

A one-dimensional zigzag coordination polymer: *catena*-poly[[[triaqua-cadmium(II)]- $[\mu$ -2,2'-(5-methyl-1,3-phenylenedioxy)diacetato- $\kappa^4 O, O': O'', O''']$] monohydrate]

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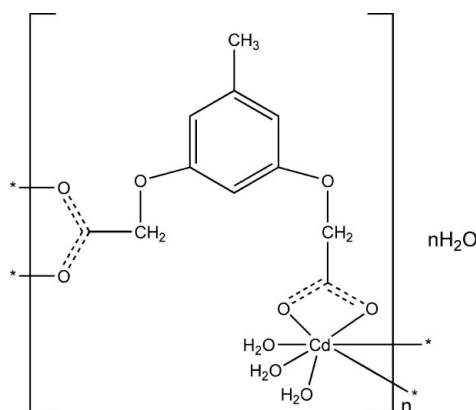
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 Key indicators: single-crystal X-ray study; $T = 298$ K; mean $\sigma(\text{C}-\text{C}) = 0.007$ Å; R factor = 0.040; wR factor = 0.093; data-to-parameter ratio = 13.2.

In the title one-dimensional coordination polymer, $\{[\text{Cd}(\text{C}_{11}\text{H}_{10}\text{O}_6)(\text{H}_2\text{O})_3] \cdot \text{H}_2\text{O}\}_n$, the 2,2'-(5-methyl-1,3-phenylenedioxy)diacetate dianions connect Cd^{II} ions in a head-to-tail fashion to generate zigzag chains. The coordination geometry of the Cd atom is distorted pentagonal bipyramidal. There are $\text{O}-\text{H} \cdots \text{O}$ hydrogen bonds between the carboxyl O atoms, the aqua ligands and the uncoordinated water molecules.

Related literature

For related literature on coordination polymers, see: Burrows *et al.* (2004); Hong *et al.* (2006); Janiak (2000, 2003); Kitagawa *et al.* (2004); Moulton & Zaworotko (2001); Russell *et al.* (2001).



Experimental

Crystal data

$[\text{Cd}(\text{C}_{11}\text{H}_{10}\text{O}_6)(\text{H}_2\text{O})_3] \cdot \text{H}_2\text{O}$
 $M_r = 422.66$
 Triclinic, $P\bar{1}$
 $a = 7.3792$ (11) Å
 $b = 8.6946$ (12) Å
 $c = 11.9495$ (17) Å
 $\alpha = 85.294$ (19)°
 $\beta = 82.52$ (2)°

$\gamma = 88.44$ (2)°
 $V = 757.47$ (19) Å³
 $Z = 2$
 Mo $K\alpha$ radiation
 $\mu = 1.49$ mm⁻¹
 $T = 298$ (2) K
 $0.24 \times 0.08 \times 0.02$ mm

Data collection

Bruker SMART CCD area-detector diffractometer
 Absorption correction: multi-scan (*SADABS*; Sheldrick, 1996)
 $T_{\text{min}} = 0.717$, $T_{\text{max}} = 0.971$

5397 measured reflections
 2647 independent reflections
 2121 reflections with $I > 2\sigma(I)$
 $R_{\text{int}} = 0.037$

Refinement

$R[F^2 > 2\sigma(F^2)] = 0.040$
 $wR(F^2) = 0.093$
 $S = 0.98$
 2647 reflections

200 parameters
 H-atom parameters constrained
 $\Delta\rho_{\text{max}} = 0.41$ e Å⁻³
 $\Delta\rho_{\text{min}} = -0.75$ e Å⁻³

Table 1

Selected geometric parameters (Å, °).

Cd—O9	2.262 (4)	Cd—O8	2.381 (4)
Cd—O7	2.297 (4)	Cd—O5 ⁱ	2.432 (4)
Cd—O1	2.303 (3)	Cd—O2	2.573 (4)
Cd—O6 ⁱ	2.331 (4)		
O9—Cd—O7	167.09 (14)	O6 ⁱ —Cd—O8	80.48 (13)
O9—Cd—O1	85.66 (14)	O9—Cd—O5 ⁱ	88.92 (13)
O9—Cd—O8	92.24 (14)	O9—Cd—O2	83.86 (14)
O7—Cd—O8	88.31 (14)	O8—Cd—O2	77.69 (12)
O1—Cd—O8	130.96 (13)	O5 ⁱ —Cd—O2	146.90 (12)

 Symmetry code: (i) $x - 1, y, z + 1$.

