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Bis(μ -biphenyl-2,2'-dicarboxylato)bis-[aqua(4,4'-dimethyl-2,2'-bipyridine- κ^2N,N')copper(II)]

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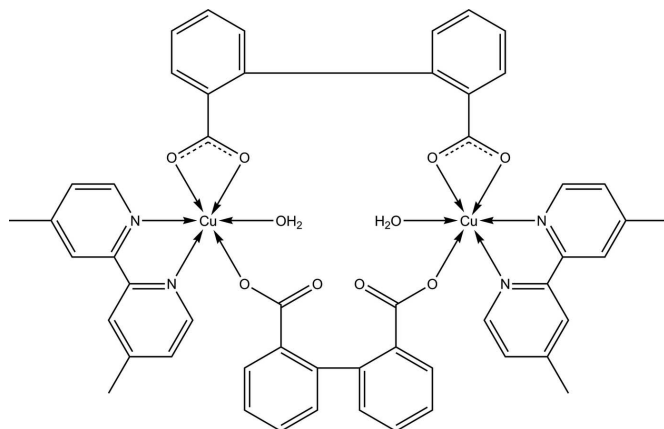
Received 27 September 2009; accepted 6 October 2009

Key indicators: single-crystal X-ray study; $T = 296$ K; mean $\sigma(\text{C}-\text{C}) = 0.007$ Å; R factor = 0.047; wR factor = 0.109; data-to-parameter ratio = 14.8.

The molecule of the title binuclear copper(II) complex, $[\text{Cu}_2(\text{C}_{14}\text{H}_8\text{O}_4)_2(\text{C}_{12}\text{H}_{12}\text{N}_2)_2(\text{H}_2\text{O})_2]$, is bisected by a crystallographic twofold axis. Each Cu^{II} atom is coordinated in a distorted octahedral geometry by three O atoms from two biphenyl-2,2'-dicarboxylate anions, one aqua O atom and two N atoms of a 4,4'-dimethyl-2,2'-bipyridine ligand. Intramolecular $\text{O}-\text{H}\cdots\text{O}$ hydrogen bonds between the coordinated water molecules and the carboxylate O atoms are also present.

Related literature

For related structures, see: Li *et al.* (2009); Jiang & Feng (2009); Xu *et al.* (2009); Zhang *et al.* (2009); Rizal & Ng (2009); Zhang (2009).



Experimental

Crystal data

$[\text{Cu}_2(\text{C}_{14}\text{H}_8\text{O}_4)_2(\text{C}_{12}\text{H}_{12}\text{N}_2)_2(\text{H}_2\text{O})_2]$ $V = 4664.2$ (13) Å³
 $M_r = 1109.71$ $Z = 4$
 Monoclinic, $C2/c$ Mo $K\alpha$ radiation
 $a = 17.104$ (3) Å $\mu = 0.98$ mm⁻¹
 $b = 15.395$ (2) Å $T = 296$ K
 $c = 18.289$ (3) Å $0.26 \times 0.24 \times 0.22$ mm
 $\beta = 104.413$ (3)°

Data collection

Bruker SMART APEXII CCD 13512 measured reflections
 area-detector diffractometer 4594 independent reflections
 Absorption correction: multi-scan 3108 reflections with $I > 2\sigma(I)$
 (SADABS; Bruker, 2005) $R_{\text{int}} = 0.060$
 $T_{\text{min}} = 0.785$, $T_{\text{max}} = 0.814$

Refinement

$R[F^2 > 2\sigma(F^2)] = 0.047$ 12 restraints
 $wR(F^2) = 0.109$ H-atom parameters constrained
 $S = 0.93$ $\Delta\rho_{\text{max}} = 0.87$ e Å⁻³
 4594 reflections $\Delta\rho_{\text{min}} = -0.33$ e Å⁻³
 310 parameters

Table 1

Hydrogen-bond geometry (Å, °).

$D-H\cdots A$	$D-H$	$H\cdots A$	$D\cdots A$	$D-H\cdots A$
$\text{O1W}-\text{H1WB}\cdots\text{O4}^i$	0.85	1.78	2.632 (4)	174
$\text{O1W}-\text{H1WA}\cdots\text{O1}^i$	0.85	1.95	2.782 (4)	164

Symmetry code: (i) $-x, y, -z + \frac{1}{2}$.

Data collection: APEX2 (Bruker, 2005); cell refinement: SAINT (Bruker, 2005); data reduction: SAINT; program(s) used to solve structure: SHELXS97 (Sheldrick, 2008); program(s) used to refine structure: SHELXL97 (Sheldrick, 2008); molecular graphics: SHELXTL (Sheldrick, 2008); software used to prepare material for publication: SHELXTL.

