Two poly-oxo cluster complexes of tetravalent neptunium (Np(IV)), \( \text{Np}_{38} \text{O}_{56} \text{Cl}_{42} \text{bz}_{24}(\text{THF})_{8} \text{nTHF} \) and \( \text{Np}_{38} \text{O}_{56} \text{Cl}_{42} \text{ipa}_{20} \text{mpia} \text{ (bz = benzoate, THF = tetrahydrofuran, and ipa = isopropanol), were obtained via solvothermal synthesis and structurally characterised by single-crystal X-ray diffraction. The \( \text{Np}_{38} \) clusters are comparable to the analogous \( \text{U}_{38} \) and \( \text{Pu}_{38} \) 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.\(^1\)\(^-\)\(^5\) In particular, the poly-oxo/hydroxo polymer and cluster complexes of tetravalent actinides (An(IV)) have received considerable attention,\(^6\)\(^-\)\(^{20}\) as the formation of these complexes stems primarily from their hydrolysis\(^21\)\(^-\)\(^{24}\) and, hence, it would have significant implications for the natural and engineered aqueous systems associated with the nuclear industry.\(^2\)\(^5\) The largest poly-oxo An(IV) cluster reported thus far is the \( \text{An}_{38} \) complex, consisting of 38 An(IV) centres bridged by 56 oxygens. This type of cluster complex has been synthesised and characterised for U(IV)\(^9\)\(^,\)\(^{26}\) and Pu(IV),\(^1\)\(^3\)\(^,\)\(^1\)\(^5\) and this cluster motif could be stable in solution in a colloidal form.\(^1\)\(^3\)\(^,\)\(^1\)\(^5\)\(^,\)\(^2\)\(^1\)\(^\)\(^2\)\(^4\) In the periodic table, neptunium (\( 93\text{Np} \)) is present and this cluster motif could be stable in solution in a colloidal form.\(^1\)\(^3\)\(^,\)\(^1\)\(^5\)\(^,\)\(^2\)\(^1\)\(^\)\(^2\)\(^4\) The \( \text{Np}_{38} \) motif shown in Fig. 2a. The second \( \text{Np}_{38} \) complex, \( \text{Np}_{38} \text{O}_{56} \text{Cl}_{42} \text{ipa}_{20} \text{mpia} \) (2, ipa = isopropanol), was synthesised solvothermally from a mixture of NpCl\(_4\), benzoic acid in THF and deionised water (Sections 1 and 2 in the ESI\(^\dagger\)). Heating the sample mixture at 370 K for 3 days resulted in the formation of 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 \( \text{Np}_{38} \) motif shown in Fig. 2a. The second \( \text{Np}_{38} \) complex, \( \text{Np}_{38} \text{O}_{56} \text{Cl}_{42} \text{ipa}_{20} \text{mpia} \) (2, ipa = isopropanol), was synthesised solvothermally from a mixture of NpCl\(_4\) and benzoic acid in isopropanol (Sections 1–3 in the ESI\(^\dagger\)). 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 \( \text{Np}_{38} \) motif, as illustrated in Fig. 2b.

Both compounds 1 and 2 crystallise in the tetragonal space group \( \text{I} / \text{m} / \text{C} \) (Table S1 in the ESI\(^\dagger\)), which is the same as that observed for the analogous \( \text{U}_{38} \) clusters \( \text{U}_{38} \text{O}_{56} \text{Cl}_{42} \text{bz}_{24}(\text{THF})_{8} 8\text{THF} \) and \( \text{U}_{38} \text{O}_{56} \text{Cl}_{42} (\text{H}_2\text{O})_{2} (\text{ipa})_{20} \text{ipa.} \(^{26}\) Both the \( \text{Np}_{38} \) clusters are composed of a \( \text{Np}_{34} \) core (light green polyhedra in Fig. 2) surrounded by six tetranuclear \( \text{Np}_4 \) subunits (blue and dark

