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# Introduction

*trans*-9,10-Dihydro-9,10-ethanoanthracene-11,12-dicarboxylic acid (deadeH<sub>2</sub>, Scheme 1) has been the subject of various studies of its capacity to form, sometimes selectively, crystalline clathrates with small organic molecules.<sup>1-4</sup> Less

# Uranyl ion coordination polymers with the dibenzobarrelene-based *rac*- and (*R*,*R*)-*trans*-9,10-dihydro-9,10-ethanoanthracene-11,12-dicarboxylate ligands<sup>†</sup>

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trans-9,10-Dihydro-9,10-ethanoanthracene-11,12-dicarboxylic acid (deadcH<sub>2</sub>), in its racemic or *R*,*R* enantiomeric forms, has been used to synthesize eight uranyl ion complexes under solvo-hydrothermal conditions. [UO<sub>2</sub>(deadc)]·1.5CH<sub>3</sub>CN (1) and [H<sub>2</sub>NMe<sub>2</sub>]<sub>2</sub>[(UO<sub>2</sub>)<sub>2</sub>(deadc)<sub>3</sub>]·2H<sub>2</sub>O (2) crystallize as monoperiodic coordination polymers in which deadc<sup>2-</sup> forms both 4- and 7-membered chelate rings. Although synthesized in the same conditions as 2, the enantiomerically pure complex [H<sub>2</sub>NMe<sub>2</sub>]<sub>4</sub>[(UO<sub>2</sub>)<sub>2</sub>(O)(*R*,*R*-deadc)<sub>2</sub>]<sub>2</sub> (3) is a discrete tetranuclear complex containing two  $\mu_3$ -oxo anions. Association with the zwitterion Ni(tpyc)<sub>2</sub>, where tpyc<sup>-</sup> is 2,2':6',2''-terpyridine-4'-carboxylate, gives [(UO<sub>2</sub>)<sub>2</sub>(deadc)(deadcH)(NO<sub>3</sub>) Ni(tpyc)<sub>2</sub>]·CH<sub>3</sub>CN·2H<sub>2</sub>O (4), a rake-shaped monoperiodic assembly. [UO<sub>2</sub>(deadc)(DMA)] (5), [UO<sub>2</sub>(deadc)] (6) and [PPh<sub>4</sub>]<sub>2</sub>[(UO<sub>2</sub>)<sub>2</sub>(*R*,*R*-deadc)<sub>3</sub>] (7) crystallize as diperiodic networks with the **fes, sql** and **hcb** topologies, respectively, the thick layers formed being coated on both sides by protruding, hydrophobic dibenzobarrelene groups. Finally, [(UO<sub>2</sub>)<sub>2</sub>Ag<sub>2</sub>(deadc)<sub>3</sub>(CH<sub>3</sub>CN)<sub>2</sub>]·0.5H<sub>2</sub>O (8) contains monoperiodic uranyl-deadc<sup>2-</sup> subunits which are assembled into a triperiodic framework by bridging silver(i) cations, the latter interacting with both carboxylate groups and aromatic rings. Except for **6**, all these complexes are emissive with photoluminescence quantum yields of 2–26%, and most spectra display the usual vibronic fine structure of uranyl emission.

well established are the roles of its conjugate anions in coordination chemistry, though both deadcH<sup>-</sup> and deadc<sup>2-</sup> have been shown to be useful counteranions for crystallization of organometallic cations<sup>5</sup> and structural studies of complexes formed with Co<sup>II</sup> and Cd<sup>II</sup> have shown deadc<sup>2-</sup> to be suited to the formation of coordination polymers.<sup>6</sup> The acid deadcH<sub>2</sub> is chiral and while several methods are known for its resolution into enantiomers,<sup>7</sup> only in one instance has resolved deadc<sup>2-</sup> been used as a ligand in an osmium complex, not structurally characterized, employed as a catalyst for asymmetric synthesis.<sup>8</sup>

The ligand deadc<sup>2-</sup> has carboxylate donor groups in a similar spatial disposition to those of *trans*-1,2-



Scheme 1 The diacid deadcH<sub>2</sub>

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cyclohexanedicarboxylate (t-1,2-chdc<sup>2-</sup>). The latter ligand has been shown, both as its racemate and pure enantiomers, to give uranyl ion complexes of several forms including closed tetranuclear species as well as mixed-ligand complexes,9 with one of them involving both anionic  $(R, R-t-1, 2-chdc^{2-})$  and zwitterionic carboxylate donors.9g Modelling of the isolated diacid deadcH<sub>2</sub> (at MM2 level using Chem3D<sup>10</sup>) provides a disposition of the carboxyl groups (C···C distance, 3.52 Å) intermediate between those found experimentally, for example, in crystal structures of the racemic acid  $(3.62 \text{ Å})^{1a}$ and its S,S isomer  $(3.40 \text{ Å})^{1b}$  (as its hexane adduct), indicating, along with moderate torsion angle differences, that there is rather limited flexibility of the molecule. The same distance and torsion angles found in organometallic complexes of deadcH<sup>-</sup> and deadc<sup>2-</sup> anions not directly coordinated to a metal ion<sup>5</sup> show that deprotonation has little obvious effect on this degree of flexibility. The carboxylate C…C separation for deadc<sup>2-</sup> here is intermediate between those of the diaxial isomer of t-1,2-chdc<sup>2-</sup> (3.86–3.88 Å)<sup>9c</sup> and its more commonly observed diequatorial form  $(2.96-3.12 \text{ Å})^{9a,b}$  but closer to the former, indicating that it should favour polymer formation over that of closed oligomers. In the known transition metal ion complexes of deadc<sup>2-</sup>, which are in fact both polymers,<sup>6</sup> the values for this separation (3.35-3.54 Å) fall within the range of the values given above. Another obvious difference between deadc<sup>2-</sup> and t-1,2-chdc<sup>2-</sup> is the much greater bulkiness of the dibenzobarrelene compared to the cyclohexane platform, and the presence of aromatic rings with the variety of interactions that they entail.

Although an impressive variety of polycarboxylates have been used for the synthesis of coordination polymers and frameworks based on the uranyl ion,11 there has been no report of a complex involving deadc<sup>2-</sup>. Since the peculiarities of this ligand, *i.e.* both the closeness of its coordination sites to those of 1,2-chdc<sup>2-</sup>, and the hydrophobicity of the dibenzobarrelene platform, led us to anticipate novel types of arrangements, we synthesized eight uranyl ion complexes involving either rac- (racemic) or R,R-deadc2- under solvohydrothermal conditions. As in previous work, we have used various additional species, either neutral in the form of coordinating cosolvents, or cationic as structure-directing counterions. Additional metal cations were also used in two cases, silver(I) for its ability to interact with aromatic rings, and nickel(II) in association with 2,2':6',2"-terpyridine-4'carboxylic acid (tpycH), a combination producing the zwitterionic, "expanded" ligand Ni(tpyc)<sub>2</sub>. All these complexes have been characterized by their crystal structure and emission properties in the solid state.

