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## {Np<sub>38</sub>} clusters: the missing link in the largest poly-oxo cluster series of tetravalent actinides†

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**Two poly-oxo cluster complexes of tetravalent neptunium (Np<sup>IV</sup>), Np<sub>38</sub>O<sub>56</sub>Cl<sub>18</sub>(bz)<sub>24</sub>(THF)<sub>8</sub>·nTHF and Np<sub>38</sub>O<sub>56</sub>Cl<sub>42</sub>(ipa)<sub>20</sub>·mipa (bz = benzoate, THF = tetrahydrofuran, and ipa = isopropanol), were obtained via solvothermal synthesis and structurally characterised by single-crystal X-ray diffraction. The {Np<sub>38</sub>} clusters are comparable to the analogous {U<sub>38</sub>} and {Pu<sub>38</sub>} motifs, filling the gap in this largest poly-oxo cluster series of tetravalent actinides.**

Owing to its diversity as well as its technological importance in the nuclear industry, the polymer and cluster chemistry of actinides (An) has been flourishing since the last decade.<sup>1–5</sup> In particular, the poly-oxo/hydroxo polymer and cluster complexes of tetravalent actinides (An<sup>IV</sup>) have received considerable attention,<sup>6–20</sup> as the formation of these complexes stems primarily from their hydrolysis<sup>21–24</sup> and, hence, it would have significant implications for the natural and engineered aqueous systems associated with the nuclear industry.<sup>25</sup> The largest poly-oxo An<sup>IV</sup> cluster reported thus far is the {An<sub>38</sub>} complex, consisting of 38 An<sup>IV</sup> centres bridged by 56 oxygens. This type of cluster complex has been synthesised and characterised for U<sup>IV</sup><sup>9,26</sup> and Pu<sup>IV</sup>,<sup>13,15</sup> and this cluster motif could be stable in solution in a colloidal form.<sup>13,15,27</sup> In the periodic table, neptunium (<sup>93</sup>Np) is present between <sup>92</sup>U and <sup>94</sup>Pu. Despite its importance in nuclear fuel reprocessing and radioactive waste management,<sup>25</sup> the research on Np always lags behind those on U and Pu,<sup>28</sup> which is also the case for the chemistry of poly-oxo/hydroxo polymers and clusters. In order to fill the gap between the {U<sub>38</sub>} and {Pu<sub>38</sub>} clusters and to complete a series of these largest poly-oxo An<sup>IV</sup>

cluster complexes, we report herein the synthesis and characterisation of the first {Np<sub>38</sub>} cluster complexes, which are also the largest single molecules of Np reported thus far.

The first {Np<sub>38</sub>} complex, Np<sub>38</sub>O<sub>56</sub>Cl<sub>18</sub>(bz)<sub>24</sub>(THF)<sub>8</sub>·nTHF‡ (1, bz = benzoate, THF = tetrahydrofuran), was obtained via solvothermal synthesis with NpCl<sub>4</sub>, benzoic acid, THF and deionised water (Sections 1 and 2 in the ESI†), which is a slight modification of the synthesis of the {U<sub>38</sub>} complex.<sup>9</sup> After 1 day of heating at 400 K, purple/brown crystals with an octahedral shape were deposited (Fig. 1, left). Single-crystal X-ray diffraction (SC-XRD) measurements on the obtained crystals reveal the discrete {Np<sub>38</sub>} motif shown in Fig. 2a. The second {Np<sub>38</sub>} complex, Np<sub>38</sub>O<sub>56</sub>Cl<sub>42</sub>(ipa)<sub>20</sub>·mipa‡ (2, ipa = isopropanol), was synthesised solvothermally from a mixture of NpCl<sub>4</sub> and benzoic acid in isopropanol (Sections 1–3 in the ESI†). Heating the sample mixture at 370 K for 3 days resulted in the formation of purple/brown crystals with an octahedral shape (Fig. 1, right). The complex structure determined by SC-XRD shows another {Np<sub>38</sub>} motif, as illustrated in Fig. 2b.