Supplementary data and figures for this paper are available from the IUCr electronic archives (Reference: JH2107).

References

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supplementary materials

Acta Cryst. (2009). E65, m1360 [doi:10.1107/S1600536809040628]

Bis(μ -biphenyl-2,2'-dicarboxylato)bis[aqua(4,4'-dimethyl-2,2'-bipyridine- κ^2N,N')copper(II)]

X.-Y. Dong, X.-J. Xu and L. Yang

Comment

The title binuclear copper(II) complex, $[\text{Cu}_2(\text{C}_{14}\text{H}_8\text{O}_4)_2(\text{C}_{10}\text{H}_8\text{N}_2)_2(\text{H}_2\text{O})_2]$, is a centrosymmetric dimer. The asymmetric unit consists of one Cu^{II} atom, one 4,4'-dimethyl-2,2'-bipyridine (dbpy) ligand, one [1,1'-biphenyl]-2,2'-dicarboxylate dianion (bpdc²⁻) and a coordinated water molecule.

The Cu^{II} atom is six-coordinated by two N atoms from bpy and four O atoms, three from two bpdc²⁻ anions and one from coordinated H_2O , in a distorted octahedron coordination geometry. And it is noteworthy that the two Cu^{II} ions in the complex are bridged by two bpdc²⁻ dianions, one is in a bis-monodentate mode whereas the other is in a bis-bidentate mode.

Experimental

The title compound was synthesized hydrothermally in a Teflon-lined autoclave (25 ml) by heating a mixture of H_2bpdc (0.2 mmol), dbpy (0.4 mmol) and $\text{CuSO}_4 \cdot 5\text{H}_2\text{O}$ (0.2 mmol) in water (10 ml) at 393 K for 3 d. Crystals suitable for X-ray analysis were obtained.

Refinement

All H atoms were included in calculated positions, with C—H bond lengths fixed at 0.96 Å (methyl CH_3), 0.93 Å (aryl group) and O—H = 0.85 Å and were refined in the riding-model approximation. $U_{\text{iso}}(\text{H})$ values were calculated at 1.5 $U_{\text{eq}}(\text{C})$ for methyl groups and 1.2 $U_{\text{eq}}(\text{C})$ otherwise.

Figures

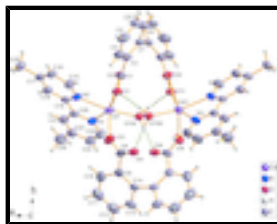


Fig. 1. The molecular structure of the title compound with the atom-labelling scheme. Displacement ellipsoids are drawn at the 30% probability level. H atoms are presented as small spheres of arbitrary radius. Hydrogen-bond interactions are drawn with dashed lines. Atoms labeled(i) are generated by the symmetry code $-x, y, -z + 1/2$.

Bis(μ -biphenyl-2,2'-dicarboxylato)bis[aqua(4,4'-dimethyl-2,2'- bipyridine)copper(II)]

Crystal data

$[\text{Cu}_2(\text{C}_{14}\text{H}_8\text{O}_4)_2(\text{C}_{12}\text{H}_{12}\text{N}_2)_2(\text{H}_2\text{O})_2]$

$F_{000} = 2240$

supplementary materials

$M_r = 1109.71$

Monoclinic, $C2/c$

Hall symbol: $-C\ 2yc$

$a = 17.104\ (3)\ \text{\AA}$

$b = 15.395\ (2)\ \text{\AA}$

$c = 18.289\ (3)\ \text{\AA}$

$\beta = 104.413\ (3)^\circ$

$V = 4664.2\ (13)\ \text{\AA}^3$

$Z = 4$

$D_x = 1.580\ \text{Mg m}^{-3}$

Mo $K\alpha$ radiation, $\lambda = 0.71073\ \text{\AA}$

Cell parameters from 3688 reflections

$\theta = 2.7\text{--}27.3^\circ$

$\mu = 0.98\ \text{mm}^{-1}$

$T = 296\ \text{K}$

Block, colourless

$0.26 \times 0.24 \times 0.22\ \text{mm}$

Data collection

Bruker SMART APEXII CCD area-detector diffractometer

Radiation source: fine-focus sealed tube

Monochromator: graphite

$T = 296\ \text{K}$

φ and ω scans

Absorption correction: multi-scan (SADABS; Bruker, 2005)