### Experimental

#### Synthesis

**Synthesis of the ligands.** All chemicals were purchased from Sigma-Aldrich and Tokyo Kasei and used without further purification. The <sup>1</sup>H NMR spectrum of deadcH<sub>2</sub> was

recorded on a Bruker AVANCE III HD 400 (400.13 MHz for <sup>1</sup>H) spectrometer at ambient temperature.

11R\*,12R\*-9,10-Dihydro-9,10-ethanoanthracene-11,12-

*dicarboxylic acid (rac-deadcH*<sub>2</sub>). This racemate was prepared by a literature method<sup>12</sup> (see ESI† for details). <sup>1</sup>H NMR (DMSO-d<sub>6</sub>):  $\delta$  7.42–7.40 (m, 2H), 7.28–7.26 (m, 2H), 7.14–7.07 (m 4H), 4.73 (s, 2H), 3.13 (s, 2H) (Fig. S1†).

(11R,12R)-(+)-9,10-Dihydro-9,10-ethanoanthracene-11,12dicarboxylic acid (R,R-deadcH<sub>2</sub>). The racemate was resolved into each enantiomer as in the literature by formation of its diastereomeric inclusion complexes with S-proline<sup>7c</sup> (see ESI<sup>†</sup> for details).

**Synthesis of the complexes. 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%),  $Ni(NO_3)_2 \cdot 6H_2O$ , and AgNO<sub>3</sub> were purchased from Prolabo; 2,2':6',2"-terpyridine-4'-carboxylic acid (tpycH) and guanidinium nitrate were from Alfa-Aesar. 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.

 $[UO_2(deadc)]$ -1.5CH<sub>3</sub>CN (1). deadcH<sub>2</sub> (15 mg, 0.05 mmol),  $[UO_2(NO_3)_2(H_2O)_2]$ ·4H<sub>2</sub>O (25 mg, 0.05 mmol), and guanidinium nitrate (12 mg, 0.10 mmol) were dissolved in a mixture of water (0.4 mL) and acetonitrile (0.2 mL). Yellow crystals of complex 1 were obtained within one week (11 mg, 36%). Chemical analysis indicates the loss of 0.5 acetonitrile molecule with respect to the formula derived from crystal structure determination. Anal. calcd for C<sub>20</sub>H<sub>15</sub>NO<sub>6</sub>U: C, 39.81; H, 2.51; N, 2.32. Found: C, 39.50; H, 2.50; N, 2.26%.

 $[H_2NMe_2]_2[(UO_2)_2(deadc)_3]\cdot 2H_2O$  (2). deadcH<sub>2</sub> (15 mg, 0.05 mmol) and  $[UO_2(NO_3)_2(H_2O)_2]\cdot 4H_2O$  (25 mg, 0.05 mmol) were dissolved in a mixture of water (0.5 mL) and *N*,*N*-dimethylformamide (0.2 mL). Yellow crystals of complex **2** were obtained overnight (19 mg, 49%). Anal. calcd for  $C_{58}H_{56}N_2O_{18}U_2$ : C, 45.09; H, 3.65; N, 1.81. Found: C, 44.76; H, 3.67; N, 1.82%.

 $[H_2NMe_2]_4[(UO_2)_2(O)(R,R-deadc)_2]_2$  (3). R,R-deadcH<sub>2</sub> (15 mg, 0.05 mmol) and  $[UO_2(NO_3)_2(H_2O)_2]\cdot 4H_2O$  (25 mg, 0.05 mmol) were dissolved in a mixture of water (0.5 mL) and *N,N*-dimethylformamide (0.2 mL). Yellow crystals of complex 3 were obtained within four days (7 mg, 23%). Chemical analysis indicates the presence of about three water molecules in excess of the formula derived from crystal structure determination. Anal. calcd for C<sub>80</sub>H<sub>80</sub>N<sub>4</sub>O<sub>26</sub>U<sub>4</sub> + 3H<sub>2</sub>O: C, 38.14; H, 3.44; N, 2.22. Found: C, 38.06; H, 3.22; N, 2.15%.

 $[(UO_2)_2(deadc)(deadcH)(NO_3)Ni(tpyc)_2]$ ·CH<sub>3</sub>CN·2H<sub>2</sub>O (4). deadcH<sub>2</sub> (15 mg, 0.05 mmol),  $[UO_2(NO_3)_2(H_2O)_2]$ ·4H<sub>2</sub>O (25 mg, 0.05 mmol), Ni(NO<sub>3</sub>)<sub>2</sub>·6H<sub>2</sub>O (10 mg, 0.03 mmol), and tpycH (14 mg, 0.05 mmol), were dissolved in a mixture of water (0.6 mL) and acetonitrile (0.2 mL). Orange crystals of complex 4 were obtained within four days (40 mg, 85%). Anal. calcd for C<sub>70</sub>H<sub>52</sub>N<sub>8</sub>NiO<sub>21</sub>U<sub>2</sub>: C, 44.82; H, 2.79; N, 5.97. Found: C, 44.20; H, 2.84; N, 5.86%.

 $[UO_2(deadc)(DMA)]$  (5). deadcH<sub>2</sub> (15 mg, 0.05 mmol) and  $[UO_2(NO_3)_2(H_2O)_2]$ ·4H<sub>2</sub>O (25 mg, 0.05 mmol) were dissolved in a mixture of water (0.5 mL) and *N*,*N*-dimethylacetamide (0.2 mL). Yellow crystals of complex 5 were obtained overnight (22 mg, 68%). Anal. calcd for C<sub>22</sub>H<sub>21</sub>NO<sub>7</sub>U: C, 40.69; H, 3.26; N, 2.16. Found: C, 40.71; H, 3.27; N, 2.10%.

*[UO<sub>2</sub>(deadc)]* (6). deadcH<sub>2</sub> (15 mg, 0.05 mmol),  $[UO_2(NO_3)_2(H_2O)_2]$ ·4H<sub>2</sub>O (25 mg, 0.05 mmol), and CsNO<sub>3</sub> (20 mg, 0.10 mmol) were dissolved in a mixture of water (0.6 mL) and acetonitrile (0.2 mL). Yellow crystals of complex 6 were obtained within three days (20 mg, 71%). Chemical analysis indicates the presence of 0.5 acetonitrile in excess of the formula derived from crystal structure determination. Anal. calcd for  $C_{18}H_{12}O_6U + 0.5CH_3CN$ : C, 39.16; H, 2.33; N, 1.20. Found: C, 39.16; H, 2.36; N, 1.12%.

