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# Triple-armed aliphatic tricarboxylic acids as sources of ligands for uranyl ion: influence of bridgehead functionalization†

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The two triple-armed species tris(2-carboxyethyl)nitromethane ( $H_3tce$ ) and tris(2-carboxyethyl)phosphine ( $H_3tcep$ ) have been used to synthesize seven uranyl ion complexes under (solvo)-hydrothermal conditions and in the presence of various structure-directing cations. The three carboxylate groups chelate three different cations and the nitro group is uncoordinated in  $[H_2NMe_2][UO_2(tce)] \cdot 3H_2O$  (**1**),  $[C(NH_2)_3][UO_2(tce)] \cdot 0.5H_2O$  (**2**) and  $[PPh_3Me][UO_2(tce)]$  (**3**), which crystallize as diperiodic coordination polymers with the **hcb** topology and minor variations in shape depending on the counterion. The two isomorphous complexes  $[UO_2(tce)M(bipy)_2][UO_2(tce)] \cdot 3H_2O$ , with  $M = Ni$  (**4**) or  $Cu$  (**5**) and  $bipy = 2,2'$ -bipyridine, display the same arrangement, with a  $M(bipy)_2^{2+}$  group bridging two adjoining carboxylate donors in one uranyl equatorial plane.  $[(UO_2)_2(tce)Cu(R,S-Me_6cyclam)] \cdot 2H_2O$  (**6**), where  $R,S-Me_6cyclam = 7(R),14(S)-5,5,7,12,14$ -hexamethyl-1,4,8,11-tetraazacyclotetradecane, is the only triperiodic framework in the series, with the **tcs** topology resulting from  $Cu^{II}$  pillaring diperiodic, uranyl-based **sql** networks.  $H_3tcep$  is oxidized *in situ* to give the phosphine oxide  $H_3tcepo$ , which is partially deprotonated in  $[UO_2(Htcepo)]$  (**7**); the phosphine oxide and the two carboxylate groups are coordinated, and the diperiodic, three-dimensional network formed has the point symbol  $\{6^6\}$  and the vertex symbol  $6_2 \cdot 6_2 \cdot 6_3 \cdot 6_6 \cdot 6_4 \cdot 6_4$ , with the rings involved in Hopf links formation. Only complexes **3** and **7** are significantly emissive in the solid state, with photoluminescence quantum yields of 9%, and the emission maxima positions are in agreement with the number of uranyl equatorial donors.

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## Introduction

Although tripodal ligands possessing a hydrogen bond donor bridgehead able to interact with one uranyl oxo group were proposed long ago for selective uranyl recognition through so-called stereognostic coordination,<sup>1</sup> and other tripodal ligands have been investigated for use in uranyl solvent extraction,<sup>2</sup> they are not the most obvious choice for the synthesis of uranyl-based coordination polymers of frameworks.<sup>3</sup> These known species are ligands with long and often rather rigid arms intended to be convergent about a single uranyl ion and thus to give mononuclear, discrete complexes; as such, they provide little prospect for use in uranyl-based coordination polymer synthesis. Of other ligands which can be viewed as tripodal due to their capacity to adopt a conformation

displaying three convergent functional groups, most are of small size, such as the conformationally restricted tricarboxylate derived from Kemp's triacid (*cis,cis*-1,3,5-trimethylcyclohexane-1,3,5-tricarboxylic acid), and this is a ligand known to give a wide range of discrete polynuclear or polymeric species.<sup>4</sup> A more commonly encountered, flexible tripodal ligand, with cation encapsulation ability and a potentially coordinating bridgehead, is nitrilotriacetate,<sup>5</sup> for which, besides cases in which the central nitrogen atom is protonated<sup>5a,b</sup> or coordinated to an additional nickel(II) cation,<sup>5c</sup> *ONO*-chelation of uranyl is known,<sup>5d</sup> with however the third carboxylate arm being diverted away so as to give a coordination polymer, the size of the tripodal cavity being definitely too small for uranyl ion encapsulation.<sup>1a</sup> A semi-rigid, tripodal tricarboxylate ligand based on the triazine platform has also been used to synthesize various cage-like or polymeric, diperiodic complexes,<sup>6</sup> and a very recent publication has shown that a large coordination cage can be obtained with a calix[3]arene-carboxylate tripod rigidified by uranyl ion coordination as a bridgehead.<sup>7</sup> Recently, several larger tripodal, zwitterionic tricarboxylates have been investigated,<sup>8</sup> and  $1,1',1''-[(2,4,6\text{-trimethylbenzene-1,3,5-triyl})\text{tris(methylene)}]\text{tris(pyridin-1-ium-4-carboxylate)}$  in particular

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was shown to retain its convergent shape when encapsulating a bromide ion, while a more irregular conformation yielded a triperiodic uranyl-based framework.<sup>8d</sup> To further explore these diverse indications of competition between encapsulation and polymer formation, we have now synthesized a series of seven complexes from the triple-armed species tris(2-carboxyethyl)nitromethane ( $H_3tce nm$ ; 4-(2'-hydroxycarbonylethyl)-4-nitroheptane-1,7-dioic acid; Scheme 1) and tris(2-carboxyethyl)phosphine ( $H_3tce p$ ; 4-(2'-hydroxycarbonylethyl)-4-phosphahexane-1,7-dioic acid), which have been characterized by their crystal structure and emission spectrum in the solid state. Under the solvo-hydrothermal conditions used for the syntheses, the phosphine group in  $H_3tce p$  is oxidized to give  $H_3tce po$  (Scheme 1), and the nitromethane and phosphine oxide bridgeheads appear to have very different effects on the charge of the bound ligand and the structures of the complexes formed, another instance of structural control through internal functionality of the bridging ligands.<sup>9</sup>

## Experimental

### Synthesis

**Caution!** Uranium is a radioactive and chemically toxic element, and uranium-containing samples must be handled with suitable care and protection. Small quantities of reagents and solvents were employed to minimize any potential hazards arising both from the presence of uranium and the use of pressurized vessels for the syntheses.

$[UO_2(NO_3)_2(H_2O)_2] \cdot 4H_2O$  (RP Normapur, 99%) and  $Ni(NO_3)_2 \cdot 6H_2O$  were purchased from Prolabo. Tris(2-carboxyethyl)nitromethane ( $H_3tce nm$ ), tris(2-carboxyethyl)phosphine hydrochloride ( $H_3tce p \cdot HCl$ ), and  $Cu(NO_3)_2 \cdot 2.5H_2O$  were from Sigma-Aldrich, 2,2'-bipyridine (bipy) was from Fluka, and guanidinium nitrate was from Alfa-Aesar. *R,S*-Me<sub>6</sub>cyclam (*meso* isomer of 7(*R*),14(*S*)-5,5,7,12,12,14-hexamethyl-1,4,8,11-tetraazacyclotetradecane) was prepared as described in the literature,<sup>10</sup> and 7(*R*),14(*S*)-5,5,7,12,12,14-hexamethyl-1,4,8,11-tetraazacyclotetradecane)copper(II) dinitrate,  $[Cu(R,S-Me_6cyclam)](NO_3)_2$ , was synthesized as previously reported.<sup>11</sup> Elemental analyses were performed by MEDAC Ltd. For all

syntheses, the solutions were placed in 10 mL tightly closed glass vessels (Pyrex culture tubes with SVL15 stoppers and Teflon-coated seals, provided by VWR) and heated at 140 °C in a sand bath (Harry Gestigkeit ST72). The crystals were grown in the hot, pressurized solutions and not as a result of a final return to ambient conditions.

**$[H_2NMe_2][UO_2(tce nm)] \cdot 3H_2O$  (1).**  $H_3tce nm$  (28 mg, 0.10 mmol) and  $[UO_2(NO_3)_2(H_2O)_2] \cdot 4H_2O$  (50 mg, 0.10 mmol) were dissolved in a mixture of water (0.5 mL) and *N,N*-dimethylformamide (0.2 mL). Yellow crystals of complex 1 were obtained within three days (15 mg, 23%). Anal. Calcd for  $C_{12}H_{26}N_2O_{13}U$ : C, 22.37; H, 4.07; N, 4.35. Found: C, 22.65; H, 3.75; N, 4.20%.