Table 2

Hydrogen-bond geometry (Å, °).

$D-\text{H} \cdots A$	$D-\text{H}$	$\text{H} \cdots A$	$D \cdots A$	$D-\text{H} \cdots A$
O7—H7B ⁱⁱ ···O8 ⁱⁱ	0.85	2.08	2.875 (5)	156
O7—H7C ⁱⁱⁱ ···O5 ⁱⁱⁱ	0.85	2.00	2.846 (5)	174
O8—H8C ^{iv} ···O9 ^{iv}	0.85	2.51	3.263 (5)	148
O8—H8C ^v ···O10 ^v	0.85	2.38	3.008 (7)	131
O8—H8D ^{vi} ···O2 ^{vi}	0.85	2.14	2.808 (5)	136
O9—H9E ^{vii} ···O1 ^{vii}	0.85	1.92	2.752 (5)	165
O9—H9F ^{vii} ···O10 ^{vii}	0.85	2.10	2.659 (6)	123
O10—H10C ^{vii} ···O6	0.85	2.08	2.731 (6)	133
O10—H10D ^{vii} ···O2 ^{vii}	0.85	1.95	2.793 (6)	170

 Symmetry codes: (ii) $-x - 1, -y, -z + 2$; (iii) $-x, -y + 1, -z + 1$; (iv) $-x, -y, -z + 2$; (v) $-x, -y, -z + 1$; (vi) $-x, -y + 1, -z + 2$; (vii) $x, y, z + 1$.

Data collection: *SMART* (Bruker, 2000); cell refinement: *SAINTE* (Bruker, 2000); data reduction: *SAINTE*; program(s) used to solve structure: *SHELXS97* (Sheldrick, 1997); program(s) used to refine structure: *SHELXL97* (Sheldrick, 1997); molecular graphics: *SHELXTL* (Sheldrick, 2000); software used to prepare material for publication: *SHELXTL*.

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Supplementary data and figures for this paper are available from the IUCr electronic archives (Reference: OB2104).

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A one-dimensional zigzag coordination polymer: catena-poly[[[triaquacadmium(II)]- $[\mu$ -2,2'-(5-methyl-1,3-phenylenedioxy)diacetato- κ^4 O,O':O'',O''']] monohydrate]

Y. Lin, Y. Wei, X.-W. Wei and H.-T. Zhang

Comment

Supramolecular self-assembly of coordination polymers have been one of the areas of rapid growth in chemistry recently, owing to their intriguing molecular topologies and useful properties, such as molecular recognition, electronic, optical, magnetic and catalytic properties (Janiak, 2003; Kitagawa *et al.*, 2004). Generally, the structure of such a molecular architecture is governed by coordination interaction and other non-covalent interactions such as hydrogen bonding and π - π stacking as well as the conformations of ligands depending on their rigidity and flexibility (Russell *et al.*, 2001; Moulton & Zaworotko, 2001; Burrows *et al.*, 2004). To date, there is a great research interest focused on the coordination interaction, hydrogen bonding and π - π stacking as well as the rigidity of ligands, whereas there is scant attention to the influence of the flexibility of ligands on the structure of coordination polymer. In order to further understand the role of the flexibility of ligands in the self-assembly of coordination polymers, we have designed and synthesized a ligand bearing the flexible group, 2,2'-(5-methyl-1,3-phenylenedioxy)diacetic acid (abbreviated to H₂5-mpdoa), and employed it with Cd^{II} ion to assemble the title coordination polymer, (I), [Cd(5-mpdoa)(H₂O)₃]_n·nH₂O.

As shown in Fig. 1, the coordination geometry of the Cd^{II} atom is a distorted pentagonal bipyramid. The equatorial positions are occupied by four carboxyl O atoms (O1, O2, O5ⁱ, O6ⁱ, symmetry code (i) $x - 1, y, z + 1$) from two symmetry related 5-mpdoa ligands and the aqua O8 atom. The aqua O7 and O9 atoms are located at the axial vertices of the pentagonal bipyramid. Both two carboxylate groups of the 5-mpdoa ligands, O1/C1/O2 and O5ⁱ/C11ⁱ/O6ⁱ, chelate the Cd^{II} atom in the same mode. The O5/C11/O6 carboxylate is almost coplanar with the benzene ring C3—C8, whereas the O1/C1/O2 carboxylate has a slightly twisted from the benzene ring with the dihedral angle of 10.7 (7)°. Due to the flexibility of the molecule induced by the σ -rotation of the C—O bond (selected torsion angles are listed in Table 1), the 5-mpdoa ligand adopts a W-shape conformation.