 $[PPh_4]_2[(UO_2)_2(R,R-deadc)_3]$  (7). R,R-deadcH<sub>2</sub> (15 mg, 0.05 mmol),  $[UO_2(NO_3)_2(H_2O)_2]$ ·4H<sub>2</sub>O (25 mg, 0.05 mmol), and PPh<sub>4</sub>Br (21 mg, 0.05 mmol) were dissolved in a mixture of water (0.5 mL) and *N*,*N*-dimethylacetamide (0.2 mL). Yellow crystals of complex 7 were obtained overnight (15 mg, 43%). Anal. calcd for C<sub>102</sub>H<sub>76</sub>O<sub>16</sub>P<sub>2</sub>U<sub>2</sub>: C, 58.46; H, 3.66. Found: C, 58.11; H, 3.71%.

 $[(UO_2)_2Ag_2(deadc)_3(CH_3CN)_2]\cdot 0.5H_2O$  (8). deadcH<sub>2</sub> (15 mg, 0.05 mmol),  $[UO_2(NO_3)_2(H_2O)_2]\cdot 4H_2O$  (25 mg, 0.05 mmol), and AgNO<sub>3</sub> (17 mg, 0.10 mmol) were dissolved in a mixture of water (0.6 mL) and acetonitrile (0.2 mL). Yellow crystals of complex 8 were obtained within four days (20 mg, 46%). Chemical analysis indicates the possible presence of one water molecule in excess of the formula derived from crystal structure determination. Anal. calcd for C<sub>58</sub>H<sub>43</sub>Ag<sub>2</sub>N<sub>2</sub>O<sub>16.5</sub>U<sub>2</sub> + H<sub>2</sub>O: C, 40.00; H, 2.60; N, 1.61. Found: C, 39.78; H, 2.48; N, 1.38%.

#### Crystallography

Data collections were performed at 100(2) K on a Bruker D8 Quest diffractometer using an Incoatec Microfocus Source (IµS 3.0 Mo) and a PHOTON III area detector, and operated with APEX3.<sup>13</sup> The data were processed with SAINT,<sup>14</sup> and empirical absorption corrections were made with SADABS.<sup>15</sup> The structures were solved by intrinsic phasing with SHELXT,<sup>16</sup> and refined by full-matrix least-squares on  $F^2$  with SHELXL,<sup>17</sup> using the ShelXle interface.<sup>18</sup> The hydrogen atoms bound to oxygen and nitrogen atoms in 2 and 4 were retrieved from residual electron density maps and they were refined with geometric restraints. All other 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 CH<sub>3</sub>). In 1, one of the solvent acetonitrile molecules has been given an occupancy factor of 0.5 to account for its closeness to its image by symmetry. In 3, one dimethylammonium cation is disordered over two positions which have been refined with occupancy parameters constrained to sum to unity and with restraints on bond lengths and displacement parameters. In 8, the solvent water molecule has been given partial occupancy so as to retain an acceptable displacement parameter. For compounds 3 and 4, the SOUEEZE<sup>19</sup> software was used to subtract the contribution of disordered solvent molecules to the structure factors; in compound 3, about 15 electrons per formula unit were thus added in the solventaccessible volume, which is less than the number corresponding to the three water molecules found from elemental analysis (see above), the volume being however sufficiently large to accommodate them (121 Å<sup>3</sup> per formula unit); in compound 4, about 25 electrons per formula unit were added in a volume of 118 Å<sup>3</sup>, possibly corresponding to about 2.5 water molecules, these being apparently lost during drying of the crystals before elemental analysis. Crystal data and structure refinement parameters are given in Table 1. Drawings were made with ORTEP-3,20 and VESTA,21 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 mm<sup>-1</sup> 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. Gaussian deconvolution of the spectrum of 4 was made with the Origin software. 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

All the complexes presently described were obtained from reaction mixtures in which the  $U^{VI}$ /deadcH<sub>2</sub> ratio was 1:1 and in 5 of the 8 crystalline solids obtained, this stoichiometry was retained, though in a considerable variety of structures. If it is assumed that crystal deposition is not kinetically controlled, the crystals deposited have to be simply those of the least soluble component of the reaction mixture. This would explain why, when additional non-coordinating cations (H<sub>2</sub>NMe<sub>2</sub><sup>+</sup>, PPh<sub>4</sub><sup>+</sup>) are present in the mixture, it is possible to obtain crystals incorporating these cations in which  $U^{VI}$  is present in an anionic component

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Table 1 Crystal data and structure refinement details

	1	2	3	4	ſŪ	9	7	8
Chemical formula <i>M/9</i> mol <sup>-1</sup>	$C_{21}H_{16.5}N_{1.5}O_6U$ 623.89	$C_{58}H_{56}N_2O_{18}U_2$ 1545.10	$C_{80}H_{80}N_4O_{26}U_4$ 2465.60	$C_{70}H_{52}N_8NiO_{21}U_2$ 1875.96	C <sub>22</sub> H <sub>21</sub> NO <sub>7</sub> U 649.43	$C_{18}H_{12}O_{6}U$ 562.31	$C_{102}H_{76}O_{16}P_2U_2$ 2095.62	$C_{58}H_{43}Ag_2N_2O_{16.5}U_2$ 1723.74
Crystal system	Triclinic	Monoclinic	Orthorhombic	Triclinic	Monoclinic	Monoclinic	Orthorhombic	Monoclinic
Space group	$Par{1}$	C2/c	$P2_{1}2_{1}2$	$P\bar{1}$	C2/c	$P2_1/c$	$P2_{1}2_{1}2$	C2/c
ajâ	8.7652(4)	15.7139(5)	17.5195(6)	8.6348(3)	28.9120(13)	8.4166(3)	23.3097(6)	15.4596(4)
$b/ m \AA$	10.7608(5)	15.0080(5)	26.5815(7)	14.3691(5)	11.9986(5)	21.8953(7)	13.3655(3)	15.1930(4)
$c/{ m \AA}$	11.2060(5)	22.3323(6)	8.8808(3)	27.8539(11)	13.6561(5)	8.4203(3)	13.5379(3)	21.8735(5)
$\alpha^{ m o}$	103.3336(18)	06	60	93.9567(17)	06	06	60	60
<i>B</i> /°	92.5999(18)	96.4976(11)	06	98.1776(17)	95.2025(17)	93.6517(12)	06	95.5571(9)
y/o	107.6526(17)	. 06	90	94.8470(16)	06	06	90	<u> </u>
$V/{\rm \AA}^3$	972.43(8)	5232.9(3)	4135.7(2)	3397.1(2)	4717.8(3)	1548.58(9)	4217.68(17)	5113.5(2)
Z	2	4	2	2	8	4	2	4
Reflections collected	50533	85794	53405	137913	63094	19826	96 094	90 293
Independent reflections	3696	6774	7839	12911	4484	3990	10917	6585
Observed reflections $[I > 2\sigma(I)]$	3431	6126	6652	$12\ 122$	4047	3814	10575	6542
$R_{ m int}$	0.061	0.057	0.118	0.061	0.063	0.039	0.051	0.043
Parameters refined	282	375	533	935	283	226	551	371
$R_1$	0.021	0.019	0.047	0.036	0.034	0.016	0.019	0.018
$wR_2$	0.046	0.043	0.106	0.080	0.089	0.039	0.041	0.047
S	1.151	1.059	1.035	1.208	1.174	1.119	1.038	1.073
$\Delta  ho_{ m min}$ /e Å $^{-3}$	-1.73	-0.84	-1.07	-2.32	-1.31	-1.07	-0.47	-0.88
$\Delta  ho_{ m max}$ /e Å <sup>-3</sup>	0.95	0.69	1.43	2.07	2.16	1.28	0.44	1.56
Flack parameter			0.016(17)				-0.012(4)	