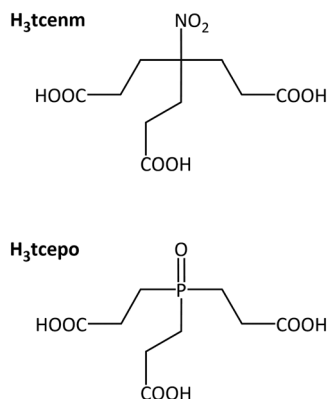
**$[C(NH_2)_3][UO_2(tce nm)] \cdot 0.5H_2O$  (2).**  $H_3tce nm$  (28 mg, 0.10 mmol),  $[UO_2(NO_3)_2(H_2O)_2] \cdot 4H_2O$  (35 mg, 0.07 mmol), and guanidinium nitrate (24 mg, 0.20 mmol) were dissolved in a mixture of water (0.6 mL) and acetonitrile (0.2 mL). Yellow crystals of complex 2 were obtained within four days (13 mg, 30%). Anal. Calcd for  $C_{22}H_{38}N_8O_{21}U_2$ : C, 21.54; H, 3.12; N, 9.13. Found: C, 21.43; H, 3.26; N, 9.16%.

**$[PPh_3Me][UO_2(tce nm)]$  (3).**  $H_3tce nm$  (28 mg, 0.10 mmol),  $[UO_2(NO_3)_2(H_2O)_2] \cdot 4H_2O$  (35 mg, 0.07 mmol), and  $PPh_3MeBr$  (36 mg, 0.10 mmol) were dissolved in a mixture of water (0.6 mL) and acetonitrile (0.2 mL). Yellow crystals of complex 3 were obtained within two weeks (45 mg, 78%). Anal. Calcd for  $C_{29}H_{30}N_2O_{10}PU$ : C, 42.40; H, 3.68; N, 1.70. Found: C, 42.23; H, 3.66; N, 1.74%.

**$[UO_2(tce nm)Ni(bipy)_2][UO_2(tce nm)] \cdot 3H_2O$  (4).**  $H_3tce nm$  (28 mg, 0.10 mmol),  $[UO_2(NO_3)_2(H_2O)_2] \cdot 4H_2O$  (35 mg, 0.07 mmol),  $Ni(NO_3)_2 \cdot 6H_2O$  (15 mg, 0.05 mmol), and 2,2'-bipyridine (24 mg, 0.15 mmol) were dissolved in a mixture of water (0.5 mL) and acetonitrile (0.2 mL). Pale pink crystals of complex 4 were obtained within three days (31 mg, 59%). Anal. Calcd for  $C_{40}H_{46}N_6NiO_{23}U_2$ : C, 31.74; H, 3.06; N, 5.55. Found: C, 31.92; H, 2.93; N, 5.75%.

**$[UO_2(tce nm)Cu(bipy)_2][UO_2(tce nm)] \cdot 3H_2O$  (5).**  $H_3tce nm$  (28 mg, 0.10 mmol),  $[UO_2(NO_3)_2(H_2O)_2] \cdot 4H_2O$  (35 mg, 0.07 mmol),  $Cu(NO_3)_2 \cdot 2.5H_2O$  (12 mg, 0.05 mmol), and 2,2'-bipyridine (24 mg, 0.15 mmol) were dissolved in a mixture of water (0.5 mL) and acetonitrile (0.2 mL). Blue-purple crystals of complex 5 were obtained within three days (21 mg, 40%). Anal. Calcd for  $C_{40}H_{46}CuN_6O_{23}U_2$ : C, 31.64; H, 3.05; N, 5.53. Found: C, 31.69; H, 2.88; N, 5.61%.

**$[UO_2(tce nm)_2Cu(R,S-Me_6cyclam)] \cdot 2H_2O$  (6).**  $H_3tce nm$  (28 mg, 0.10 mmol),  $[UO_2(NO_3)_2(H_2O)_2] \cdot 4H_2O$  (35 mg, 0.07 mmol), and  $[Cu(R,S-Me_6cyclam)](NO_3)_2$  (24 mg, 0.05 mmol) were dissolved in a mixture of water (0.6 mL) and acetonitrile (0.2 mL). Purple crystals of complex 6 were obtained within three days (16 mg, 31%). The elemental analysis results indicate the presence of approximately one additional acetonitrile molecule, in keeping with the presence of voids containing disordered solvent molecules in the structure (see below), although the number of acetonitrile molecules determined from elemental analysis is smaller than that estimated from the structure determination, probably due to the loss of part of the solvent upon drying. Anal. Calcd for



**Scheme 1** The ligands  $H_3tce nm$  and  $H_3tce po$ .



$\text{C}_{36}\text{H}_{64}\text{CuN}_6\text{O}_{22}\text{U}_2 + \text{CH}_3\text{CN}$ : C, 30.15; H, 4.46; N, 6.48. Found: C, 30.52; H, 4.18; N, 6.78%.

$[\text{UO}_2(\text{Htcepo})]$  (7).  $\text{H}_3\text{tcep}\cdot\text{HCl}$  (15 mg, 0.05 mmol) and  $[\text{UO}_2(\text{NO}_3)_2(\text{H}_2\text{O})_2]\cdot 4\text{H}_2\text{O}$  (25 mg, 0.05 mmol) were dissolved in water (0.5 mL). Yellow crystals of complex 7 were obtained within three days (17 mg, 64%). Anal. Calcd for  $\text{C}_9\text{H}_{13}\text{O}_9\text{PU}$ : C, 20.24; H, 2.45. Found: C, 19.80; H, 2.42%.

## Crystallography

Data collections were performed at 100(2) K on a Bruker D8 Quest diffractometer using an Incoatec Microfocus Source ( $\lambda$  3.0 Mo) and a PHOTON III area detector, and operated with APEX3.<sup>12</sup> The data were processed with SAINT,<sup>13</sup> and empirical absorption corrections were made with SADABS.<sup>14</sup> The structures were solved by intrinsic phasing with SHELXT,<sup>15</sup> and refined by full-matrix least-squares on  $F^2$  with SHELXL,<sup>16</sup> using the ShelXle interface.<sup>17</sup> When possible, hydrogen atoms bound to oxygen and nitrogen atoms were retrieved from residual electron density maps and they were refined either freely or with geometric restraints. The hydrogen atoms of the guanidinium cations in 2 and all carbon-bound hydrogen atoms in all compounds were introduced at calculated positions and treated as riding atoms with an isotropic displacement parameter equal to 1.2 times that of the parent atom (1.5 for  $\text{CH}_3$ ). For compound 2, the SQUEEZE<sup>18</sup> software was used to subtract the contribution of other, disordered solvent molecules to the structure factors; about 28 electrons were found in the unit cell, which could correspond to about 1.5 additional water molecules per formula unit. Use of SQUEEZE for compound

6 gave 40 electrons, corresponding to about 2 acetonitrile molecules per formula unit. For compound 2, the ADDSYM software<sup>19</sup> indicates the possible space group  $P2_1/c$  with 86% fit, but no satisfying refinement could be made in this space group. Crystal data and structure refinement parameters are given in Table 1. Drawings were made with ORTEP-3 (ref. 20) and VESTA,<sup>21</sup> and topological analyses were performed with ToposPro.<sup>22</sup>

## Luminescence measurements

Emission spectra were recorded on solid samples using an Edinburgh Instruments FS5 spectrofluorimeter equipped with a 150 W CW ozone-free xenon arc lamp, dual-grating excitation and emission monochromators (2.1 nm  $\text{mm}^{-1}$  dispersion; 1200 grooves per mm) and an R928P photomultiplier detector. The powdered compounds were pressed to the wall of a quartz tube, and the measurements were performed using the right-angle mode in the SC-05 cassette. An excitation wavelength of 420 nm was used in all cases and the emission was monitored between 450 and 600 nm. The quantum yield measurements were performed by using a Hamamatsu Quantaurus C11347 absolute photoluminescence quantum yield spectrometer and exciting the samples between 300 and 400 nm.

## Results and discussion

### Synthesis

It is quite usual to find that in solvothermal syntheses of uranyl ion complexes of carboxylates there is no need to add