Both compounds 1 and 2 crystallise in the tetragonal space group *I*4/*m* (Table S1 in the ESI†), which is the same as that observed for the analogous {U<sub>38</sub>} clusters U<sub>38</sub>O<sub>56</sub>Cl<sub>18</sub>(bz)<sub>24</sub>(THF)<sub>8</sub>·8THF<sup>9</sup> and U<sub>38</sub>O<sub>56</sub>Cl<sub>42</sub>(H<sub>2</sub>O)<sub>20</sub>·xipa.<sup>26</sup> Both the {Np<sub>38</sub>} clusters are composed of a {Np<sub>14</sub>} core (light green polyhedra in Fig. 2) surrounded by six tetrานuclear {Np<sub>4</sub>} subunits (blue and dark

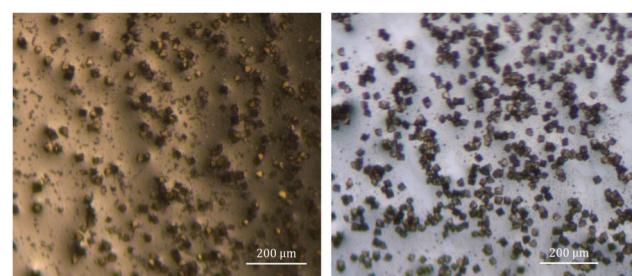


Fig. 1 Optical micrographs of single crystals of the {Np<sub>38</sub>} compounds synthesised from THF (1, left) and isopropanol (2, right).

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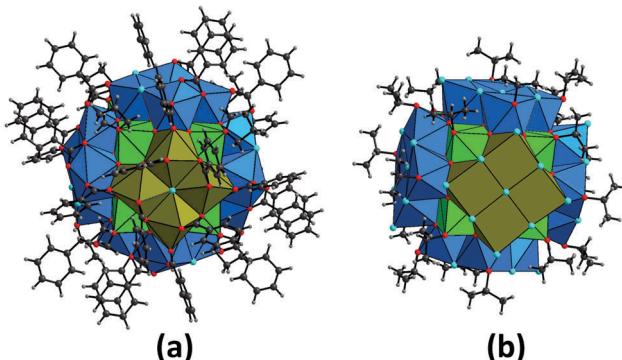


Fig. 2 Molecular structures of the two  $\{Np_{38}\}$  clusters  $Np_{38}O_{56}Cl_{18}(bz)_{24}(THF)_8$  (**1**, a) and  $Np_{38}O_{56}Cl_{42}(ipa)_{20}$  (**2**, b).<sup>§</sup> The coordination geometry around neptunium atoms (as  $Np^{4+}$ ) is illustrated with coloured polyhedra, whilst carbon (black), chlorine (light blue), hydrogen (grey) and oxygen (red) atoms are shown as balls. The cubic  $\{Np_{14}\}$  cores are depicted with light green, whilst the exterior  $\{Np_4\}^{tetra}$  and  $\{Np_4\}^m$  subunits adhering to the  $\{Np_{14}\}$  core are illustrated with blue and dark brown, respectively.

brown polyhedra in Fig. 2). In the  $\{Np_{14}\}$  core, three crystallographically distinct Np atoms (Np1, Np2 and Np3) are packed in a dense manner, forming a primitive unit of the fluorite structure (Fig. S1 in the ESI†) which is the basis of bulk  $NpO_2$ .<sup>29,30</sup> The eight corners of the cubic  $\{Np_{14}\}$  core are all occupied by the oxygen atoms of THF- or isopropanol molecules for **1** and **2**, respectively (Fig. S1 in the ESI†). Each face of the cubic  $\{Np_{14}\}$  core is further decorated with the  $\{Np_4\}$  subunits, where there is a significant difference between **1** and **2**. In both the  $\{Np_{38}\}$  compounds, two types of  $\{Np_4\}$  subunits appear. The first type (referred to as “ $\{Np_4\}^{tetra}$ ” hereafter) appears four times by symmetry operation with respect to the tetragonal axis, whilst the second type (referred to as “ $\{Np_4\}^m$ ” hereafter) appears twice around the  $\{Np_{14}\}$  core as a result of the perpendicular mirror plane operation. The coordination polyhedra of the Np atoms in the inner  $\{Np_{14}\}$  core are cubic, whilst those in the exterior  $\{Np_4\}$  subunits are square antiprismatic polyhedra (Fig. 3). This results in two different types of oxo-bridging groups in the  $\{Np_{38}\}$  clusters;  $\mu_4$ -O with Np-O distances of 2.294(9)–2.408(9) and 2.28(4)–2.42(4) Å for **1** and **2**, respectively, and  $\mu_3$ -O with Np-O distances of 2.203(9)–2.335(10) and 2.15(4)–2.305(3) Å for **1** and **2**, respectively. These Np-O distances are comparable to those in the fluorite-based  $NpO_2$  structure (2.353–2.354 Å),<sup>29,30</sup> suggesting that the Np-O framework in the  $\{Np_{38}\}$  clusters could also be comparable to that in the bulk  $NpO_2$ .