The 5-mpdoa ligands connect the neighbouring Cd^{II} atoms in a head-to-tail mode to construct an infinite zigzag chain which runs along the [101] direction. Such a supramolecular geometry could be regarded as a result of the cooperation of coordination interaction, the symmetry and the flexibility of ligand molecule as well as the hydrogen bonds. All zigzag chains are packing together through an amount of hydrogen bonding interactions between the carboxyl O atoms, the aqua ligands and the lattice water molecules (Fig. 2, Table 2). The shortest center-center distance between two adjacent benzene rings of the different chains is 4.997 (15) Å, indicating no interchain π - π interaction of 5-mpdoa (Janiak *et al.*, 2000).

Experimental

The H₂5-mpdoa ligand, 2,2'-(5-methyl-1,3-phenylenedioxy)diacetic acid, was synthesized from 5-methylbenzene-1,3-diol and 2-chloroacetic acid according to a literature method reported by Hong *et al.* (2006). Cd(CH₃COO)₂·2(H₂O) (26.8 mg, 0.10 mmol) and H₂5-mpdoa (24.0 mg, 0.10 mmol) were dissolved in 10 ml water. The resulting yellow solution was filtered

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and the filtrate was left at room temperature. Yellow column-like crystals were obtained (25.5 mg, yield *ca* 60%) after several weeks by slow evaporation of the solvent.

Refinement

All non-hydrogen atoms were refined anisotropically. H atoms bonded to C atoms were introduced at calculated positions and refined using a riding model with C—H distances of 0.93–0.97 Å. All hydrogen atoms of the water molecules were located in difference maps at an intermediate stage of the refinement and were then treated as riding, with O—H=0.85 (3) Å. In all cases, the H-atom $U_{\text{iso}}(\text{H})$ is 1.2 times U_{eq} of the parent atom.

Figures

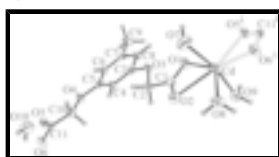


Fig. 1. A drawing of the asymmetric unit of (I) (solid line portion) with displacement ellipsoids at the 30% probability level. [symmetry code: (i) $x - 1, y, z + 1$].

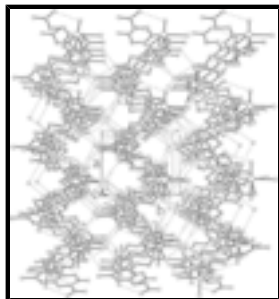


Fig. 2. A packing diagram of (I) viewed down the c axis. Dotted lines show O—H...O hydrogen bonds. All hydrogen atoms have been omitted for clarity.

catena-poly[[[triaquacadmium(II)]- $[\mu$ -2,2'-(5-methyl-1,3-phenylenedioxy)diacetato- κ^4 O,O':O'',O''']] mono-hydrate]

Crystal data

$[\text{Cd}(\text{C}_{11}\text{H}_{10}\text{O}_6)(\text{H}_2\text{O})_3] \cdot \text{H}_2\text{O}$