Table 1 Crystal data and structure refinement details

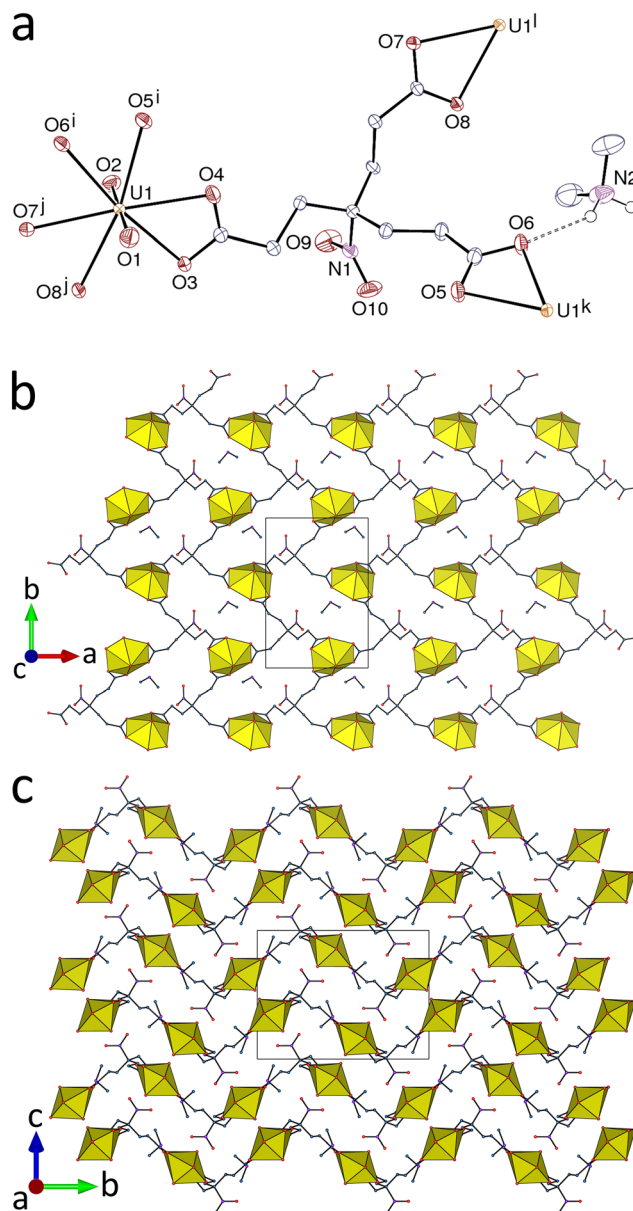
	1	2	3	4	5	6	7
Chemical formula	$\text{C}_{12}\text{H}_{26}\text{N}_2\text{O}_{13}\text{U}$	$\text{C}_{22}\text{H}_{38}\text{N}_8\text{O}_{21}\text{U}_2$	$\text{C}_{29}\text{H}_{30}\text{NO}_{10}\text{PU}$	$\text{C}_{40}\text{H}_{46}\text{N}_6\text{NiO}_{23}\text{U}_2$	$\text{C}_{40}\text{H}_{46}\text{CuN}_6\text{O}_{23}\text{U}_2$	$\text{C}_{36}\text{H}_{64}\text{CuN}_6\text{O}_{22}\text{U}_2$	$\text{C}_9\text{H}_{13}\text{O}_9\text{PU}$
$M/\text{g mol}^{-1}$	644.38	1226.66	821.54	1513.60	1518.43	1472.53	534.19
Crystal system	Monoclinic	Monoclinic	Monoclinic	Monoclinic	Monoclinic	Triclinic	Monoclinic
Space group	$P2_1/n$	$P2_1$	$P2_1/c$	$Cc$	$Cc$	$P\bar{1}$	$P2_1/c$
$a/\text{\AA}$	11.6052(7)	11.6822(6)	11.4827(4)	12.4106(7)	12.4847(7)	11.3872(4)	7.3820(5)
$b/\text{\AA}$	15.1797(10)	15.3912(8)	14.2168(5)	17.2410(9)	17.0888(11)	11.5224(4)	23.3063(18)
$c/\text{\AA}$	12.7681(13)	11.7959(5)	17.7990(6)	23.5189(14)	23.5455(16)	12.7894(5)	8.5496(7)
$\alpha/^\circ$	90	90	90	90	90	114.7971(13)	90
$\beta/^\circ$	116.773(3)	115.5602(16)	92.2934(15)	102.782(2)	103.131(2)	98.3735(14)	113.584(2)
$\gamma/^\circ$	90	90	90	90	90	110.7704(13)	90
$V/\text{\AA}^3$	2008.1(3)	1913.37(16)	2903.31(17)	4907.7(5)	4892.1(5)	1335.95(9)	1348.07(18)
$Z$	4	2	4	4	4	1	4
Reflections collected	71 359	46 015	89 833	60 642	39 817	53 010	44 893
Independent reflections	3814	7198	7497	9277	8866	6907	3476
Observed reflections [ $I > 2\sigma(I)$ ]	3589	7097	6879	9090	8727	6523	3338
$R_{\text{int}}$	0.052	0.056	0.060	0.068	0.058	0.052	0.060
Parameters refined	269	485	380	650	650	313	184
$R_1$	0.019	0.029	0.021	0.036	0.029	0.019	0.022
$wR_2$	0.046	0.075	0.055	0.093	0.071	0.042	0.052
$S$	1.049	1.040	1.054	1.069	1.044	1.074	1.096
$\Delta\rho_{\text{min}}/\text{e \AA}^{-3}$	−0.65	−0.72	−1.58	−1.15	−0.50	−0.89	−1.56
$\Delta\rho_{\text{max}}/\text{e \AA}^{-3}$	1.18	1.59	1.43	1.19	0.74	0.83	1.37
Flack parameter		0.416(11)		−0.001(10)	−0.003(7)		



a base to deprotonate the parent carboxylic acids, so that the trianionic form of the ligand found to be present in complexes 1–6 is no surprise. Incomplete deprotonation, however, is not unknown and given that this is seen in the presently unique example of the dicarboxylato species seen in complex 7, it cannot be attributed great significance, except in that this is the only complex of the present series obtained from pure water alone. The organic solvents, *N,N*-dimethylformamide and acetonitrile, used as co-solvents in the preparation of 1–6 both undergo hydrolysis under solvothermal conditions, a reaction which must lead to buffering of the medium at near-neutral pH, so that this is a factor which should be recognized as yet another influence on the composition of the crystalline deposits. A further influence of solubility is possibly reflected in the identical composition (and structure) of complexes 4 and 5, where differences were expected due to the unlike preferences of the two metal ions  $\text{Ni}^{\text{II}}$  and  $\text{Cu}^{\text{II}}$  for tris- and bis-(2,2'-bipyridine) binding and where sufficient 2,2'-bipyridine was added to form  $[\text{Ni}(\text{bipy})_3]^{2+}$ ,<sup>23</sup> but only the  $[\text{Ni}(\text{bipy})_2]^{2+}$  unit was present in the deposited crystals. Given the sensitivity of alkylphosphines in general to oxidation and the common observations of oxalate formation by oxidation of ligands during solvothermal syntheses of uranyl ion complexes, the presence of the phosphine oxide in complex 7 was perhaps to be expected, though it is of interest as a possible example of the simplest step in uranyl ion catalyzed oxidation of an organic substrate. This complex was repeatedly obtained from several different attempts at synthesis involving different additional reagents,  $[\text{Cu}(\text{R},\text{S-Me}_6\text{cyclam})](\text{NO}_3)_2$  or the couples  $\text{Ni}^{\text{II}}/\text{bipy}$  and  $\text{Zn}^{\text{II}}/\text{phen}$ , all with acetonitrile as cosolvent. While the same attempts were made with both  $\text{H}_3\text{-tcenm}$  and  $\text{H}_3\text{-tcep}$ , only complex 7 resulted in the latter case, most syntheses giving amorphous precipitates or no solid whatever, which points to the particular stability and/or insolubility of this complex.

### Crystal structures

The complex  $[\text{H}_2\text{NMe}_2][\text{UO}_2(\text{tcenm})]\cdot 3\text{H}_2\text{O}$  (**1**) includes dimethylammonium counterions formed from hydrolysis of the organic cosolvent *N,N*-dimethylformamide, as very commonly observed,<sup>24</sup> and the 1:1 metal/ligand stoichiometry used during the synthesis is retained in the solid state. The uranyl ion is tris- $\kappa^2\text{O},\text{O}'$ -chelated by three carboxylate groups from three  $\text{tcenm}^{3-}$  ligands [ $\text{U}-\text{O}(\text{oxo})$ , 1.763(3) and 1.769(3) Å;  $\text{U}-\text{O}(\text{carboxylate})$ , 2.440(2)–2.496(2) Å] (Fig. 1). The ligand  $\text{tcenm}^{3-}$  has a conformation completely lacking in symmetry and is therefore a source of chirality but both enantiomers are present since the complex crystallizes in a centrosymmetric space group. Both metal and ligand are 3-coordinated (3-c) nodes in the diperiodic network formed, which is parallel to (001) and has the  $\{6^3\}$  point symbol and the **hcb** topological type, as commonly found in tris-chelated uranyl carboxylate complexes. When viewed down [100], the homochiral layers have a zigzag profile and the nitro groups



**Fig. 1** (a) View of complex **1** with displacement ellipsoids shown at the 50% probability level. Solvent molecules and carbon-bound hydrogen atoms are omitted, and the hydrogen bond is shown as a dashed line. Symmetry codes:  $i = 3/2 - x, y + 1/2, 3/2 - z$ ;  $j = x - 1, y, z$ ;  $k = 3/2 - x, y - 1/2, 3/2 - z$ ;  $l = x + 1, y, z$ . (b) View of the diperiodic assembly showing uranium coordination polyhedra. (c) Packing with layers viewed edge-on.

are located near the peaks and troughs and thus alternate in their orientation relative to the mean plane. The nitro C–N bonds lie at an angle near 60° to the mean plane and the nitrogen atom lies at 7.127(3)–7.532(3) Å of the three  $\text{U}^{\text{VI}}$  centres bound to the same ligand, although the shortest  $\text{N}\cdots\text{U}$  separations of 4.509(3) and 5.813(3) Å involve two centres in one adjacent sheet. The  $\text{H}_2\text{NMe}_2^+$  cation forms bifurcated hydrogen bonds with two carboxylate, one nitro and one water oxygen atoms [ $\text{N}\cdots\text{O}$  distances, 2.779(6)–2.911(5) Å;  $\text{N}-\text{H}\cdots\text{O}$  angles, 113(4)–154(5)°], thus linking two





sheets, while the water molecule for which the hydrogen atoms could be located also links two sheets through hydrogen bonding to one oxo and one carboxylate oxygen donors. Due to the inability to locate the hydrogen atoms of the other water molecules, the hydrogen bonding network cannot be defined in complete detail but the identifiable bonds unite the sheets into a triperiodic assembly. The packing is quite compact, as indicated by the Kitaigorodsky packing index (KPI, evaluated with PLATON<sup>19</sup>) of 0.70.