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where the ratio U/deadc<sup>2-</sup> is 1:1.5 (complexes 2 and 7). Where an additional cation is present, such as  $C(NH_2)_3^+$  in the synthesis of complex 1 and  $Cs^+$  in that of complex 6, but is not incorporated in the precipitated crystals, any influence it may have on the solution equilibria is uncertain but it does not prevent the formation of crystals of a neutral 1:1 complex.

In general, it is unsurprising to find that different solvates may have different structures and this is seen when complex 5 is compared to 1 and 6 (see structural discussion ahead), even though all have a U/deadc<sup>2-</sup> ratio of 1:1. Where Ag<sup>I</sup> is the additional cation and interacts, along with UVI, with the deadc2- ligand, its incorporation in the isolated crystals (complex 8) leads to a  $U/deadc^{2-}$ ratio of 1:1.5, formally because its insolubility is favoured by cation...anion attraction. In contrast, where Ni<sup>II</sup> is the additional cation but is cloaked within an additional ligand (tpyc<sup>-</sup>) so as to be a neutral (zwitterionic) component not in direct interaction with deadc<sup>2-</sup>, the 1:1 ratio is regained (complex 4). In the case of complexes 2 and 3, the cation is generated through hydrolysis of the cosolvent N,N-dimethylformamide (DMF), a reaction which must result in buffering of the solution pH and thus modification of the solution equilibria. These two complexes display a marked difference in composition, although they were prepared under identical conditions except that for the former the deadcH<sub>2</sub> reactant was the racemate and for the latter the R,R enantiomer. The structure of 2 (see ahead) shows both enantiomers to be present, *i.e.* that the complex is racemic, so clearly the same structure could not be obtained with a single enantiomer. It appears that a tetranuclear species present as a result of the partial hydrolysis of uranyl ion under the reaction conditions forms the least soluble species involving coordination to R,R-deadc<sup>2-</sup>.

#### Crystal structures

The complex  $[UO_2(deadc)]$ ·1.5CH<sub>3</sub>CN (1), obtained with acetonitrile as organic cosolvent, is the simplest case in this series. The unique uranyl cation is  $\kappa^2 O_{,O'}$ -chelated by one carboxylate group from one ligand, chelated between the two carboxylate groups of a second ligand (seven-membered chelate ring) and bound to one more carboxylate oxygen atom from a third ligand (Fig. 1). The uranium environment is thus pentagonal-bipyramidal [U-O(oxo), 1.757(3) and 1.762(3) Å; U-O(carboxylate), 2.474(3) and 2.506(3) Å for the chelating group, 2.304(3)-2.404(3) Å for the others]. The deadc<sup>2-</sup> ligand has thus one carboxylate group chelating and bridging in the  $\mu_2$ - $\kappa^2 O$ , O': $\kappa^1 O$  mode, and the other syn/anti  $\mu_2$ - $\kappa^1 O$ : $\kappa^1 O'$ -bridging. Both metal and ligand are thus threecoordinated (3-c) nodes, and the coordination polymer formed is monoperiodic and directed along [100]. The uranium centers are assembled into centrosymmetric, edgesharing dimers arranged into ribbons with flanking dihydroanthracenyl groups.

The carboxylate C···C separation is 3.445(6) Å, showing that the separation in phthalate (close to 3.1 Å),<sup>23</sup> for which such chelate rings are well known,<sup>23,24</sup> does not represent a limit for 7-membered chelate ring formation on uranyl ion. The small variations of the torsional parameters in the aliphatic part of the ligand are indicative of the rigidity of deadc<sup>2-</sup> (Table S1, ESI<sup>†</sup>). The chains lie side-by-side in sheets parallel to (011) in such a way that enantiomeric dihydroanthracenyl entities confront one another, thus creating cavities. This feature of the structure has a clear similarity to those seen in inclusion complexes of deadcH<sub>2</sub> alone.<sup>1-4</sup> The acetonitrile molecules are involved in multiple weak interactions. One parallel-displaced  $\pi$ -stacking interaction between two aromatic rings pertaining to adjacent chains may be present [centroid…centroid distance, 3.870(3)



Fig. 1 (a) View of complex 1 with displacement ellipsoids shown at the 50% probability level. Solvent molecules and hydrogen atoms are omitted. Symmetry codes: i = 1 - x, 1 - y, 1 - z; j = 2 - x, 1 - y, 1 - z. (b) and (c) Two views of the monoperiodic assembly showing uranium coordination polyhedra. (d) Packing with chains viewed edge-on.

Å; dihedral angle,  $0^{\circ}$ ] and the Kitaigorodsky packing index (KPI, evaluated with PLATON<sup>25</sup>) is 0.66 (with disordered acetonitrile molecule excluded).

A dicarboxylate/U ratio of 3:2 commonly results in the formation of  $[tris(\kappa^2 O, O'-carboxylate)UO_2]^-$  centres of hexagonal-bipyramidal geometry<sup>11</sup> but yet another exception to this occurs in the complex [H<sub>2</sub>NMe<sub>2</sub>]<sub>2</sub>[(UO<sub>2</sub>)<sub>2</sub>(deadc)<sub>3</sub>]·2H<sub>2</sub>-O (2). Here, the monoperiodic polymer present has unique U<sup>VI</sup> centres with a pentagonal-bipyramidal coordination geometry and an environment similar to that in 1 [U–O(oxo), 1.7740(17) and 1.7811(17) Å; U-O(carboxylate), 2.4368(17) and 2.5298(16) Å for the chelating group, 2.3354(17)-2.3656(17) Å for the others] (Fig. 2). The deadc2- ligands adopt two different coordination modes, both of them bridging. The first ligand is involved in 7-membered chelate ring formation and further bridging  $(\mu_2 - \kappa^1 O) \cdot \kappa^1 O' \cdot \kappa^1 O'$  mode), and the second, which has twofold rotation symmetry, is  $bis(\kappa^2 O, O'$ chelating). Formation of a 7-membered chelate ring shows again deadc<sup>2-</sup> to have some resemblance to t-1,2-chdc<sup>2-.9</sup>