$M_r = 422.66$

Triclinic, $P\bar{1}$

Hall symbol: $-P\ 1$

$a = 7.3792$ (11) Å

$b = 8.6946$ (12) Å

$c = 11.9495$ (17) Å

$\alpha = 85.294$ (19)°

$\beta = 82.52$ (2)°

$\gamma = 88.44$ (2)°

$V = 757.47$ (19) Å³

$Z = 2$

$F_{000} = 424$

$D_x = 1.853$ Mg m⁻³

Mo $K\alpha$ radiation

$\lambda = 0.71073$ Å

Cell parameters from 1908 reflections

$\theta = 2.4$ – 25.5 °

$\mu = 1.49$ mm⁻¹

$T = 298$ (2) K

Column, yellow

$0.24 \times 0.08 \times 0.02$ mm

Data collection

Bruker SMART CCD area-detector diffractometer	2647 independent reflections
Radiation source: fine-focus sealed tube	2121 reflections with $I > 2\sigma(I)$
Monochromator: graphite	$R_{\text{int}} = 0.037$
$T = 274(2)$ K	$\theta_{\text{max}} = 25.0^\circ$
φ and ω scans	$\theta_{\text{min}} = 1.7^\circ$
Absorption correction: multi-scan (SADABS; Sheldrick, 1996)	$h = -8 \rightarrow 8$
$T_{\text{min}} = 0.717$, $T_{\text{max}} = 0.971$	$k = -10 \rightarrow 10$
5397 measured reflections	$l = -14 \rightarrow 14$

Refinement

Refinement on F^2	Secondary atom site location: difference Fourier map
Least-squares matrix: full	Hydrogen site location: inferred from neighbouring sites
$R[F^2 > 2\sigma(F^2)] = 0.040$	H-atom parameters constrained
$wR(F^2) = 0.093$	$w = 1/[\sigma^2(F_o^2) + (0.0479P)^2]$
$S = 0.98$	where $P = (F_o^2 + 2F_c^2)/3$
2647 reflections	$(\Delta/\sigma)_{\text{max}} = 0.001$
200 parameters	$\Delta\rho_{\text{max}} = 0.41 \text{ e } \text{\AA}^{-3}$
Primary atom site location: structure-invariant direct methods	$\Delta\rho_{\text{min}} = -0.75 \text{ e } \text{\AA}^{-3}$
	Extinction correction: none

Special details

Geometry. All e.s.d.'s (except the e.s.d. in the dihedral angle between two l.s. planes) are estimated using the full covariance matrix. The cell e.s.d.'s are taken into account individually in the estimation of e.s.d.'s in distances, angles and torsion angles; correlations between e.s.d.'s in cell parameters are only used when they are defined by crystal symmetry. An approximate (isotropic) treatment of cell e.s.d.'s is used for estimating e.s.d.'s involving l.s. planes.

Refinement. Refinement of F^2 against ALL reflections. The weighted R -factor wR and goodness of fit S are based on F^2 , conventional R -factors R are based on F , with F set to zero for negative F^2 . The threshold expression of $F^2 > \sigma(F^2)$ is used only for calculating R -factors(gt) *etc.* and is not relevant to the choice of reflections for refinement. R -factors based on F^2 are statistically about twice as large as those based on F , and R -factors based on ALL data will be even larger.

Fractional atomic coordinates and isotropic or equivalent isotropic displacement parameters (\AA^2)

	x	y	z	$U_{\text{iso}}^*/U_{\text{eq}}$
C1	-0.0333 (7)	0.3276 (6)	0.8172 (4)	0.0364 (12)
C2	0.0773 (7)	0.3784 (6)	0.7048 (4)	0.0357 (12)
H2A	0.2052	0.3511	0.7070	0.043*
H2B	0.0346	0.3267	0.6445	0.043*
C3	0.1556 (6)	0.6127 (6)	0.5904 (4)	0.0304 (11)