In compound **1**, the  $\{Np_4\}^{tetra}$  subunit (blue polyhedra in Fig. 2a and 3a) is composed of three distinct Np centres (Np4, Np5 and Np6). The four Np atoms are linked *via* either  $\mu_2/\mu_4$ -Cl atoms or bidentately coordinating the carboxylate groups of bz, forming a ninefold coordination geometry with distorted trideca- and undecahedra around Np4 and Np5, respectively, and an eightfold geometry with a distorted nonahedron around Np6 (Fig. 3a). Each  $\{Np_4\}^{tetra}$  subunit adheres to one side of the cubic  $\{Np_{14}\}$  core *via* twelve O atoms facing the core, covering the four sides of the  $\{Np_{14}\}$  cube. On the other hand, the  $\{Np_4\}^m$  subunit (dark brown polyhedra in Fig. 2a and 3b) is composed of a single type of a distinct Np centre (Np7). The four Np atoms in the

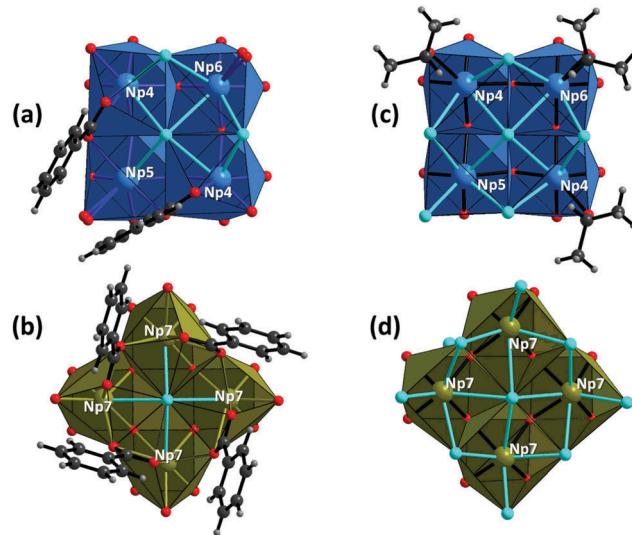


Fig. 3 Top-side views of the  $\{Np_4\}$  subunits in **1** and **2**; the  $\{Np_4\}^{tetra}$  and  $\{Np_4\}^m$  subunits in **1** (a and b, respectively) and those in **2** (c and d, respectively).<sup>§</sup> All these subunits contain unique  $\mu_4$ -bridging Cl atoms. The coordination geometry around neptunium atoms (as  $Np^{4+}$ ) is illustrated with blue and dark brown polyhedra ( $\{Np_4\}^{tetra}$  and  $\{Np_4\}^m$ , respectively), whilst carbon (black), chlorine (light blue), hydrogen (grey) and oxygen (red) atoms are shown as balls.

$\{Np_4\}^m$  subunit are linked *via* one  $\mu_4$ -Cl atom and four carboxylate groups of bz, forming a ninefold coordination geometry with a dodecahedron around the Np7 atoms (Fig. 3b). This  $\{Np_4\}^m$  subunit covered the rest of the two sides of the  $\{Np_{14}\}$  cube, and is further connected with the  $\{Np_4\}^{tetra}$  subunit by the bidentate carboxylate groups of bz to overlay the  $\{Np_{14}\}$  core (Fig. 2a).

Despite the presence of benzoic acid in the initial sample solution, no benzoate groups are involved in the formation of the  $\{Np_{38}\}$  cluster in **2**, which is in contrast to **1**. The exterior  $\{Np_4\}$  subunits in **2** are composed of the O atoms of isopropanol molecules,  $Cl^-$  ions and the  $O_{oxo}$  groups shared with the  $\{Np_{14}\}$  core. The  $\{Np_4\}^{tetra}$  subunit in **2** is composed of three distinct Np centres (Np4, Np5 and Np6), all of which form an eightfold square antiprismatic geometry (Fig. 3c). In the  $\{Np_4\}^m$  subunit of **2**, there is only a single distinct Np centre (Np7) which is surrounded by four  $O_{oxo}$  groups shared with the  $\{Np_{14}\}$  core and four  $Cl^-$  ions, eventually forming an eightfold square antiprismatic geometry as well (Fig. 3d). In a similar manner to the arrangement in **1**, the six  $\{Np_4\}$  subunits decorate the exterior of the cubic  $\{Np_{14}\}$  core to form an  $\{Np_{38}\}$  cluster.