Fig. 1 Optical micrographs of single crystals of the \( \text{Np}_{38} \) compounds synthesised from THF (1, left) and isopropanol (2, right).
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 NpO2.29,30 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 \( \{Np_{18}\} \) compounds, two types of \( \{Np_{4}\} \) subunits appear. The first type (referred to as \( \{Np_{4}\}_\text{tert} \) hereafter) appears four times by symmetry operation with respect to the tetragonal axis, whilst the second type (referred to as \( \{Np_{4}\}_\text{m} \) 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 \( \{Np_{18}\} \) clusters; \( \mu_4-O \) with Np-O distances of 2.294(9)–2.408(9) and 2.28(4)–2.42(4) \( \AA \) for 1 and 2, respectively, and \( \mu_2-O \) with Np-O distances of 2.203(9)–2.335(10) and 2.15(4)–2.305(3) \( \AA \) for 1 and 2, respectively. These Np-O distances are comparable to those in the \( \{Np_{38}\} \) clusters (2.28(4)–2.42(4) \( \AA \), respectively), and this results in two different types of oxo-bridging groups in the \( \{Np_{38}\} \) clusters (2.28(4)–2.42(4) \( \AA \), respectively). These Np–O distances are comparable to those in the bulk NpO2.

In compound 1, the \( \{Np_{4}\}_\text{tert} \) 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 \( \mu_2/\mu_4-Cl \) 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 \( \{Np_{4}\}_\text{tert} \) subunit adheres to one side of the cubic \( \{Np_{14}\} \) core \( \mu_2 \) twelve O atoms facing the core, covering the four sides of the \( \{Np_{14}\} \) cube. On the other hand, the \( \{Np_{4}\}_\text{m} \) 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 \( \{Np_{4}\}_\text{m} \) subunit are linked \( \mu_4 \) 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 \( \{Np_{4}\}_\text{m} \) subunit covered the rest of the two sides of the \( \{Np_{14}\} \) cube, and is further connected with the \( \{Np_{4}\}_\text{tert} \) 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 \( \{Np_{4}\}_\text{tert} \) 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 \( \{Np_{4}\}_\text{m} \) 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 \( [Np_{38}O_{56}Cl_{18}(bz)_2]_2(THF)_9 \) (1) and \( [Np_{38}O_{56}Cl_{18}(ipa)_{20}]_2 \) (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. Np4+). 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 (H2O) in the \( \{Np_{4}\} \) subunits and/or the replacement of the \( \mu_4-oxo \) 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 (\textit{i.e.} \{U_{38}\},9,26 \{Np_{38}\} in this study, and \{Pu_{38}\}13,15) 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,11 An(\textit{iv}) can form a variety of oligomer species\(^8\) including soluble \\(\mu_4\)-hydroxo-bridged dinuclear species,7,17,32 which could further evolve into hexanuclear species via hydrolysis.6,10,11,14 A similar evolution of such oligomer species was also observed for Ce(\textit{iv}),33,34 a chemical analogue of An(\textit{iv}). The union of di- and hexanuclear species could also lead to the formation of \{An_{38}\} complexes.8 However, to the best of our knowledge, no \{An_{14}\} complexes have been characterised for An(\textit{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(\textit{iv})35 and U(\textit{iv}),36–43 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\(^\dagger\)). A striking difference between the \{An_{38}\} subunits and the discrete tetranuclear An(\textit{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_{14}\} subunits, which are observed for the whole series of \{An_{38}\} clusters. On the other hand, none of the discrete tetranuclear An(\textit{iv}) complexes contains \\(\mu_4\)-bridging groups (Table S2 and Fig. S2 in the ESI\(^\dagger\)), except one Th(\textit{iv}) complex35 containing a \\(\mu_4\)-bridging oxo group to form a unique non-planar boat conformation arrangement (Fig. S3 in the ESI\(^\dagger\)). Instead, these discrete tetranuclear complexes are composed of \\(\mu_2\) and \\(\mu_3\)-bridging groups (Table S2 and Fig. S3 in the ESI\(^\dagger\)), 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 (\textit{e.g.} dinuclear and/or hexanuclear species),

(2) Owing to an extremely large negative charge of the primitive \{An_{14}\} core (\(\approx -72\) based on the formula \(\text{An}_{14}\text{O}_{34}\) (Fig. S1 in the ESI\(^\dagger\))), cationic species in the system (\textit{i.e.} An(\textit{iv})) are further attracted to the surface of the \{An_{14}\} core immediately after the formation of the primitive core, forming dense \{An_{14}\} 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 AnO\(_2\) structure.