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C4	0.2445 (6)	0.5368 (5)	0.5016 (4)	0.0289 (11)
H4	0.2422	0.4298	0.5022	0.035*
C5	0.3370 (6)	0.6246 (6)	0.4120 (4)	0.0301 (11)
C6	0.3419 (6)	0.7843 (6)	0.4103 (4)	0.0339 (12)
H6	0.4059	0.8411	0.3491	0.041*
C7	0.2513 (7)	0.8586 (6)	0.4997 (5)	0.0350 (12)
C8	0.1596 (7)	0.7715 (6)	0.5901 (4)	0.0358 (12)
H8	0.0999	0.8201	0.6514	0.043*
C9	0.2558 (8)	1.0326 (6)	0.5005 (5)	0.0515 (15)
H9A	0.1448	1.0770	0.4773	0.061*
H9B	0.3583	1.0715	0.4492	0.061*
H9C	0.2671	1.0595	0.5756	0.061*
C10	0.4195 (7)	0.4039 (5)	0.3083 (4)	0.0348 (12)
H10A	0.2934	0.3767	0.3065	0.042*
H10B	0.4627	0.3468	0.3732	0.042*
C11	0.5344 (6)	0.3625 (7)	0.2014 (4)	0.0363 (13)
Cd	-0.24893 (5)	0.25748 (4)	1.02124 (3)	0.03576 (16)
O1	-0.1211 (5)	0.4250 (4)	0.8748 (3)	0.0411 (9)
O2	-0.0333 (5)	0.1860 (4)	0.8455 (3)	0.0499 (10)
O3	0.0573 (5)	0.5399 (4)	0.6840 (3)	0.0387 (9)
O4	0.4307 (5)	0.5637 (4)	0.3184 (3)	0.0383 (9)
O5	0.6157 (5)	0.4624 (4)	0.1334 (3)	0.0455 (9)
O6	0.5458 (5)	0.2198 (4)	0.1862 (3)	0.0500 (10)
O7	-0.4729 (5)	0.2338 (5)	0.9073 (3)	0.0635 (12)
H7B	-0.5520	0.1705	0.9401	0.076*
H7C	-0.5233	0.3214	0.8950	0.076*
O8	-0.2346 (5)	-0.0167 (4)	1.0480 (3)	0.0570 (11)
H8C	-0.1671	-0.0518	0.9923	0.069*
H8D	-0.1896	-0.0430	1.1088	0.069*
O9	0.0139 (5)	0.2816 (4)	1.0963 (3)	0.0580 (11)
H9E	0.0262	0.3754	1.1086	0.069*
H9F	0.0093	0.2255	1.1580	0.069*
O10	0.2207 (6)	0.0730 (5)	0.1980 (4)	0.0801 (15)
H10C	0.3344	0.0701	0.1748	0.096*
H10D	0.1737	-0.0127	0.1891	0.096*

Atomic displacement parameters (\AA^2)

	U^{11}	U^{22}	U^{33}	U^{12}	U^{13}	U^{23}
C1	0.028 (3)	0.047 (3)	0.034 (3)	-0.010 (2)	-0.003 (2)	-0.001 (3)
C2	0.040 (3)	0.035 (3)	0.031 (3)	-0.001 (2)	0.000 (2)	-0.002 (2)
C3	0.030 (3)	0.036 (3)	0.024 (3)	-0.004 (2)	-0.001 (2)	0.002 (2)
C4	0.029 (3)	0.027 (3)	0.031 (3)	-0.002 (2)	-0.003 (2)	-0.003 (2)
C5	0.027 (3)	0.039 (3)	0.025 (3)	-0.001 (2)	-0.002 (2)	-0.003 (2)
C6	0.030 (3)	0.037 (3)	0.031 (3)	-0.005 (2)	0.004 (2)	0.000 (2)
C7	0.034 (3)	0.030 (3)	0.043 (3)	-0.003 (2)	-0.010 (2)	-0.003 (2)
C8	0.043 (3)	0.032 (3)	0.032 (3)	0.008 (2)	-0.001 (2)	-0.008 (2)
C9	0.050 (4)	0.035 (3)	0.067 (4)	-0.002 (3)	0.003 (3)	-0.006 (3)

C10	0.034 (3)	0.036 (3)	0.032 (3)	-0.005 (2)	0.004 (2)	-0.001 (2)
C11	0.027 (3)	0.057 (4)	0.026 (3)	0.004 (3)	-0.002 (2)	-0.007 (3)
Cd	0.0367 (2)	0.0382 (2)	0.0297 (2)	-0.00158 (16)	0.00579 (15)	-0.00233 (16)
O1	0.043 (2)	0.046 (2)	0.030 (2)	-0.0051 (18)	0.0085 (17)	0.0003 (17)
O2	0.059 (3)	0.038 (2)	0.050 (3)	-0.0106 (19)	-0.003 (2)	0.0085 (19)
O3	0.047 (2)	0.0324 (19)	0.031 (2)	0.0022 (16)	0.0131 (16)	-0.0004 (16)
O4	0.051 (2)	0.031 (2)	0.029 (2)	-0.0055 (16)	0.0096 (17)	-0.0030 (16)
O5	0.045 (2)	0.053 (2)	0.033 (2)	0.0000 (19)	0.0115 (17)	-0.0003 (18)
O6	0.058 (3)	0.040 (2)	0.050 (3)	-0.0003 (19)	0.0103 (19)	-0.0144 (19)
O7	0.060 (3)	0.059 (3)	0.072 (3)	-0.022 (2)	-0.023 (2)	0.023 (2)
O8	0.047 (2)	0.046 (2)	0.072 (3)	0.0030 (19)	0.006 (2)	0.005 (2)
O9	0.070 (3)	0.048 (2)	0.062 (3)	-0.001 (2)	-0.023 (2)	-0.013 (2)
O10	0.078 (3)	0.047 (3)	0.124 (4)	-0.006 (2)	-0.044 (3)	-0.008 (3)