The complex involving the guanidinium counterion,  $[C(NH_2)_3][UO_2(tcenm)] \cdot 0.5H_2O$  (**2**), crystallizes in the Sohncke group  $P2_1$  with two independent uranyl cations in the asymmetric unit, both in the same environment as in complex **1** [U–O(oxo), 1.763(6)–1.793(7) Å; U–O(carboxylate), 2.416(12)–2.501(6) Å] (Fig. 2). The anionic sheets formed, parallel to (10 $\bar{1}$ ) and here also with the **hcb** topology, have more a square-wave profile than those in **1** when viewed down [101], with the uranyl equatorial planes roughly parallel to the sheet mean plane instead of being tilted as in **1**. The ligand nitro groups again project in alternation to each side of the sheets, with the C–N bonds at an angle of close to 75° to the mean plane. The sheets are tightly packed so as to form channels parallel to [001], with a section of  $\sim 5$  Å  $\times$   $\sim 10$  Å, occupied by the guanidinium counterions and water molecules. There are two slightly different minimum U $\cdots$ NO<sub>2</sub> separations of 4.812(8) and 4.720(8) Å, both slightly longer than the shorter one in **1** but again between and not within sheets. In both sheets, the ligand has a pseudo-mirror plane containing one of the carboxylate arms and the nitro group, and it is thus achiral ( $C_s$  symmetry). The chirality of the crystal appears to result from the cation–water associates within the channels of the structure. The guanidinium cations form either eight or six simple or bifurcated hydrogen bonds involving carboxylate, nitro or water acceptors [N $\cdots$ O distances, 2.910(14)–3.171(15) Å; N–H $\cdots$ O angles, 125–159°]. Pairs of guanidinium cations bridged by hydrogen bonding to a water molecule form units which, considered alone, have  $C_2$  symmetry and are therefore chiral, so that the chirality of the crystal can be considered a result of the fact that within a given crystal all these units have the same configuration. Further hydrogen bonding interactions of the water molecule and both cations, mostly involving carboxylate acceptors, reduce the actual symmetry to  $C_1$  but only one cation of a given unit interacts with a nitro group, producing the inequivalence of adjacent sheets. The KPI amounts to only 0.66, some voids being occupied by unresolved solvent molecules (see Experimental).

Use of the bulky methyltriphenylphosphonium cation yields the complex  $[PPh_3Me][UO_2(tcenm)]$  (**3**), in which the uranium atom environment is the same as in **1** and **2** [U–O(oxo), 1.7721(17) and 1.7773(18) Å; U–O(carboxylate), 2.4345(18)–2.4930(17) Å] (Fig. 3). Here also, the network formed, parallel to (001), has the **hcb** topology, and its shape is close to that found in **1**. The packing of sheets is however different, as indicated by the shortest intersheet U $\cdots$ N contact at 5.823(3) Å, much larger than in **1** as a result of

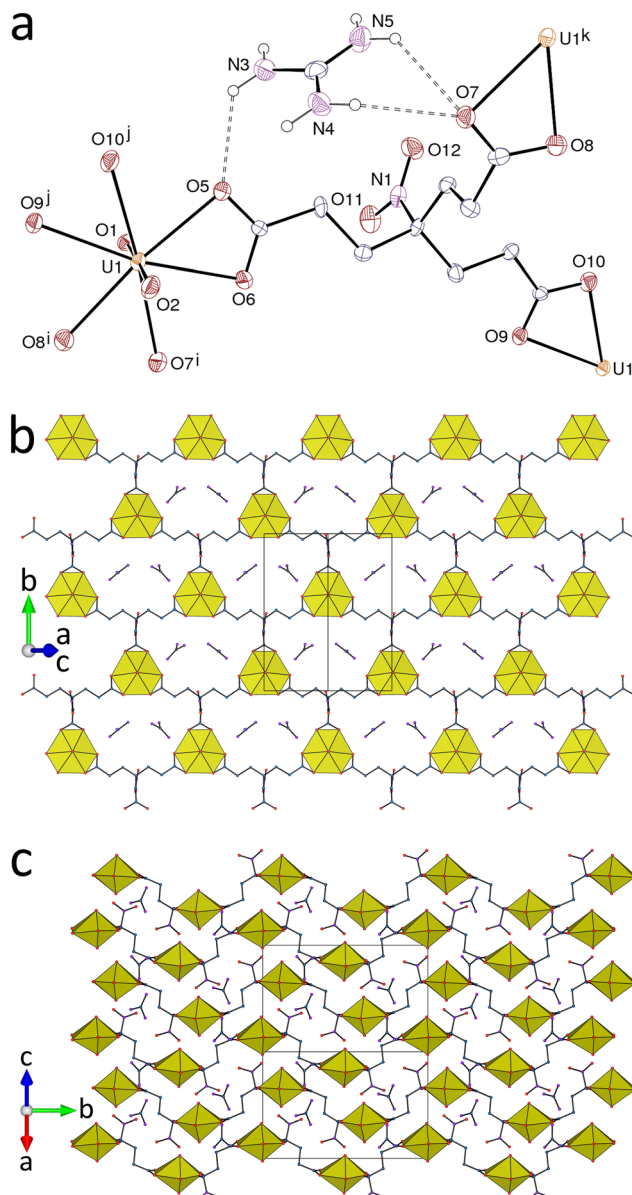


Fig. 2 (a) View of one of the two independent complex units in compound **2** with displacement ellipsoids shown at the 30% probability level. Solvent molecules and carbon-bound hydrogen atoms are omitted, and hydrogen bonds are shown as dashed lines. Symmetry codes:  $i = 1 - x, y + 1/2, 2 - z$ ;  $j = x + 1, y, z + 1$ ;  $k = 1 - x, y - 1/2, 2 - z$ ;  $l = x - 1, y, z - 1$ . (b) View of the diperiodic assembly showing uranyl coordination polyhedra. (c) Packing with layers viewed edge-on.

both the larger spacing of the sheets and a lateral displacement resulting in the uranium atoms being arranged in layers parallel to (010). The  $PPh_3Me^+$  counterions occupy the interlayer spaces, with one aromatic ring crossing a hexagonal cell, and they are too far away from one another for phenyl-embrace or  $\pi$ -stacking interactions, all centroid $\cdots$ centroid distances being larger than 5 Å. The KPI of 0.70 indicates that no significant void is present.