The carboxylate C···C distances are 3.468(3) Å in the ligand forming the 7-membered ring and 3.713(4) in the other, showing again that deadc<sup>2–</sup> does have some flexibility, if considerably less than that of t-1,2-chdc<sup>2–</sup>. The dimethylammonium counterion, produced *in situ* from hydrolysis of DMF, is hydrogen bonded to the uncoordinated atom O4 and to the water molecule. The metal cation is here also a 3-c node while the ligands are simple edges, and the coordination polymer formed is monoperiodic and directed along [101]. Packing of the zigzag polymer strands involves some degree of interdigitation within the layers parallel to (010). One parallel-displaced  $\pi$ -stacking interaction between two aromatic rings pertaining to adjacent chains may be

present here also [centroid···centroid distance, 4.0874(15) Å; dihedral angle,  $6.26(12)^{\circ}$ ], as well as two CH··· $\pi$  edge-to-face interactions involving the same two ligands and resulting in their dimerization. This interpenetration produces a very compact packing with no evidence of significant porosity (KPI, 0.72).

The complex  $[H_2NMe_2]_4[(UO_2)_2(O)(R,R-deadc)_2]_2$ (3). produced under conditions identical to those giving 2 from the racemic diacid, has a structure which shows that chirality here is in fact an important influence. Hydrolysis of DMF and the concomitant buffering of the reaction mixture may explain the partial hydrolysis of uranyl ion to give a slightly twisted tetranuclear uranate cluster. The latter has twofold rotation symmetry and it is built around two  $\mu_3$ -oxo anions and bounded by an up-down alternating garland of four R,R $deadc^{2-}$  ligands (Fig. 3). The cluster, a coordination oligomer only, is an example of a U4O2 motif with additional peripheral ligands which has previously been observed in several cases,<sup>23,24i,26</sup> with in particular one instance involving t-1,2-chdc<sup>2-</sup> in the diequatorial conformation.<sup>9e</sup> U1 is part of two 7-membered chelate rings and bound to one oxo anion (pentagonal-bipyramidal environment), and U2 is  $bis(\kappa^2 O, O'$ chelated) and bound to two oxo anions (hexagonalbipyramidal environment) [U–O(oxo), 1.748(12)–1.783(13) Å; U-O(carboxylate), 2.526(10)-2.646(11) Å for the chelating groups, 2.315(10)-2.439(11) Å for the others; U-O( $\mu_3$ -oxo), 2.243(10)-2.278(11) Å]. The sum of the three U-O-U angles around the oxo atom O13 is 353.4°, indicating that the three U-O bonds around this atom are nearly coplanar. The sum of bond valence parameters calculated with  $PLATON^{25}$  for atom O13 is 1.99, thus confirming that it is an oxo and not a hydroxo anion. The uranium coordination polyhedra share



Fig. 2 (a) View of complex 2 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 = 1/2 - x, 1/2 - y, 1 - z; j = 1 - x, y, 1/2 - z. (b) and (c) Two views of the monoperiodic assembly showing uranium coordination polyhedra. (d) Packing with chains viewed edge-on.



Fig. 3 (a) View of compound 3 with displacement ellipsoids shown at the 30% probability level. The disordered counterion and carbon-bound hydrogen atoms are omitted, and hydrogen bonds are shown as dashed lines. Symmetry code: i = 1 - x, 1 - y, z. (b) Side view of the tetranuclear complex. (c) and (d) Two views of the packing with counterions omitted.

either two or three edges with their neighbours, giving one of the more compact arrangements among those found in uranyl tetranuclear clusters.<sup>11c</sup>

While there are two inequivalent ligands in each cluster, the differences between them are very minor, the carboxylate  $C\cdots C$  separation being 3.46(2) Å for both, showing again that the conformation of deadc<sup>2-</sup> is quite insensitive to its environment. The tetranuclear anions lie in stacks aligned parallel to [001], with two disordered H<sub>2</sub>NMe<sub>2</sub><sup>+</sup> cations sandwiched between every two clusters. The ordered H<sub>2</sub>NMe<sub>2</sub><sup>+</sup> cation forms two bifurcated hydrogen bonds with two carboxylate groups bound to the same uranyl group. While no significant  $\pi$ -stacking interaction is present, each of the two methyl groups of the ordered counterion forms two  $CH\cdots\pi$  interactions involving the two rings of one ligand. These counterions are thus included in the cavity formed by the concave parts of two dihydroanthracenyl units from adjacent cluster columns (KPI, 0.65).

To explore the consequences of the combination of deadc<sup>2-</sup> with carboxylate zwitterions, [Ni(tpyc)<sub>2</sub>], in which tpyc<sup>-</sup> is 2,2':6',2"-terpyridine-4'-carboxylate, was used to synthesize [(UO<sub>2</sub>)<sub>2</sub>(deadc)(deadcH)(NO<sub>3</sub>)Ni(tpyc)<sub>2</sub>]·CH<sub>3</sub>CN·2H<sub>2</sub>-O (4), providing the first instance in the present study of a complex involving both deadc<sup>2-</sup> and deadcH<sup>-</sup>. The two independent uranium atoms are both tris( $\kappa^2 O$ ,O'-chelated) and in hexagonal-bipyramidal environments. While U1 is bound to two carboxylate groups from two deadc<sup>2-</sup> ligands and one from Ni(tpyc)<sub>2</sub>, U2 is bound to one carboxylate from deadcH<sup>-</sup>, one from Ni(tpyc)<sub>2</sub>, and one nitrate anion [U-

O(oxo), 1.765(4)-1.779(4) Å; U-O(carboxylate), 2.410(3)-2.523(4) Å; U-O(nitrate), 2.500(4) Å for both] (Fig. 4). Complex 4 is a monoperiodic polymer in which  $\{UO_2(deadcH)(NO_3)\}$ Ni(tpyc)<sub>2</sub>} units can be considered as pendent substituents on a  $\{UO_2(deadc)\}\$  main chain directed along [100]. However, if hydrogen bonding between the carboxylic group of deadcH<sup>-</sup> and the uranyl oxo group O4 is recognized, the polymer has a ladder-like, double stranded form. The essential structural unit is a U<sub>4</sub>Ni<sub>2</sub> metallacycle closely similar to that seen in several other mixed-ligand complexes involving the  $[Ni(tpyc)_2]$  zwitterion.<sup>9g</sup> There is also a parallel the complex  $[(UO_2)_2(1,2-pda)(1,2-pdaH)]$ with Ni(tpyc)<sub>2</sub>(NO<sub>3</sub>)]·CH<sub>3</sub>CN (1,2-pda<sup>2-</sup> = 1,2-phenylenediacetate),  $^{9g}$ considered as a monoperiodic meander polymer where hexagonal-bipyramidal  $\boldsymbol{U}^{\text{VI}}$  centres bound to the two nonzwitterion species 1,2-pda<sup>2-</sup> and 1,2-pdaH<sup>-</sup> alternate along the chain rather than forming separate chains as here. Unlike 1,2-pda<sup>2-</sup> and 1,2-pdaH<sup>-</sup>, which can adopt chiral but labile conformations,9g,27 deadc2- and deadcH- exist as kinetically stable enantiomers (as shown indirectly by the isolation of the cis and trans isomers,5 as well as through chiral resolution<sup>7</sup>). The carboxylate C···C separations are 3.673(7) Å in deadc<sup>2-</sup> and 3.468(7) Å in deadcH<sup>-</sup>, much longer than those (~2.97 Å) found for the inequivalent, non-disordered t-R,R-chdc<sup>2-</sup> units in the complex  $[(UO_2)_4(t-R,R-chdc)_4 Ni_2(tpyc)_4$ <sup>9g</sup> where the conformation is diequatorial, indicating that deadc2- and deadcH- could be considered as rigidified analogues of diaxial t-1,2-chdc<sup>2-</sup>. The quite different compositions and structures of the complexes of