Geometric parameters (Å, °)

C1—O1	1.249 (6)	C10—O4	1.409 (5)
C1—O2	1.250 (6)	C10—C11	1.503 (7)
C1—C2	1.517 (7)	C10—H10A	0.9700
C1—Cd	2.766 (5)	C10—H10B	0.9700
C2—O3	1.413 (6)	C11—O5	1.249 (6)
C2—H2A	0.9700	C11—O6	1.268 (6)
C2—H2B	0.9700	C11—Cd ⁱ	2.715 (5)
C3—O3	1.370 (6)	Cd—O9	2.262 (4)
C3—C4	1.379 (7)	Cd—O7	2.297 (4)
C3—C8	1.382 (7)	Cd—O1	2.303 (3)
C4—C5	1.378 (7)	Cd—O6 ⁱⁱ	2.331 (4)
C4—H4	0.9300	Cd—O8	2.381 (4)
C5—O4	1.374 (5)	Cd—O5 ⁱⁱ	2.432 (4)
C5—C6	1.388 (7)	Cd—O2	2.573 (4)
C6—C7	1.382 (7)	O7—H7B	0.8489
C6—H6	0.9300	O7—H7C	0.8491
C7—C8	1.379 (7)	O8—H8C	0.8499
C7—C9	1.515 (7)	O8—H8D	0.8492
C8—H8	0.9300	O9—H9E	0.8495
C9—H9A	0.9600	O9—H9F	0.8482
C9—H9B	0.9600	O10—H10C	0.8492
C9—H9C	0.9600	O10—H10D	0.8499
O1—C1—O2	123.5 (5)	O7—Cd—O1	84.25 (13)
O1—C1—C2	120.1 (5)	O9—Cd—O6 ⁱⁱ	100.20 (15)
O2—C1—C2	116.4 (5)	O7—Cd—O6 ⁱⁱ	92.61 (15)
O1—C1—Cd	55.6 (3)	O1—Cd—O6 ⁱⁱ	148.09 (13)
O2—C1—Cd	68.0 (3)	O9—Cd—O8	92.24 (14)
C2—C1—Cd	175.5 (4)	O7—Cd—O8	88.31 (14)
O3—C2—C1	109.0 (4)	O1—Cd—O8	130.96 (13)
O3—C2—H2A	109.9	O6 ⁱⁱ —Cd—O8	80.48 (13)
C1—C2—H2A	109.9	O9—Cd—O5 ⁱⁱ	88.92 (13)