The two complexes  $[UO_2(tcenm)M(bipy)_2][UO_2(tcenm)] \cdot 3H_2O$  with  $M = Ni$  (**4**) or  $Cu$  (**5**) are isomorphous, despite anticipation



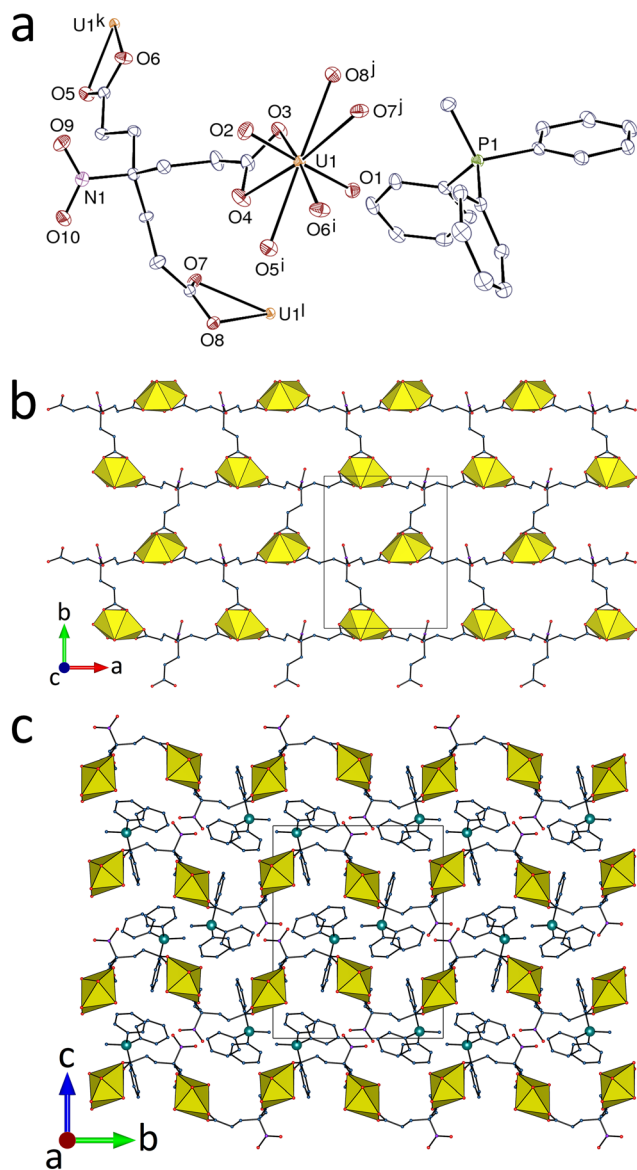


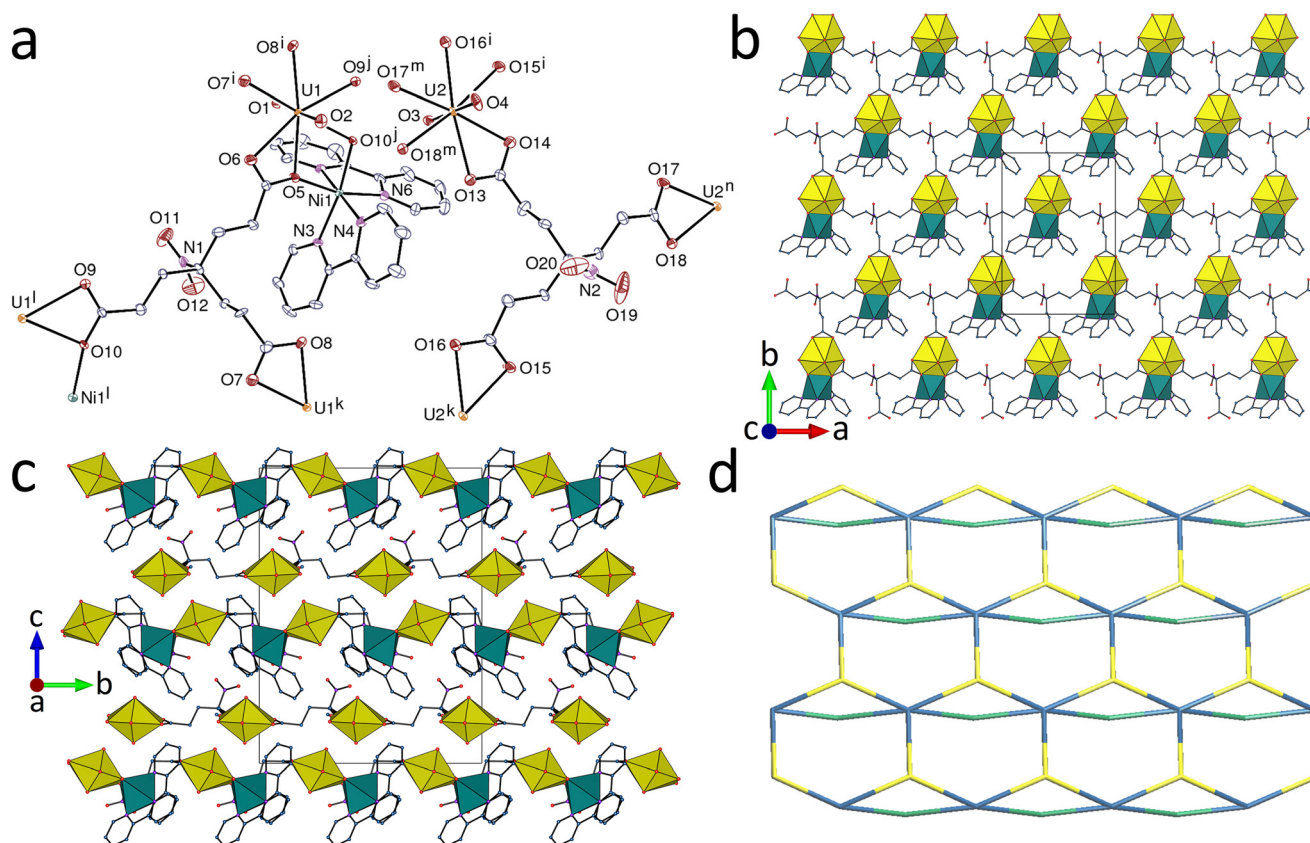
Fig. 3 (a) View of compound **3** with displacement ellipsoids shown at the 50% probability level and hydrogen atoms omitted. Symmetry codes:  $i = x + 1, y, z$ ;  $j = 1 - x, y + 1/2, 3/2 - z$ ;  $k = x - 1, y, z$ ;  $l = 1 - x, y - 1/2, 3/2 - z$ . (b) The diperic assembly with uranium coordination polyhedra yellow. (c) Packing with layers viewed edge-on.

that the different stereochemical preferences of  $\text{Ni}^{\text{II}}$  and  $\text{Cu}^{\text{II}}$  might have been of influence, and they crystallize in the non-centrosymmetric space group  $Cc$ . As shown in Fig. 4 in the case of complex **4**, the crystal contains two independent, cationic and anionic units which are hetero- and homometallic, respectively. In both units, the uranium environment is that found in the previous complexes [U–O(oxo), 1.762(10)–1.787(10) Å; U–O(carboxylate), 2.424(7)–2.515(10) Å, including both compounds], and, as in **2**, the  $\text{tcenm}^{3-}$  ligand has either pseudo-mirror symmetry in the cation, or is not far from it in the anion (with the nitro group slightly tilted). The transition metal cation is bound to two carboxylate oxygen atoms, thus forming a four-membered  $\text{UMO}_2$  ring, and to two chelating bipy

ligands, its environment being approximately octahedral, with slightly more distortion with Cu than with Ni [Ni–N, 2.030(10)–2.061(10) Å; Ni–O, 2.130(8) and 2.156(9) Å; Cu–N, 1.953(7)–2.076(8) Å; Cu–O, 2.362(7) and 2.450(6) Å]. The transition metal achieves 6-coordination by binding to  $\text{UO}_2(\text{carboxylate})_3^-$  units in a manner similar to that long known in the case of  $\text{Na}[\text{UO}_2(\text{CH}_3\text{CO}_2)_3]$ ,<sup>25</sup> where the uranate anion behaves as a metalloligand *via* chelation of  $\text{Na}^+$  by O–U–O units derived from two adjacent acetates. Both cation and anion form uranyl-based **hcb** networks parallel to (001), with the  $\text{M}(\text{bipy})_2^{2+}$  groups forming an additional link between two  $\text{tcenm}^{3-}$  nodes in the cation (Fig. 4d). Viewed down [001], the triuranacyclic units have a near-rectangular form very similar to that in complex **2**, but when viewed down [100] both sheets have a flattened sawtooth profile in which all nitro groups lie to one side of their sheet. The nitro groups of the cationic sheet confront those of the anionic, so that an alternative view of the structure is as of one where neutral double sheets stack down [001].  $\text{U} \cdots \text{N}(\text{nitro})$  separations are considerably longer than in complexes **1** and **2**, with the shortest being  $\text{U2} \cdots \text{N1}$  of 6.037(13) Å in **4** and 6.104(9) Å in **5**, again an intersheet contact, all the others being larger than 6.7 Å. This may in part be a consequence of the fact that while the nitro groups retain their capacity to become involved in hydrogen bonding with water molecules, there is also an insertion of one nitro group of the anionic sheet into the space between bipy ligands in an adjacent cationic sheet producing  $\text{O19} \cdots \pi$  interactions [O  $\cdots$  centroid distances, 3.062(19) and 3.353(19) Å in **4**, 2.904(12) Å in **5**; N–O  $\cdots$  centroid angles, 157 and 126° in **4**, 155° in **5**]; these interactions appear as exceeding dispersion on the Hirshfeld surface (HS) calculated with CrystalExplorer.<sup>26</sup> The 6-coordinate  $\text{M}^{\text{II}}$  unit is chiral and each cationic sheet is enantiomerically pure but the chirality alternates from one sheet to the next. The  $\text{M}(\text{bipy})_2^{2+}$  substituents largely occupy the space defined by the triuranacycles, so that although the projection of sheets onto one another is very similar to that in complex **2**, no channels are apparent because they are blocked by the  $\text{M}(\text{bipy})_2^{2+}$  attachments. This does mean, however, that, because the anionic sheets have cyclic units occupied only by presumably replaceable water molecules, the complete structure can be regarded as containing cavities with caps defined by enantiomeric  $\text{M}(\text{bipy})_2(\text{O})_2$  entities at a  $\text{M} \cdots \text{M}$  separation of 11.8301(8) and 11.8006(8) Å in **4** and **5**, respectively. With KPIs of 0.68 and 0.69, the packings in **4** and **5** do not contain significant free spaces.