**Fig. 4** (a) View of compound **4** 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 = x + 1, *y*, *z*; j = x - 1, *y*, *z*. (b) The heterometallic monoperiodic assembly with uranium coordination polyhedra yellow and those of nickel blue, and hydrogen bonds shown as dotted blue lines. (c) Packing with ribbon-like chains viewed end-on.

the anthracene and cyclohexane derivatives must be at least in part a reflection of this conformational factor. Although there are examples for *t*-1,2-chdc<sup>2-</sup> where chirality does not seem to be an important influence,<sup>9*a*</sup> complexes **2** and **3** show that this is not necessarily the case with deadc<sup>2-</sup>. Neighbouring Ni(tpyc)<sub>2</sub> units in **4** are possibly associated by one intrachain parallel-displaced  $\pi$ -stacking interaction [centroid…centroid distance, 3.607(3) Å; dihedral angle, 9.4(2)°], while two weaker ones involving deadcH<sup>-</sup> anions and tpyc<sup>-</sup> may be found between adjacent chains. Both intra- and interchain CH… $\pi$  interactions are present as well, and the packing has a KPI of 0.67.

Direct coordination of a cosolvent molecule, here *N*,*N*-dimethylacetamide (DMA), to  $U^{VI}$  produces a significant structural change in  $[UO_2(deadc)(DMA)]$  (5). The unique

uranium centre is  $\kappa^2 O_{,O'}$ -chelated by one carboxylate group and bound to two more carboxylate donors from two different ligands, and to the DMA molecule (Fig. 5). The uranium environment is thus pentagonal-bipyramidal [U-O(oxo), 1.769(5) and 1.773(5) Å; U–O(carboxylate), 2.421(5) and 2.462(5) Å for the chelating group, 2.316(5) and 2.320(5) Å for the others; U–O(DMA), 2.374(5) Å]. The deadc<sup>2–</sup> ligand is chelating and *syn/anti* bridging ( $\mu_3$ - $\kappa^2 O, O'; \kappa^1 O'': \kappa^1 O'''$ ), and the carboxylate C···C separation [3.456(9) Å] is essentially unchanged, as are the torsional parameters. Both metal and ligand are thus 3-c nodes, and the coordination polymer formed, which is diperiodic and parallel to (100), has the  $\{4\cdot8^2\}$  point symbol and the common **fes** topological type. When viewed down [001], the thick layers display a central, zigzag layer of uranyl cations surrounded on both sides by dihydroanthracenyl units pointing outward, each surface comprising a racemic mixture of ligands. Enantiomeric ligands in adjacent sheets are united pairwise through one parallel-displaced  $\pi$ -stacking interaction [centroid...centroid distance, 3.808(4) Å; dihedral angle, 0°] and two edge-to-face  $CH\cdots\pi$  interactions. The KPI of 0.62 shows that disordered and unresolved solvent molecules are possibly present.

Although its composition differs from that of 1 only by the latter incorporating acetonitrile solvent molecules, the complex [UO<sub>2</sub>(deadc)] (6) has a different structure, possibly as a result of subtle effects of the additional reagents, guanidinium nitrate for 1 and cesium nitrate here, although these species are not present in the final compound (the difference in concentration of the reactants may also play a part). The single, hexagonal-bipyramidal uranium atom is  $\kappa^2 O, O'$ -chelated by two carboxylate groups (one of them very unsymmetrical) and bound to two more carboxylate donors in trans positions [U-O(oxo), 1.7586(17) and 1.7652(17) Å; U-O(carboxylate), 2.4440(19)-2.5874(17) Å for the chelating groups, 2.4582(17) and 2.4727(17) Å for the others] (Fig. 6). The ligand is bis(chelating/bridging) [bis( $\mu_2$ - $\kappa^2 O$ , O': $\kappa^1 O$ ] and is thus a 4-c node, as the metal cation. The diperiodic, uninodal coordination polymer formed is parallel to (010) and it has the  $\{4^4 \cdot 6^2\}$  point symbol and the simple sql topological type. The layers contain strands of edge-sharing UO<sub>6</sub> groups running parallel to [001] which are crosslinked by the ligands. Here also, the dihydroanthracenyl units form upper and lower faces to each sheet. The packing displays dimerization of enantiomeric ligands pertaining to adjacent sheets similar to that found in 5, with both one paralleldisplaced  $\pi$ -stacking interaction [centroid…centroid distance, 3.7429(15) Å; dihedral angle, 0°] and two edge-to-face CH $\cdots\pi$ interactions. The packing is however more compact here and no solvent-accessible space is present (KPI, 0.71).

In the enantiomerically pure complex  $[PPh_4]_2[(UO_2)_2(R,R-deadc)_3]$  (7), the uranium atom is tris( $\kappa^2 O, O'$ -chelated) by three carboxylate groups  $[U-O(\infty o), 1.776(3) \text{ and } 1.781(3) \text{ Å}; U-O(carboxylate), 2.458(2)-2.484(2) \text{ Å}]$  (Fig. 7). As often found in complexes in which the uranyl cation is a 3-c node and the ligand a simple edge, the diperiodic network formed, parallel to (001) has the  $\{6^3\}$  point symbol and the **hcb** topological

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**Fig. 5** (a) View of compound **5** with displacement ellipsoids shown at the 50% probability level and hydrogen atoms omitted. Symmetry codes: i = x, 1 - y, z + 1/2; j = 3/2 - x, y - 1/2, 3/2 - z; k = x, 1 - y, z - 1/2; l = 3/2 - x, y + 1/2, 3/2 - z. (b) View of the **fes** diperiodic assembly. (c) Packing with layers viewed edge-on. (d) Nodal representation of the network (yellow, uranium nodes; blue, deadc<sup>2-</sup> nodes; view down [100] with [001] horizontal).