supplementary materials

O3—C2—H2B	109.9	O7—Cd—O5 ⁱⁱ	99.79 (15)
C1—C2—H2B	109.9	O1—Cd—O5 ⁱⁱ	93.99 (12)
H2A—C2—H2B	108.3	O6 ⁱⁱ —Cd—O5 ⁱⁱ	55.16 (12)
O3—C3—C4	123.9 (4)	O8—Cd—O5 ⁱⁱ	135.00 (13)
O3—C3—C8	114.8 (4)	O9—Cd—O2	83.86 (14)
C4—C3—C8	121.3 (5)	O7—Cd—O2	83.67 (14)
C3—C4—C5	117.8 (4)	O1—Cd—O2	53.34 (12)
C3—C4—H4	121.1	O6 ⁱⁱ —Cd—O2	157.94 (13)
C5—C4—H4	121.1	O8—Cd—O2	77.69 (12)
O4—C5—C4	123.7 (4)	O5 ⁱⁱ —Cd—O2	146.90 (12)
O4—C5—C6	114.6 (4)	O9—Cd—C11 ⁱⁱ	94.48 (14)
C4—C5—C6	121.7 (5)	O7—Cd—C11 ⁱⁱ	97.62 (15)
C5—C6—C7	119.8 (5)	O1—Cd—C11 ⁱⁱ	121.12 (15)
C5—C6—H6	120.1	O6 ⁱⁱ —Cd—C11 ⁱⁱ	27.78 (14)
C7—C6—H6	120.1	O8—Cd—C11 ⁱⁱ	107.90 (16)
C8—C7—C6	118.9 (4)	O5 ⁱⁱ —Cd—C11 ⁱⁱ	27.38 (14)
C8—C7—C9	120.1 (5)	O2—Cd—C11 ⁱⁱ	174.26 (15)
C6—C7—C9	120.9 (5)	O9—Cd—C1	84.11 (15)
C7—C8—C3	120.5 (5)	O7—Cd—C1	83.27 (14)
C7—C8—H8	119.7	O1—Cd—C1	26.58 (14)
C3—C8—H8	119.7	O6 ⁱⁱ —Cd—C1	173.43 (14)
C7—C9—H9A	109.5	O8—Cd—C1	104.43 (15)
C7—C9—H9B	109.5	O5 ⁱⁱ —Cd—C1	120.41 (15)
H9A—C9—H9B	109.5	O2—Cd—C1	26.77 (13)
C7—C9—H9C	109.5	C11 ⁱⁱ —Cd—C1	147.67 (18)
H9A—C9—H9C	109.5	C1—O1—Cd	97.9 (3)
H9B—C9—H9C	109.5	C1—O2—Cd	85.3 (3)
O4—C10—C11	109.4 (4)	C3—O3—C2	119.1 (4)
O4—C10—H10A	109.8	C5—O4—C10	118.4 (4)
C11—C10—H10A	109.8	C11—O5—Cd ⁱ	89.0 (3)
O4—C10—H10B	109.8	C11—O6—Cd ⁱ	93.2 (3)
C11—C10—H10B	109.8	Cd—O7—H7B	109.4
H10A—C10—H10B	108.3	Cd—O7—H7C	109.2
O5—C11—O6	122.5 (5)	H7B—O7—H7C	109.7
O5—C11—C10	121.9 (5)	Cd—O8—H8C	109.3
O6—C11—C10	115.6 (5)	Cd—O8—H8D	109.2
O5—C11—Cd ⁱ	63.6 (3)	H8C—O8—H8D	109.6
O6—C11—Cd ⁱ	59.0 (3)	Cd—O9—H9E	109.2
C10—C11—Cd ⁱ	173.6 (4)	Cd—O9—H9F	109.1
O9—Cd—O7	167.09 (14)	H9E—O9—H9F	109.7
O9—Cd—O1	85.66 (14)	H10C—O10—H10D	109.6
O1—C1—C2—O3	-0.6 (7)	O2—C1—O1—Cd	0.1 (6)
O2—C1—C2—O3	177.9 (4)	C2—C1—O1—Cd	178.5 (4)
O3—C3—C4—C5	179.3 (4)	O9—Cd—O1—C1	85.5 (3)

C8—C3—C4—C5	-0.3 (7)	O7—Cd—O1—C1	-86.4 (3)
C3—C4—C5—O4	-179.9 (4)	O6 ⁱⁱ —Cd—O1—C1	-172.1 (3)
C3—C4—C5—C6	0.1 (7)	O8—Cd—O1—C1	-3.7 (4)
O4—C5—C6—C7	179.5 (4)	O5 ⁱⁱ —Cd—O1—C1	174.1 (3)
C4—C5—C6—C7	-0.5 (7)	O2—Cd—O1—C1	-0.1 (3)
C5—C6—C7—C8	0.9 (7)	C11 ⁱⁱ —Cd—O1—C1	178.1 (3)
C5—C6—C7—C9	179.4 (5)	O1—C1—O2—Cd	-0.1 (5)
C6—C7—C8—C3	-1.1 (7)	C2—C1—O2—Cd	-178.5 (4)
C9—C7—C8—C3	-179.6 (5)	O9—Cd—O2—C1	-89.1 (3)
O3—C3—C8—C7	-178.9 (4)	O7—Cd—O2—C1	87.6 (3)
C4—C3—C8—C7	0.8 (7)	O1—Cd—O2—C1	0.1 (3)
O4—C10—C11—O5	-3.2 (7)	O6 ⁱⁱ —Cd—O2—C1	168.9 (3)
O4—C10—C11—O6	175.4 (4)	O8—Cd—O2—C1	177.3 (3)
O1—C1—Cd—O9	-92.1 (3)	O5 ⁱⁱ —Cd—O2—C1	-10.6 (4)
O2—C1—Cd—O9	88.0 (3)	C4—C3—O3—C2	14.2 (7)
O1—C1—Cd—O7	90.6 (3)	C8—C3—O3—C2	-166.1 (4)
O2—C1—Cd—O7	-89.3 (3)	C1—C2—O3—C3	175.6 (4)
O2—C1—Cd—O1	-179.9 (5)	C4—C5—O4—C10	5.6 (7)
O1—C1—Cd—O8	177.1 (3)	C6—C5—O4—C10	-174.4 (4)
O2—C1—Cd—O8	-2.8 (3)	C11—C10—O4—C5	-179.0 (4)
O1—C1—Cd—O5 ⁱⁱ	-6.8 (4)	O6—C11—O5—Cd ⁱ	-2.4 (5)
O2—C1—Cd—O5 ⁱⁱ	173.3 (3)	C10—C11—O5—Cd ⁱ	176.1 (4)
O1—C1—Cd—O2	179.9 (5)	O5—C11—O6—Cd ⁱ	2.5 (5)
O1—C1—Cd—C11 ⁱⁱ	-3.0 (5)	C10—C11—O6—Cd ⁱ	-176.1 (4)
O2—C1—Cd—C11 ⁱⁱ	177.1 (3)		