While complexes **1–5** have structures based on a common **hcb** network topology, introduction of the  $\text{Cu}(\text{R},\text{S}\text{-Me}_6\text{-cyclam})^{2+}$  species results in formation of a heterometallic complex with a different connectivity,  $[(\text{UO}_2)_2(\text{tcenm})_2\text{Cu}(\text{R},\text{S}\text{-Me}_6\text{-cyclam})] \cdot 2\text{H}_2\text{O}$  (**6**). The unique uranium atom is here in a pentagonal-bipyramidal environment, being chelated by one carboxylate group and bound to three more oxygen donors from three different ligands [U–O(oxo), 1.7803(18) and 1.7899(17) Å; U–O(carboxylate), 2.4560(17) and 2.4850(16) Å for the chelating group and 2.3149(17)–2.3269(17) Å for the others] (Fig. 5). The  $\text{tcenm}^{3-}$  ligand has thus one  $\kappa^2\text{O},\text{O}'$ -





**Fig. 4** (a) View of compound **4** with displacement ellipsoids shown at the 50% probability level. Solvent molecules and hydrogen atoms are omitted. Symmetry codes:  $i = x + 1/2, y + 1/2, z; j = x + 1, y, z; k = x - 1/2, y - 1/2, z; l = x - 1, y, z; m = x - 1/2, y + 1/2, z; n = x + 1/2, y - 1/2, z$ . (b) The heterometallic dimeric assembly with uranium coordination polyhedra yellow and those of nickel green. (c) Packing with layers viewed edge-on. (d) Nodal representation of the heterometallic network (yellow, uranium nodes; blue,  $\text{tcnm}^{3-}$  nodes; green, nickel edges; same orientation as in b).

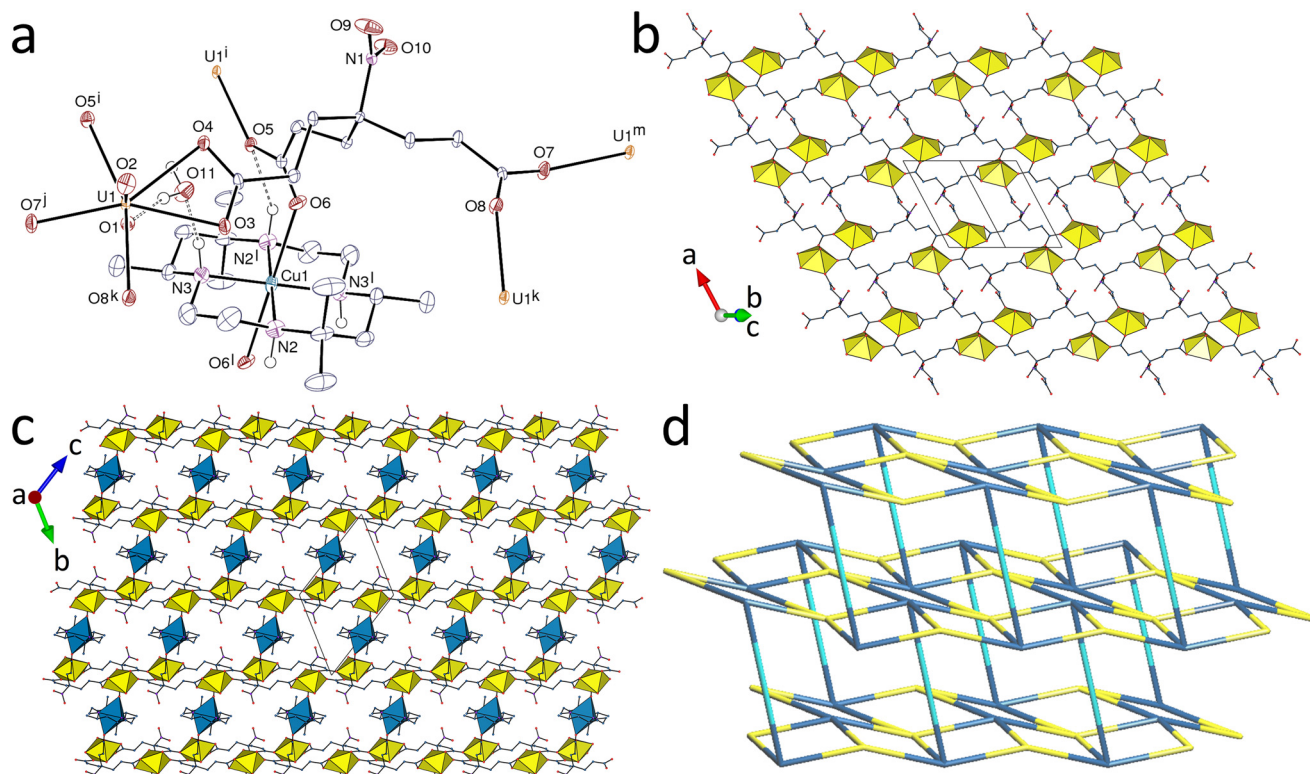
chelating, one  $\mu_2\text{-}\kappa^1\text{O}:\kappa^1\text{O}'$ -bridging and one monodentate carboxylate groups in its coordination to uranyl. If only uranium and  $\text{tcnm}^{3-}$  are considered, both are 4-c nodes and the coordination polymer formed is dimeric and parallel to  $(01\bar{1})$ , having the  $\{4^4\cdot 6^2\}$  point symbol and the common **sql** topological type. With respect to the ideal square lattice, two kinds of cells, either 8- or 20-membered, are however present here. These subunits are further linked through  $\text{Cu}^{\text{II}}$  bridges, the transition metal cation being bound to the four nitrogen atoms of the centrosymmetric azamacrocycle and to two axial carboxylate oxygen atoms to give an axially elongated octahedral environment [ $\text{Cu}-\text{N}$ , 2.020(2) and 2.061(2) Å;  $\text{Cu}-\text{O}$ , 2.5581(18) Å]. The  $\text{tcnm}^{3-}$  ligand thus becomes a 5-c node, with one chelating and two  $\mu_2\text{-}\kappa^1\text{O}:\kappa^1\text{O}'$ -bridging carboxylate groups. The resulting assembly is a 4,5-c binodal triperiodic framework with the  $\{4^4\cdot 6^2\}\{4^4\cdot 6^6\}$  point symbol (first symbol for U and second for  $\text{tcnm}^{3-}$ ) and the **tcs** ( $\text{ThCr}_2\text{Si}_2$ ) topological type, and it is to the best of our knowledge the first occurrence of this somewhat uncommon topology<sup>27</sup> in uranyl chemistry. As usually found when transition metal azamacrocyclic complexes are used as structure-directing species in uranyl cation coordination polymers, one ammonium group is hydrogen bonded to the

carboxylate group bound to the transition metal cation, while the other is linked to the water molecule, itself hydrogen bonded to oxo and carboxylate groups [ $\text{N}\cdots\text{O}$  distances, 2.979(3)–3.373(3) Å;  $\text{N}-\text{H}\cdots\text{O}$  angles, 115–161°;  $\text{O}\cdots\text{O}$  distances, 2.914(3) and 2.919(3) Å;  $\text{O}-\text{H}\cdots\text{O}$  angles, 161(3) and 165(3)°]. The shortest  $\text{U}\cdots\text{N}(\text{nitro})$  contact is here between atoms pertaining to the same dimeric subunit, at 5.987(2) Å, due to the large intersheet separation associated with the  $\text{Cu}^{\text{II}}$  links. The KPI is 0.63 only, some voids being occupied by disordered acetonitrile molecules (see Experimental).

