(\text{aq})$ . Toward this end, a more concentrated solution of  $\text{NaClO}_{2(\text{aq})}$  (2.6 M) was prepared by dissolving  $\text{NaClO}_{2(\text{s})}$  (298 mg, 2.6 mmol) in  $\text{H}_2\text{O}$  (1 mL). Then, the  $\text{NaClO}_{2(\text{aq})}$  (40  $\mu\text{L}$ , 2.6 M, 104  $\mu\text{mol}$ ) and  $\text{Np}^{4+}(\text{aq})$  (615  $\mu\text{L}$ , 1.6 mM, 1  $\mu\text{mol}$ ) solutions were combined in a cuvette. **CAUTION!** See precautions described above that were implemented to mitigate this hazardous combination activity. The reagents were mixed by pumping the solution (from bottom to top) with a transfer pipette at least 4 times. This mixing action marked zero time for the reaction. The reagent concentrations at the zero time for this solution (655  $\mu\text{L}$ ) were 1.5 mM for  $\text{Np}^{4+}(\text{aq})$ , 160 mM for  $\text{NaClO}_{2(\text{aq})}$  (approximate), and 0.02 M for  $\text{HCl}_{(\text{aq})}$ . The cuvette

was inserted into the UV-vis holder, the apparatus covered (to exclude ambient light), and the spectra were collected repeatedly over the course of a 5 day monitoring period.

### Evaluating how four variables impacted oxidation of neptunium by sodium chlorite; $\text{Np}^{4+}(\text{aq}) + \text{NaClO}_{2(\text{aq})}$

In a fume hood and with no attempt to exclude air and moisture, oxidation of  $\text{Np}^{4+}(\text{aq})$  by  $\text{NaClO}_{2(\text{aq})}$  was investigated in  $\text{HCl}$  as a function of the following four variables, (1)  $\text{Np}^{4+}(\text{aq})$  reagent concentration, (2)  $\text{NaClO}_{2(\text{aq})}$  oxidant concentration, (3)  $\text{Cl}^{-}(\text{aq})$  complexant concentration, and (4)  $\text{H}^{+}(\text{aq})$  concentration. This was achieved by modifying the experimental conditions described above in the antecedent section titled, "Using Dilute Aqueous Sodium Chlorite Solutions,  $\text{NaClO}_{2(\text{aq})}$ , to Generate Oxidation State Pure Solutions of Neptunyl(v)." See Table 1 for a summary of those tested parameters and the primary neptunium oxidation states that formed within 1 h and within 24 h. These solutions were prepared by modifying the concentration of the ingoing reagents, oxidant, complexing agents, and acid. The ingoing  $\text{Np}^{4+}(\text{aq})$  concentration was varied (1) by combining 15  $\mu\text{L}$  of the  $\text{Np}^{4+}(\text{aq})$  (66 mM) stock solution with the other reagents in a cuvette to generate a final solution that was dilute in  $\text{Np}^{4+}(\text{aq})$  (1.5 mM) or (2) by combining 30  $\mu\text{L}$  of the  $\text{Np}^{4+}(\text{aq})$  (66 mM) stock solution with the other reagents in a cuvette to generate a final solution that was highly concentrated in  $\text{Np}^{4+}(\text{aq})$  (3.1 mM). The ingoing  $\text{NaClO}_{2(\text{aq})}$  concentration was varied (1) by combining 40  $\mu\text{L}$  of a  $\text{NaClO}_{2(\text{aq})}$  (520 mM) stock solution with the other reagents in a cuvette to generate a final solution that was intermediately concentrated in  $\text{NaClO}_{2(\text{aq})}$  (31 or 32 mM) or (2) by combining 40  $\mu\text{L}$  of a  $\text{NaClO}_{2(\text{aq})}$  (2.6 M) stock solution with the other reagents in a cuvette to generate a final solution that was highly concentrated in  $\text{NaClO}_{2(\text{aq})}$  (160 mM). The ingoing  $\text{HCl}_{(\text{aq})}$  concentration was varied (1) by combining either 600  $\mu\text{L}$  of an  $\text{HCl}_{(\text{aq})}$  (6 to 8 M) stock solution with the other reagents in a cuvette to generate a final solution that was more concentrated in  $\text{HCl}_{(\text{aq})}$  (5.3 to 7.1 M) or (2) by adding an aliquot (15 or 30  $\mu\text{L}$ , described above) from the neptunium stock solution [1 M  $\text{HCl}_{(\text{aq})}$ ] to a cuvette containing Teflon distilled  $\text{H}_2\text{O}$  (600  $\mu\text{L}$ ). This gave final volumes and concentrations of either 615  $\mu\text{L}$  at 0.02 M  $\text{HCl}_{(\text{aq})}$  or 630  $\mu\text{L}$  at 0.05 M  $\text{HCl}_{(\text{aq})}$ . Then,  $\text{NaClO}_{2(\text{aq})}$  was added. These dilute  $\text{HCl}_{(\text{aq})}$  solutions also served as low ionic strength solutions, relatively speaking. High ionic strength solutions were prepared by combining 600  $\mu\text{L}$  of a  $\text{LiCl}_{(\text{aq})}$  (10 M) stock solution with the other reagents in a cuvette to generate a final solution that was 9.0 to 9.2 M in  $\text{LiCl}_{(\text{aq})}$ .

## Conflicts of interest

There are no conflicts to declare.

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