Symmetry codes: (i) $x+1, y, z-1$; (ii) $x-1, y, z+1$.

Hydrogen-bond geometry (Å, °)

<i>D</i> —H... <i>A</i>	<i>D</i> —H	H... <i>A</i>	<i>D</i> ... <i>A</i>	<i>D</i> —H... <i>A</i>
O7—H7B...O8 ⁱⁱⁱ	0.85	2.08	2.875 (5)	156
O7—H7C...O5 ^{iv}	0.85	2.00	2.846 (5)	174
O8—H8C...O9 ^v	0.85	2.51	3.263 (5)	148
O8—H8C...O10 ^{vi}	0.85	2.38	3.008 (7)	131
O8—H8D...O2 ^v	0.85	2.14	2.808 (5)	136
O9—H9E...O1 ^{vii}	0.85	1.92	2.752 (5)	165
O9—H9F...O10 ^{viii}	0.85	2.10	2.659 (6)	123
O10—H10C...O6	0.85	2.08	2.731 (6)	133
O10—H10D...O2 ^{vi}	0.85	1.95	2.793 (6)	170

Symmetry codes: (iii) $-x-1, -y, -z+2$; (iv) $-x, -y+1, -z+1$; (v) $-x, -y, -z+2$; (vi) $-x, -y, -z+1$; (vii) $-x, -y+1, -z+2$; (viii) $x, y, z+1$.

Fig. 1

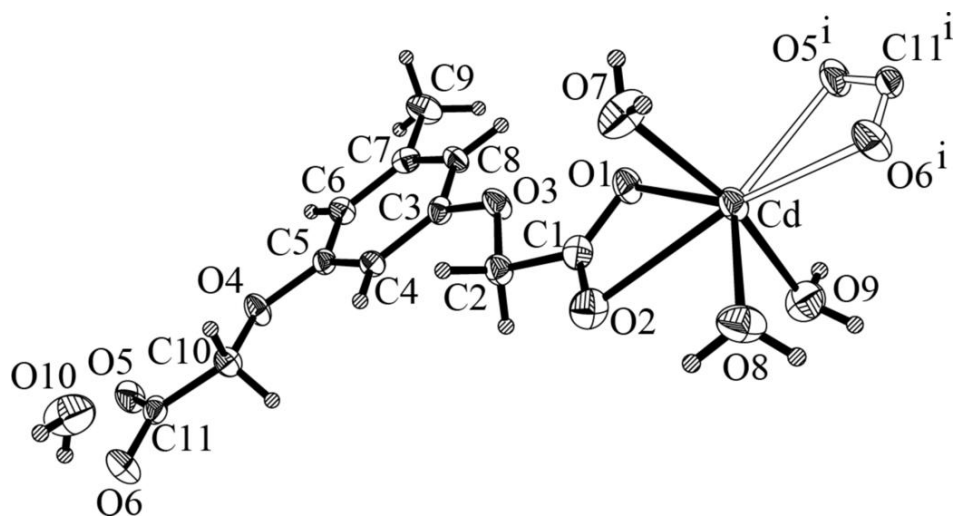


Fig. 2