Replacing  $\text{H}_3\text{tcnm}$  by  $\text{H}_3\text{tcep}$  has remarkable effects on the complexes produced.  $\text{H}_3\text{tcep}$  is oxidized *in situ* to give the corresponding phosphine oxide, which retains one of its carboxylic protons in the complex  $[\text{UO}_2(\text{Htcepo})]$  (**7**). The asymmetric unit contains a single uranium atom in a pentagonal-bipyramidal environment resulting from chelation by one carboxylate group and bonding to two more carboxylate donors and one oxide group from three different ligands [ $\text{U}-\text{O}(\text{oxo})$ , 1.774(3) and 1.778(3) Å;  $\text{U}-\text{O}(\text{carboxylate})$ , 2.439(3) and 2.544(3) Å for the chelating group, 2.337(3) and 2.349(3) Å for the others;  $\text{U}-\text{O}(\text{phosphine oxide})$ , 2.302(3) Å] (Fig. 6). 98 crystal structures of complexes with uranyl cations







**Fig. 5** (a) View of compound **6** with displacement ellipsoids shown at the 50% probability level. Carbon-bound hydrogen atoms are omitted and hydrogen bonds are shown as dashed lines. Symmetry codes:  $i = 1 - x, 1 - y, 1 - z$ ;  $j = x, y + 1, z + 1$ ;  $k = 2 - x, 1 - y, 1 - z$ ;  $l = 1 - x, -y, 1 - z$ ;  $m = x, y - 1, z - 1$ . (b) The uranyl-based dimeric subunit. (c) View of the triperiodic framework with uranium coordination polyhedra yellow and those of copper blue. (d) Nodal representation of the framework (yellow, uranium; dark blue,  $\text{tcenm}^{3-}$  nodes; light blue, copper edges).

bound to phosphine oxides are reported in the Cambridge Structural Database (CSD, Version 5.43),<sup>28</sup> with U–O(P) bond lengths in the range of 2.257–2.462 Å [mean value, 2.36(4) Å]. The uranium pentagonal-bipyramidal coordination found here contrasts with that in the complexes of  $\text{tcenm}^{3-}$ , where hexagonal-bipyramidal coordination predominates, though with one example of pentagonal-bipyramidal coordination in **6**, so no real conclusion can be drawn from what is seen in **7**, even if it is tempting to believe that the strong donor capacity of PO may be a factor favouring a lower coordination number. Although  $\text{H}_3\text{tcep}$  and its anions have previously been used as ligands, with 14 crystal structures of complexes reported in the CSD,  $\text{H}_3\text{tcepo}$  or its deprotonated forms are unknown as ligands, and only the crystal structure of the uncomplexed molecule has been described.<sup>29</sup> The  $\text{Htcepo}^{2-}$  ligand has one  $\kappa^2\text{O}, \text{O}'$ -chelating and one  $\mu_2\text{-}\kappa^1\text{O}:\kappa^1\text{O}'$ -bridging carboxylate groups, and one uncoordinated carboxylic group, and it is thus a 4-c node, as is also the uranium atom. The uninodal dimeric network formed, parallel to (010), has the point symbol  $\{6^6\}$ . This point symbol is the same as that of the triperiodic **dia** topology, but the vertex symbols<sup>30</sup> are different,  $6_2\cdot 6_2\cdot 6_3\cdot 6_6\cdot 6_4\cdot 6_4$  here and  $6_2\cdot 6_2\cdot 6_2\cdot 6_2\cdot 6_2\cdot 6_2$  for the **dia** net (and also the less common **lon** net). Instead of containing adamantane-like units as found in the **dia** network, there are here crossings giving Hopf links with a

multiplicity of 2 (Fig. 6d). The sheets formed display two layers of uranyl ions and the carboxylic groups are directed outward on each side, giving a width of  $\sim 14$  Å between the planes defined by the outermost oxygen atoms, the sheets being slightly interdigitated. The carboxylic group makes an intersheet hydrogen bond with the uranium-bound carboxylate oxygen atom O4 [O $\cdots$ O distance, 2.783(4) Å; O–H $\cdots$ O angle, 150(6)°], which induces a further level of organization leading to formation of an intricate and compact (KPI, 0.72) 4,5-c triperiodic network with the point symbol  $\{6^6\}\{6^9\cdot 8\}$  (with hydrogen bonds considered as edges between  $\text{Htcepo}^{2-}$  nodes) and the presence of eight crossings (Hopf links).

### Discussion of the crystal structures

While the role of metal ions in coordination polymers and frameworks can be purely structural, particular properties of the metal ions such as magnetism, redox activity, electronic absorption, luminescence, stereochemical preferences, and catalytic activity can also be critical in regard to any applications, as is evident in numerous recent research articles and reviews.<sup>31</sup> A further tool of structural control can arise from internal functionality of the bridging ligands,<sup>9</sup> simple examples being where an additional donor atom provides a site for chelation<sup>32</sup> or, more commonly, where stacking arrays<sup>33</sup> are formed by small aromatic units,<sup>34</sup>





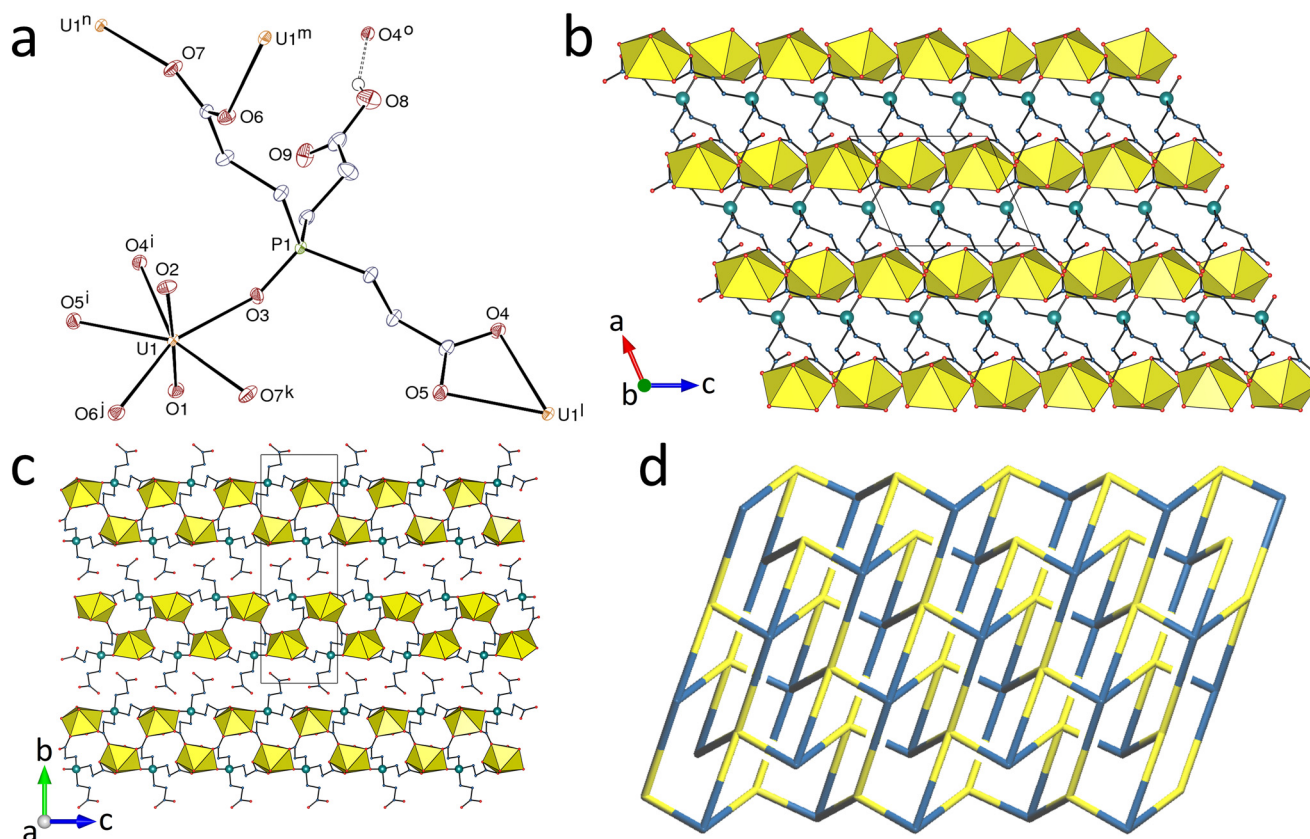


Fig. 6 (a) View of compound 7 with displacement ellipsoids shown at the 50% probability level. Carbon-bound hydrogen atoms are omitted and the hydrogen bond is shown as a dashed line. Symmetry codes:  $i = x, y, z - 1$ ;  $j = x - 1, 3/2 - y, z - 1/2$ ;  $k = x - 1, y, z$ ;  $l = x, y, z + 1$ ;  $m = x + 1, 3/2 - y, z + 1/2$ ;  $n = x + 1, y, z$ ;  $o = 2 - x, 1 - y, 2 - z$ . (b) The diperiodic assembly with uranium coordination polyhedra yellow. (c) Packing with layers viewed edge-on. (d) Nodal representation of the diperiodic network (yellow, uranium; blue, Htcepo<sup>2-</sup> nodes).