The chemical formulae of  $[Np_{38}O_{56}Cl_{18}(bz)_{24}(THF)_8]$  (**1**) and  $[Np_{38}O_{56}Cl_{42}(ipa)_{20}]$  (**2**) derived from SC-XRD both exhibit an excess negative charge of  $-2$ , assuming that the Np atoms in the clusters are all tetravalent (*i.e.*  $Np^{4+}$ ). No cationic species are identified in the crystal packing of **1** and **2** either, indicating that the  $\{Np_{38}\}$  cluster units should be neutral. This discrepancy can be, however, explained by the partial replacement of chloride anions ( $Cl^-$ ) with neutral water molecules ( $H_2O$ ) in the  $\{Np_4\}$  subunits and/or the replacement of the  $\mu_3$ -oxo groups with the hydroxo ones in the inner  $\{Np_{14}\}$  core. Further detailed discussions are provided in the ESI† (Section 3).

Albeit a series of  $\{An_{38}\}$  clusters (*i.e.*  $\{U_{38}\}$ ,<sup>9,26</sup>  $\{Np_{38}\}$ ) in this study, and  $\{Pu_{38}\}$ <sup>13,15</sup>) were successful characterised, their formation mechanism has been unrevealed even to date. All the  $\{An_{38}\}$  clusters characterised thus far are composed of an inner  $\{An_{14}\}$  core decorated with six exterior  $\{An_4\}$  subunits. Given this construction manner, it is reasonable to infer that the formation of the primary  $\{An_{14}\}$  core is followed by the attachment of the  $\{An_4\}$  subunits to the surface of the cubic  $\{An_{14}\}$  core. Owing to their strong olation/oxolation tendency,<sup>31</sup>  $Al_{(IV)}$  can form a variety of oligomer species<sup>2</sup> including soluble  $\mu_2$ -hydroxo-bridged dinuclear species,<sup>7,17,32</sup> which could further evolve into hexanuclear species *via* hydrolysis.<sup>6,10,11,14</sup> A similar evolution of such oligomer species was also observed for  $Ce_{(IV)}$ ,<sup>33,34</sup> a chemical analogue of  $Al_{(IV)}$ . The union of di- and hexanuclear species could also lead to the formation of  $\{An_{12}\}$  complexes.<sup>8</sup> However, to the best of our knowledge, no  $\{An_{14}\}$  complexes have been characterised for  $Al_{(IV)}$  thus far, indicating that the  $\{An_{14}\}$  motif is probably not stable at least as a discrete species. In contrast, several discrete  $\{An_4\}$  complexes have been characterised for  $Th_{(IV)}$ <sup>35</sup> and  $U_{(IV)}$ ,<sup>36-43</sup> seven of which exhibit a planar tetranuclear arrangement that is structurally comparable with the  $\{An_4\}$  subunits in the  $\{An_{38}\}$  clusters (Section 4 in the ESI†). A striking difference between the  $\{An_4\}$  subunits and the discrete tetranuclear  $Al_{(IV)}$  complexes is, however, the presence/absence of  $\mu_4$ -bridging groups. As illustrated in Fig. 3, there are unique  $\mu_4$ -bridging Cl atoms situated at the centre of the  $\{Np_4\}$  subunits, which are observed for the whole series of  $\{An_{38}\}$  clusters. On the other hand, none of the discrete tetranuclear  $Al_{(IV)}$  complexes contains  $\mu_4$ -bridging groups (Table S2 and Fig. S2 in the ESI†), except one  $Th_{(IV)}$  complex<sup>35</sup> containing a  $\mu_4$ -bridging oxo group to form a unique non-planar boat conformation arrangement (Fig. S3 in the ESI†). Instead, these discrete tetranuclear complexes are composed of  $\mu_2$ - and  $\mu_3$ -bridging groups (Table S2 and Fig. S3 in the ESI†), suggesting that the planar  $\{An_4\}$  motif with a  $\mu_4$ -bridging group may also not be stable as a discrete species. Given all these facts, the formation process of the  $\{An_{38}\}$  clusters could be surmised as follows:

(1) the primitive but transient  $\{An_{14}\}$  core is formed *via* the olation/oxolation of precursor species (*e.g.* dinuclear and/or hexanuclear species),

(2) owing to an extremely large negative charge of the primitive  $\{An_{14}\}$  core (−72 based on the formula  $Al_{14}O_{64}$  (Fig. S1 in the ESI†)), cationic species in the system (*i.e.*  $Al^{4+}$ ) are further attracted to the surface of the  $\{An_{14}\}$  core immediately after the formation of the primitive core, forming dense  $\{An_4\}$  subunits containing  $\mu_4$ -bridging groups on the surface, and

(3) the attachment of six  $\{An_4\}$  subunits on each face of the cubic  $\{An_{14}\}$  core neutralises the negative charge of the  $\{An_{14}\}$  core, eventually stabilising the whole unit as  $\{An_{38}\}$ .