type. The sheets are strongly puckered, with PPh<sub>4</sub><sup>+</sup> cations embedded within due to multiple CH···O interactions, and there is no significant free space (KPI, 0.67). In the case of complexes of the enantiomers of *t*-1,2-chdc<sup>2-</sup> prepared under conditions of 2:3 U/dicarboxylate stoichiometry,<sup>9c</sup> these materials have a 1:1 stoichiometry, though the ratio U/ carboxylate of 1:3 is achieved due to the coordination of formate deriving from DMF hydrolysis; they are, nonetheless, monoperiodic polymers in which the *t*-1,2-chdc<sup>2-</sup> ligands have diaxial conformations with inter-carboxylate C···C separations of 3.86–3.88 Å, significantly longer than those in 7, 3.672(5) and 3.575(7) Å. The rigidity of deadc<sup>2-</sup> thus appears to severely limit the extent to which it may be considered an analogue of diaxial *t*-1,2-chdc<sup>2-</sup>.

The second heterometallic complex in this series,  $[(UO_2)_2-Ag_2(deadc)_3(CH_3CN)_2]\cdot 0.5H_2O$  (8), crystallizes in the same space group as complex 2 and with very close unit cell parameters, so that these two species can be considered to be isomorphous, although not isostructural. The pentagonal-bipyramidal uranium atom is  $\kappa^2 O_i O'$ -chelated by one carboxylate group, chelated by two groups of another ligand (7-membered ring), and bound to one more carboxylate donor from a third ligand [U-O(oxo), 1.7732(17) and 1.7778(17) Å; U-O(carboxylate), 2.4214(17) and 2.5071(17) Å for the chelating group, 2.3107(17)-2.3680(17) Å for the others] (Fig. 8). The metal centre is thus a 3-c node, while both ligands are only bound to two uranium atoms, the unsymmetrical one in a bridging mode and that with twofold

rotation symmetry in the bis(chelating) mode. Ignoring interactions with Ag<sup>I</sup>, the monoperiodic polymer formed is essentially identical with that present in 2. As might be expected of Ag<sup>1,28</sup> its coordination sphere is irregular and heteroleptic, interactions here involving two adjacent aromatic carbon atoms, the acetonitrile nitrogen donor, one carboxylate donor and possibly one water molecule (with partial occupancy) [Ag-C, 2.367(2) and 2.464(2) Å; Ag-N, 2.167(3) Å; Ag-O(carboxylate), 2.3389(17) Å; Ag-O(water), 2.493(9) Å]. While it may appear that  $Ag^{I}$  and  $H_2NMe_2^+$  are two very different cations, the latter is involved in weak interactions in addition to the obvious NH···O bond formation and these include  $CH\cdots\pi$  interactions, so that in sum the interactions of both are quite similar and this may explain the retention of the same form of the associated polymer anion. However, a difference between the weak interactions involving H<sub>2</sub>NMe<sub>2</sub><sup>+</sup> and the coordination bonds formed by silver(I) is that the latter connect the monoperiodic, uranyl-only coordination polymers into a compact triperiodic framework (KPI, 0.73). Dimerization of deadc<sup>2-</sup> ligands involving one parallel-displaced  $\pi$ -stacking interaction [centroid…centroid distance, 3.6826(13) Å; dihedral angle, 11.60(11)°], and two CH··· $\pi$  edge-to-face interactions is found here also.

Notwithstanding its closeness to t-1,2-chdc<sup>2-</sup> regarding the relative position of the two donor groups, deadc<sup>2-</sup> appears in the present series as a ligand giving only mono- and diperiodic uranyl ion complexes, except for the tetranuclear

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**Fig. 6** (a) View of compound **6** with displacement ellipsoids shown at the 50% probability level and with hydrogen atoms omitted. Symmetry codes: i = x, 3/2 - y, z - 1/2; j = x - 1, y, z; k = x - 1, 3/2 - y, z - 1/2; l = x, 3/2 - y, z + 1/2; m = x + 1, y, z; n = x + 1, 3/2 - y, z + 1/2. (b) View of the **sql** diperiodic assembly. (c) Packing with layers viewed edge-on.

species 3 formed with R,R-deadc<sup>2-</sup> and the triperiodic assembly resulting from silver-bridging of chains in 8. In particular, no cage-like complex has been obtained with deadc<sup>2-</sup>, while such species are quite common in the case of 1,2-chdc<sup>2-</sup>, the bulkiness of the dibenzobarrelene group possibly having an effect here. The weak interactions involving the dibenzobarrelene platform are also distinctive and introduce an element of novelty. The carboxylate groups being located in the half of the molecule opposite to the aromatic-lined concave side, the latter is thus naturally pointing outward from the main body of the coordination polymer, resulting in a hydrophobic coating of the chains or networks reminiscent of that found with phosphonates<sup>29</sup> or Kemp's tricarboxylate,<sup>30</sup> for example. Such an arrangement favors interactions with neighbouring polymeric units or with solvent molecules or counterions, and it may thus have a

structure-directing effect. Three of the association modes found in the present complexes are shown in Fig. 9. Interactions with the acetonitrile molecules in 1 seem to be loose at best: the methyl group of the well-ordered acetonitrile molecule is located on the border of the cavity defined by the two facing deadc<sup>2-</sup> ligands, and the Hirshfeld surface<sup>31</sup> does not reveal any significant interaction (the nitrogen atom, in contrast, is involved in two CH…N hydrogen bonds with other anions). Interactions with the  $H_2NMe_2^+$  counterion in 3 are much more extensive and they clearly illustrate the hydrophilic/hydrophobic nature of the ligand: the ammonium protons form two bifurcated hydrogen bonds with the carboxylate groups of two ligands, while the methyl groups are involved in four  $CH \cdots \pi$ interactions with the aromatic rings of two more ligands facing one another and thus defining a hydrophobic cavity. The arrangement most often found however, in complexes 2, 5, 6 and 8, is that bringing two deadc<sup>2-</sup> anions in close interaction to form a dimer held by three aromatic ... aromatic interactions, one parallel and the other two edge-to-face. It is notable that the formation of the only triperiodic framework among these complexes is due to bridging by silver(1), a cation able to interact with both the hydrophilic and hydrophobic parts of the ligand.