$T_{\min} = 0.785$, $T_{\max} = 0.814$

13512 measured reflections

4594 independent reflections

3108 reflections with $I > 2\sigma(I)$

$R_{\text{int}} = 0.060$

$\theta_{\max} = 26.0^\circ$

$\theta_{\min} = 1.8^\circ$

$h = -21 \rightarrow 20$

$k = -18 \rightarrow 12$

$l = -22 \rightarrow 22$

Refinement

Refinement on F^2

Least-squares matrix: full

$R[F^2 > 2\sigma(F^2)] = 0.047$

$wR(F^2) = 0.109$

$S = 0.93$

4594 reflections

310 parameters

12 restraints

Primary atom site location: structure-invariant direct methods

Secondary atom site location: difference Fourier map

Hydrogen site location: inferred from neighbouring sites

H-atom parameters constrained

$w = 1/[\sigma^2(F_o^2) + (0.0562P)^2]$

where $P = (F_o^2 + 2F_c^2)/3$

$(\Delta/\sigma)_{\max} = 0.001$

$\Delta\rho_{\max} = 0.87\ \text{e \AA}^{-3}$

$\Delta\rho_{\min} = -0.33\ \text{e \AA}^{-3}$

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)

	<i>x</i>	<i>y</i>	<i>z</i>	$U_{\text{iso}}^*/U_{\text{eq}}$
Cd1	0.13088 (2)	0.599652 (19)	0.344239 (17)	0.05361 (14)
N1	0.1810 (2)	0.5542 (2)	0.4686 (2)	0.0615 (10)
N2	0.2592 (2)	0.6574 (2)	0.39215 (19)	0.0516 (9)
O1	0.11652 (18)	0.67905 (18)	0.23068 (16)	0.0602 (8)
O2	0.0871 (2)	0.74927 (19)	0.32546 (17)	0.0611 (8)
O3	0.1387 (2)	0.4729 (2)	0.2927 (3)	0.0977 (12)
O4	0.0538 (2)	0.41826 (19)	0.1917 (2)	0.0701 (10)
O1W	0.00362 (18)	0.57274 (19)	0.35468 (17)	0.0611 (8)
H1WA	-0.0326	0.6100	0.3365	0.073*
H1WB	-0.0137	0.5234	0.3371	0.073*
C1	0.0942 (3)	0.7474 (3)	0.2590 (3)	0.0527 (11)
C2	0.0782 (3)	0.8273 (3)	0.2111 (2)	0.0537 (11)
C3	0.1193 (3)	0.8370 (3)	0.1546 (3)	0.0677 (14)
H3	0.1545	0.7937	0.1473	0.081*
C4	0.1087 (4)	0.9095 (3)	0.1096 (3)	0.0825 (17)
H4	0.1387	0.9161	0.0740	0.099*
C5	0.0541 (4)	0.9725 (3)	0.1166 (3)	0.0813 (16)
H5	0.0455	1.0205	0.0848	0.098*
C6	0.0121 (3)	0.9632 (3)	0.1716 (3)	0.0685 (14)
H6	-0.0245	1.0060	0.1769	0.082*
C7	0.0232 (3)	0.8913 (2)	0.2195 (3)	0.0540 (11)
C8	0.0959 (3)	0.4138 (3)	0.2577 (4)	0.0675 (15)
C9	0.0975 (3)	0.3293 (3)	0.3005 (3)	0.0578 (12)
C10	0.1638 (4)	0.3135 (4)	0.3615 (4)	0.0935 (19)
H10	0.2047	0.3546	0.3747	0.112*
C11	0.1687 (5)	0.2361 (5)	0.4026 (4)	0.114 (2)
H11	0.2144	0.2245	0.4411	0.137*
C12	0.1072 (5)	0.1780 (4)	0.3865 (4)	0.101 (2)
H12	0.1096	0.1276	0.4151	0.121*
C13	0.0428 (3)	0.1938 (3)	0.3290 (3)	0.0691 (14)
H13	0.0007	0.1538	0.3194	0.083*
C14	0.0357 (3)	0.2671 (2)	0.2827 (2)	0.0491 (10)
C15	0.2960 (3)	0.7114 (3)	0.3534 (2)	0.0619 (12)
H15	0.2725	0.7195	0.3023	0.074*
C16	0.3658 (3)	0.7551 (3)	0.3848 (3)	0.0577 (12)
H16	0.3882	0.7925	0.3556	0.069*
C17	0.4026 (3)	0.7432 (3)	0.4598 (2)	0.0523 (11)
C18	0.3652 (3)	0.6872 (3)	0.4998 (2)	0.0514 (11)
H18	0.3882	0.6778	0.5508	0.062*
C19	0.2946 (3)	0.6456 (2)	0.4655 (2)	0.0446 (10)
C20	0.2522 (3)	0.5850 (2)	0.5076 (3)	0.0484 (11)
C21	0.2842 (3)	0.5611 (3)	0.5819 (3)	0.0575 (12)