Based on this hypothetical formation process, it is obvious that the faces of the primitive  $\{An_{14}\}$  core provide a sort of “substrate” for the formation of unique planar  $\mu_4$ -bridged  $\{An_4\}$  subunits. In other words, such a planar  $\mu_4$ -bridged  $\{An_4\}$  motif could be formed and stabilised only when an appropriate substrate exists in the system.

As previously mentioned, the inner  $\{An_{14}\}$  core in the  $\{An_{38}\}$  cluster is a primitive unit of the fluorite-based  $AlO_2$  structure.

The  $\{An_4\}$  subunits in the cluster also show a closer structural conformity to the bulk  $AlO_2$  structure as compared with the discrete tetranuclear  $Al_{(IV)}$  complexes (Fig. S4 in Section 4, ESI†). All these facts indicate that the poly-oxo  $\{An_{38}\}$  clusters can be the origin of the formation of  $AlO_2$  in solution. As a matter of fact, the  $\{An_{38}\}$  clusters can easily evolve into bulk  $AlO_2$  by moderate hydrolysis.<sup>9,26</sup> Given the electroneutrality of the  $\{An_{38}\}$  clusters, the  $\{An_{38}\}$  motif will no longer attract additional metal cations (*i.e.*  $Al^{4+}$ ) to form larger clusters. Hence, the  $\{An_{38}\}$  unit is probably the largest poly-oxo  $Al_{(IV)}$  cluster stabilised in solution, which is also the limiting point that defines the border between discrete poly-oxo oligomer/cluster complexes and insoluble bulk  $AlO_2$ , including colloidal species or aggregates.

In summary, this study succeeded in synthesising and structurally characterising the two poly-oxo  $Np_{(IV)}$  cluster complexes **1** and **2** which are composed of 38  $Np$  atoms. The characterised  $\{Np_{38}\}$  clusters are the largest discrete  $Np$  complexes reported thus far, filling the gap between the already reported  $\{U_{38}\}$  and  $\{Pu_{38}\}$  clusters to complete a series of  $\{An_{38}\}$  cluster complexes. Although hexavalent  $Al$  cations (as actinyl( $VI$ ) cations:  $AlO_2^{2+}$ ) can form poly-oxo clusters larger than 100-mer based on the cage structure which is not directly comparable with open clusters,<sup>3</sup> the  $\{An_{38}\}$  motif is probably the largest open cluster unit for the poly-oxo  $Al_{(IV)}$  complexes that can be stabilised as a discrete species in solution. A series of poly-oxo  $\{An_{38}\}$  clusters ( $Al = U$ ,  $Np$  and  $Pu$ ) exhibits a significant structural similarity, suggesting a similar olation/oxolation (*i.e.* hydrolysis) behaviour amongst these lighter  $Al_{(IV)}$  series. A comparison of the series of  $\{An_{38}\}$  clusters with the large poly-oxo clusters of other metals (*e.g.*  $Ti^{44}$  or  $Bi^{45}$ ) will further highlight the peculiarity or generality of the chemistry of  $Al_{(IV)}$  on the periodic table, particularly in terms of hydrolysis and coordination chemistry.

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## Conflicts of interest

There are no conflicts to declare.

## Notes and references

‡ The Fourier maps of electronic density residues on the collected SC-XRD data show diffused peaks in the void between the  $\{Np_{38}\}$  units. These peaks probably correspond to neutral solvent molecules (*i.e.* THF in **1** and isopropanol in **2**) intercalated in the crystal structures, as observed in the analogous  $\{U_{38}\}$  compounds.<sup>9,26</sup> However, owing to the limited quality of the collected SC-XRD data, it is difficult to determine the positions of these intercalated solvent molecules. For this reason, the number of solvent molecules is not specified in the chemical formulae of **1** and **2**.

§ The carbon atoms of some benzoate (bz) molecules in **1** show disorder, statistically populating at two close sites with equivalent occupancy. This disorder is, however, not illustrated in Fig. 2a and 3a for clarity.

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