#### Luminescence properties

Except for 6, all the present complexes are emissive, with however strong disparities in photoluminescence quantum yields (PLQYs), these being 2% for 1 and 4, 6% for 8, 8% for 7, 12% for 3, 23% for 5 and 26% for 2. These variations might be related to differences in aromatic interactions and intermetallic distances, the presence of solvent molecules also possibly playing a part. The emission spectra measured in the solid state under excitation at 420 nm for complexes 1, 2, 4, 5, 7 and 8 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>32</sup> (Fig. 10). Complexes 1, 2, 5 and 8, all with single uranyl sites and pentagonal-bipyramidal uranium environments, have quasi-identical positions for the main four emission maxima, at 491-492, 511-513, 533-536 and 559-561 nm. These values are typical of uranyl carboxylate complexes with O<sub>5</sub> equatorial environments.<sup>33</sup> Complex 7, with a unique uranyl cation in a tris-chelated, hexagonalbipyramidal environment, has maxima at 480, 499, 521 and 544 nm, as expected for such an O<sub>6</sub> environment.<sup>33</sup> Complex 4 contains two inequivalent uranium centres, both in hexagonal-bipyramidal environments, but one of them chelated by three carboxylate groups and the other by two carboxylate and one nitrate groups, and its emission spectrum obviously results from the superposition of two emission series which can be separated by deconvolution (blue and orange lines in Fig. 11), except for the last, very weak and unresolved maximum corresponding to a wavelength near 573 nm.



**Fig. 7** (a) View of compound 7 with displacement ellipsoids shown at the 50% probability level and with hydrogen atoms omitted. Symmetry codes: i = 3/2 - x, y - 1/2, 2 - z; j = 1 - x, 1 - y, z; k = 3/2 - x, y + 1/2, 2 - z. (b) View of the **hcb** diperiodic assembly. (c) Packing with layers viewed edge-on and counterions omitted for clarity. (d) Nodal representation of the network (yellow, uranium nodes; blue, deadc<sup>2-</sup> edges; view down [001] with [010] horizontal).

The wavelengths of the four main maxima for the two series are at 480, 499, 523 and 543 nm for the most blueshifted one, and 489, 511, 532 and 556 nm for the other, major one. The first series matches exactly that for complex 7 and thus may reasonably be attributed to the tris(carboxylate)-coordinated uranyl centre (U1). The other



**Fig. 8** (a) View of compound **8** with displacement ellipsoids shown at the 50% probability level. The water molecule and the hydrogen atoms are omitted. Symmetry codes: i = 1/2 - x, 1/2 - y, 1 - z; j = 1 - x, 1 - y, 1 - z; k = -x, y, 3/2 - z. (b) and (c) Two views of the uranyl-based monoperiodic subunit. (d) View of the triperiodic framework with uranium coordination polyhedra yellow and silver atoms shown as blue spheres.

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**Fig. 9** Intermolecular associations of the deadc<sup>2-</sup> ligand. (a) Loose association with acetonitrile in **1** (only one position of the disordered solvent is shown). (b) NH···O (dashed lines) and CH··· $\pi$  (dotted lines) interactions with H<sub>2</sub>NMe<sub>2</sub><sup>+</sup> in **3**. (c) Dimerization through one parallel-displaced  $\pi$ -stacking interaction (dashed line) and two edge-to-face interactions (dotted lines) in **2**, **5**, **6** and **8**.

component, which is red-shifted and with maxima positions in the highest range measured for complexes with O<sub>6</sub> equatorial environments,<sup>33</sup> should thus be attributed to the uranyl cation chelated by two carboxylate and one nitrate groups (U2). It may be noted that the maxima in the spectrum of uranyl nitrate hexahydrate measured under the same conditions are at 486, 508, 532 and 557 nm,<sup>34</sup> i.e. very close to those of the second component here. Calculation of bond valence parameters<sup>35</sup> with PLATON<sup>25</sup> gives axial/equatorial values of 3.47/2.589 for U1 and 3.446/2.683 for U2. If these values are taken as an indication of bond strength, the donor strength in the equatorial plane is slightly larger for U2, which induces a decrease of the oxo bond order to uranyl,<sup>36</sup> as reflected in the maxima positions. Finally, the spectrum of complex 3, which is reproducible when measured on different samples, displays only a very broad emission peak centered at about 535 nm (Fig. S2, ESI<sup>†</sup>). Although unusual and of uncertain origin, similar spectra have previously been observed in other uranyl ion complexes with carboxylates.37



Fig. 10 Emission spectra of complexes 1, 2, 4, 5, 7 and 8 in the crystalline state upon excitation at 420 nm.



Fig. 11 Deconvolution of the emission spectrum of 4 (red) showing the two series of peaks (dashed blue and orange).

## Conclusions

We have reported the synthesis, crystal structure and emission spectrum of eight uranyl ion complexes with deadc<sup>2–</sup>, a bulky aromatic dicarboxylate ligand. Depending on the organic cosolvent and the additional species present, the periodicity of the complexes varies from 0 to 3, the extreme values being obtained for a  $\mu_3$ -oxo-bridged tetranuclear complex and a framework in which additional bridging is provided by silver(1) cations, respectively, while the diperiodic networks present the **fes**, **sql** or **hcb** topologies. All but two complexes have emission spectra in the solid state with maxima positions matching those usual for O<sub>5</sub> or O<sub>6</sub> equatorial coordination numbers, with a mixture of two distinct components in one case.

The deadc<sup>2-</sup> ligand has the attractive feature of adopting a fairly predictable conformation, though the capacity of the carboxylate groups to adopt various coordination modes still adds variety to the structure of their complexes. The capacity of the acid itself to form inclusion complexes through aggregation defining cavities, seems to be partly retained in the anion complexes but the formation of cage-like oligomeric uranyl ion complexes, anticipated on the basis of similarity of the anions to those derived from trans-1,2cyclohexanedicarboxylic acid, has not yet been observed. The chirality of the anions appears to be an important influence upon the structures of their complexes, the fact that in all cases so far known, use of the racemic acid always results in structures which are also racemic meaning that the use of isolated enantiomers necessarily results in quite different species.

Segregation of the hydrophobic dibenzobarrelene units on the outside of the coordination polymers formed, together with the existence of weak  $\pi$ -stacking or CH $\cdots\pi$  interactions involving these units, either with one another or with counterions, are common features of these complexes and they play a prominent role in defining the packing of the polymeric moieties. Due to their ability to be bonded to both carboxylates and aromatic rings, silver(1) cations are able to bridge the hydrophilic and hydrophobic parts of different polymeric units and thus increase the overall periodicity, yet another example of the versatility of this cation and its interest in the synthesis of uranyl-based coordination polymers.<sup>38</sup>

In conclusion, these results show that, although deadc<sup>2-</sup> is fairly rigid and has a relative positioning of the coordination sites closely similar to that in trans-1,2cyclohexanedicarboxylate, it has failed up to now to produce cage-like complexes with uranyl ion, a disappointing fact because this was one of the anticipated properties of these ligands. Its amphiphilic nature can lead to enhanced periodicity through the addition of particular metal ions, but here its reduced denticity when compared with the amphiphilic Kemp's tricarboxylate<sup>30</sup> prevents formation of closed species such as nanotubes or cages.

# Conflicts of interest

There are no conflicts of interest to declare.

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