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H21	0.3336	0.5840	0.6080	0.069*
C22	0.2441 (3)	0.5040 (3)	0.6179 (3)	0.0589 (12)
C23	0.1719 (3)	0.4713 (3)	0.5762 (3)	0.0750 (15)
H23	0.1435	0.4311	0.5974	0.090*
C24	0.1423 (3)	0.4981 (3)	0.5035 (3)	0.0761 (15)
H24	0.0927	0.4763	0.4766	0.091*
C25	0.4800 (3)	0.7895 (3)	0.4962 (3)	0.0717 (14)
H25A	0.4703	0.8509	0.4960	0.108*
H25B	0.5196	0.7774	0.4685	0.108*
H25C	0.4994	0.7698	0.5473	0.108*
C26	0.2776 (3)	0.4777 (4)	0.6985 (3)	0.0843 (16)
H26A	0.3136	0.4294	0.7006	0.126*
H26B	0.2342	0.4612	0.7203	0.126*
H26C	0.3066	0.5256	0.7263	0.126*

Atomic displacement parameters (\AA^2)

	U^{11}	U^{22}	U^{33}	U^{12}	U^{13}	U^{23}
Cd1	0.0639 (3)	0.0397 (2)	0.0639 (2)	-0.00834 (15)	0.02867 (18)	-0.00204 (16)
N1	0.068 (3)	0.050 (2)	0.073 (3)	-0.012 (2)	0.031 (2)	0.012 (2)
N2	0.062 (2)	0.044 (2)	0.057 (2)	-0.0098 (17)	0.0290 (18)	0.0006 (17)
O1	0.074 (2)	0.0420 (17)	0.073 (2)	0.0006 (15)	0.0356 (17)	-0.0078 (15)
O2	0.078 (2)	0.0476 (17)	0.068 (2)	0.0056 (15)	0.0378 (17)	0.0039 (15)
O3	0.087 (2)	0.070 (2)	0.150 (3)	-0.021 (2)	0.057 (2)	-0.040 (2)
O4	0.084 (3)	0.0377 (18)	0.106 (3)	-0.0016 (17)	0.058 (2)	-0.0019 (19)
O1W	0.065 (2)	0.0444 (16)	0.080 (2)	-0.0043 (15)	0.0306 (17)	-0.0008 (16)
C1	0.055 (3)	0.039 (2)	0.073 (3)	-0.0036 (19)	0.033 (2)	0.001 (2)
C2	0.063 (3)	0.042 (2)	0.065 (3)	-0.007 (2)	0.032 (2)	-0.002 (2)
C3	0.082 (4)	0.054 (3)	0.080 (3)	-0.004 (3)	0.045 (3)	0.003 (3)
C4	0.113 (5)	0.073 (4)	0.078 (4)	-0.014 (3)	0.054 (3)	0.009 (3)
C5	0.115 (5)	0.050 (3)	0.087 (4)	-0.005 (3)	0.040 (3)	0.017 (3)
C6	0.088 (4)	0.041 (3)	0.083 (3)	-0.002 (2)	0.035 (3)	0.008 (2)
C7	0.063 (3)	0.035 (2)	0.068 (3)	-0.005 (2)	0.025 (2)	0.000 (2)
C8	0.062 (3)	0.044 (3)	0.116 (5)	-0.009 (2)	0.057 (3)	-0.029 (3)
C9	0.058 (3)	0.053 (3)	0.069 (3)	0.005 (2)	0.027 (2)	-0.026 (2)
C10	0.067 (4)	0.093 (5)	0.117 (5)	0.003 (3)	0.017 (4)	-0.044 (4)
C11	0.116 (5)	0.116 (5)	0.093 (4)	0.045 (5)	-0.008 (4)	-0.020 (4)
C12	0.125 (6)	0.089 (5)	0.082 (4)	0.030 (4)	0.013 (4)	-0.001 (4)
C13	0.090 (4)	0.052 (3)	0.069 (3)	0.017 (3)	0.026 (3)	0.003 (3)
C14	0.061 (3)	0.035 (2)	0.058 (3)	0.0070 (19)	0.0269 (19)	-0.0053 (19)
C15	0.072 (3)	0.065 (3)	0.055 (3)	-0.009 (3)	0.029 (2)	0.012 (2)
C16	0.066 (3)	0.049 (3)	0.068 (3)	-0.010 (2)	0.037 (3)	0.008 (2)
C17	0.066 (3)	0.036 (2)	0.065 (3)	-0.003 (2)	0.036 (2)	-0.004 (2)
C18	0.062 (3)	0.047 (2)	0.052 (2)	-0.003 (2)	0.027 (2)	-0.003 (2)
C19	0.056 (3)	0.032 (2)	0.055 (3)	0.0018 (19)	0.031 (2)	-0.0006 (19)
C20	0.056 (3)	0.036 (2)	0.062 (3)	0.0020 (19)	0.032 (2)	0.001 (2)
C21	0.066 (3)	0.052 (3)	0.064 (3)	-0.001 (2)	0.034 (2)	0.008 (2)
C22	0.069 (3)	0.054 (3)	0.068 (3)	0.006 (2)	0.042 (3)	0.010 (2)