although once again such functionality in more sophisticated forms,<sup>35,36</sup> has been exploited to engender special properties in the structures formed. In the particular case of tripodal ligands, the nature of the bridgehead is of prime importance,<sup>37</sup> as shown in the present case. The ligand Htcepo<sup>2-</sup> produced by oxidation of the reactant H<sub>3</sub>tcep under the reaction conditions and found in the neutral crystalline complex 7 provides a significant contrast with the similar tripod ligand tcecm<sup>3-</sup> in its mode of coordination. While valence bond representations of both NO<sub>2</sub> and PO units mean that they can both be considered as “vicinal zwitterions”,<sup>38</sup> and thus as sources of negatively charged oxygen donors, the known coordination chemistry of phosphine oxides is vastly more extensive<sup>39</sup> than that of nitroalkanes, which are commonly considered as rather poor ligands,<sup>40</sup> although nitro coordination and chelation are well-known for functionalized nitroarenes.<sup>41</sup> The present work confirms an apparently major difference in donor capacity of the two units in that only in 7 is coordination other than of carboxylate donors observed. More significantly, the  $\kappa^1\text{O}$  U–OP bond length of 2.302(3) Å is shorter than either of those of the  $\kappa^1\text{O}$  U–O(carboxylate) bonds of 2.337(3) and 2.349(3) Å. This may explain why one carboxylic acid group of H<sub>3</sub>tcep retains its proton and does not bind to U<sup>VI</sup>, though such a

conclusion must be qualified by recognizing that what is observed in the solid state is determined to some extent by solubility and does not necessarily represent the dominant species in solution. Coordination of the bridgehead unit of Htcepo<sup>2-</sup> does not produce a ligand conformation which is radically different from that adopted by tcecm<sup>3-</sup> and both can be considered as flattened, extended tripods where each of the donor units binds to a separate U<sup>VI</sup> centre. It has been estimated that a minimum of 6-atom spacing (in addition to the bonding groups) is required for coordination to one uranyl centre of the three carboxylate groups of a tripodal ligand with a tertiary ammonium bridgehead hydrogen bonded to one uranyl oxo group.<sup>1a</sup> As with nitrilotriacetate,<sup>5</sup> the two present ligands fall short of this limit and it is thus unsurprising that the coordinating carboxylate groups in both tcecm<sup>3-</sup> and Htcepo<sup>2-</sup> are divergently oriented and bound to different cations. The non-coordinating nature of the nitro group in the first case leads in most cases (complexes 1–5) to threefold uranyl  $\kappa^2\text{O},\text{O}'$ -chelation (with additional bridging to transition metal cations in 4 and 5) and formation of **hcb** networks. Only in complex 6 is a different connectivity of tcecm<sup>3-</sup> observed, with carboxylate axial bonding to copper(II) disrupting the previous pattern and making the ligand a  $\kappa^2\text{O},\text{O}';\text{bis}(\mu_2-\kappa^1\text{O}:\kappa^1\text{O}')$ -bonded



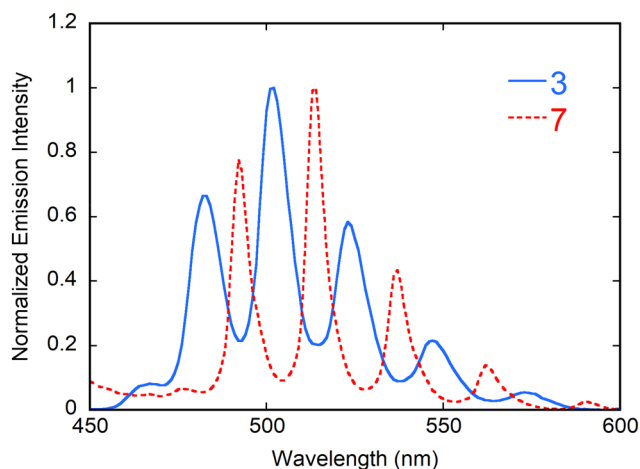


Fig. 7 Emission spectra of complexes **3** and **7** in the crystalline state upon excitation at 420 nm.

node; a diperiodic uranyl coordination polymer, here of **sql** topology, is however formed here also, with triperiodic extension ensured by the  $\text{Cu}^{\text{II}}$  pillars. Bridgehead coordination with the  $\text{Htcepo}^{2-}$  ligand, although it does not increase much the overall number of bonded cations due to the carboxylic group being uncoordinated, yields nevertheless a diperiodic network which is not quasi-planar or able to be embedded into a plane, but displays crossings of rings and can thus be considered as three-dimensional diperiodic.<sup>42</sup>

### Luminescence properties

Emission spectra of complexes **1–7** have been measured in the solid state under excitation at 420 nm. Complexes **2** and **4–6** are non-emissive, unsurprisingly for the 3d metal cation-containing complexes **4–6**, for which this is usual and probably results from either preferential absorption by the transition metal cation or energy transfer and non-radiative relaxation,<sup>43</sup> but less so for **2**, although weak emission in guanidinium-containing species has previously been observed.<sup>44</sup> Although emission from **1** was detectable with maxima very close to those of complex **3**, it was particularly weak and showed an anomalous sharp peak at 514 nm possibly due to an impurity. The spectra of complexes **3** and **7**, which both have a photoluminescence quantum yield (PLQY) of 9%, indicating no special efficacy of either the nitro or PO substituent as an antenna for  $\text{U}^{\text{VI}}$  excitation, are shown in Fig. 7 ( $\text{U} \cdots \text{N}(\text{nitro})$  separations in the  $\text{tcenm}^{3-}$  complexes were estimated (see above) to see if they might correlate with PLQY values and indicate a significant role as an antenna for the nitro group but this was clearly not the case). These spectra are well-resolved and show the typical vibronic progression due to the  $S_{11} \rightarrow S_{00}$  and  $S_{10} \rightarrow S_{0v}$  ( $v = 0–4$ ) transitions of the uranyl ion.<sup>45</sup> The main maxima positions for **3** (482, 502, 523 and 547 nm) and **7** (492, 514, 537 and 562 nm) are in agreement with the red-shift usually associated with the passage from six- to five-coordinated equatorial environment in carboxylate uranyl ion complexes.<sup>46</sup>

## Conclusions

Two triple-armed ligands,  $\text{H}_3\text{tcenm}$  and  $\text{H}_3\text{tcepo}$ , the latter oxidized *in situ* into the corresponding phosphine oxide  $\text{H}_3\text{tcepo}$ , have been used to synthesize seven uranyl ion complexes under (solvo)-hydrothermal conditions and in the presence of various structure-directing cations. The tricarboxylate  $\text{tcenm}^{3-}$  appears to be much more sensitive to the presence of these additional species than the dicarboxylate  $\text{Htcepo}^{2-}$  since six of the complexes involve the former, while only one complex could be obtained, under different conditions, with the latter. Five of the six  $\text{tcenm}^{3-}$  complexes share however the common feature of threefold  $\kappa^2\text{O},\text{O}'$ -chelation of three uranyl ions to give **hcb** networks, some geometric variety being introduced by the different counterions or additional coordinated transition metal cations, with the nitro substituent being involved in various hydrogen bonding interactions ( $\text{O} \cdots \text{HN}$  and  $\text{O} \cdots \text{HC}$ ) presumably of some influence. Pillaring of **sql** networks by  $\text{Cu}(\text{R},\text{S}\text{-Me}_6\text{cyclam})^{2+}$  provides the only triperiodic framework in the series, which has the **tcs** topology. In no case does the nitro group participate in complexation, in contrast to the phosphine oxide group which is bound to uranyl in the  $\text{Htcepo}^{2-}$  complex, yielding a diperiodic, three-dimensional network with rings forming Hopf links. Broad considerations of the donor capacity of phosphine-oxide and carboxylate towards  $\text{U}^{\text{VI}}$  indicate that the former is the more effective,<sup>47</sup> explaining the unsymmetrical bonding and incomplete ionization of  $\text{Htcepo}$ , while the poor coordinating ability of an aliphatic nitro group explains the action of  $\text{tcenm}^{3-}$  as a tricarboxylate donor. Both these triple-armed ligands are too small to encompass a single uranyl cation bound to all three carboxylate groups, and indeed adopt conformations describable as rather flattened tripods, so that they act instead as divergent ligands with either an uncoordinated or a coordinated bridgehead. The conformation restrictions due to the sophisticated design of triple-armed ligands forming mononuclear uranyl ion complexes<sup>2</sup> therefore appear to have been critical in enforcing a convergent, tripodal form.

## Conflicts of interest

There are no conflicts of interest to declare.

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