C23	0.083 (4)	0.068 (3)	0.087 (4)	-0.013 (3)	0.045 (3)	0.025 (3)
C24	0.068 (3)	0.070 (3)	0.096 (4)	-0.016 (3)	0.030 (3)	0.020 (3)
C25	0.077 (4)	0.068 (3)	0.077 (3)	-0.023 (3)	0.033 (3)	-0.007 (3)
C26	0.106 (4)	0.083 (4)	0.079 (4)	-0.003 (3)	0.050 (3)	0.022 (3)

Geometric parameters (Å, °)

Cd1—O3	2.186 (4)	C10—H10	0.9300
Cd1—O1W	2.270 (3)	C11—C12	1.357 (9)
Cd1—N2	2.327 (3)	C11—H11	0.9300
Cd1—N1	2.329 (4)	C12—C13	1.342 (8)
Cd1—O1	2.369 (3)	C12—H12	0.9300
Cd1—O2	2.420 (3)	C13—C14	1.397 (6)
N1—C20	1.335 (6)	C13—H13	0.9300
N1—C24	1.343 (5)	C14—C14 ⁱ	1.482 (8)
N2—C19	1.339 (5)	C15—C16	1.366 (6)
N2—C15	1.346 (5)	C15—H15	0.9300
O1—C1	1.272 (5)	C16—C17	1.371 (6)
O2—C1	1.252 (5)	C16—H16	0.9300
O3—C8	1.240 (6)	C17—C18	1.386 (5)
O4—C8	1.244 (6)	C17—C25	1.505 (6)
O1W—H1WA	0.8498	C18—C19	1.373 (6)
O1W—H1WB	0.8499	C18—H18	0.9300
C1—C2	1.495 (6)	C19—C20	1.506 (5)
C2—C3	1.395 (6)	C20—C21	1.382 (6)
C2—C7	1.397 (6)	C21—C22	1.380 (6)
C3—C4	1.372 (6)	C21—H21	0.9300
C3—H3	0.9300	C22—C23	1.375 (7)
C4—C5	1.374 (7)	C22—C26	1.499 (6)
C4—H4	0.9300	C23—C24	1.364 (7)
C5—C6	1.382 (7)	C23—H23	0.9300
C5—H5	0.9300	C24—H24	0.9300
C6—C7	1.394 (6)	C25—H25A	0.9600
C6—H6	0.9300	C25—H25B	0.9600
C7—C7 ⁱ	1.524 (8)	C25—H25C	0.9600
C8—C9	1.515 (7)	C26—H26A	0.9600
C9—C10	1.400 (8)	C26—H26B	0.9600
C9—C14	1.403 (6)	C26—H26C	0.9600
C10—C11	1.400 (9)		
O3—Cd1—O1W	92.06 (12)	C9—C10—H10	119.9
O3—Cd1—N2	110.50 (13)	C12—C11—C10	120.3 (6)
O1W—Cd1—N2	150.70 (11)	C12—C11—H11	119.8
O3—Cd1—N1	96.06 (15)	C10—C11—H11	119.8
O1W—Cd1—N1	89.48 (12)	C13—C12—C11	119.4 (6)
N2—Cd1—N1	70.32 (12)	C13—C12—H12	120.3
O3—Cd1—O1	95.00 (13)	C11—C12—H12	120.3
O1W—Cd1—O1	106.00 (11)	C12—C13—C14	123.5 (6)
N2—Cd1—O1	90.88 (11)	C12—C13—H13	118.2

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N1—Cd1—O1	160.61 (12)	C14—C13—H13	118.2
O3—Cd1—O2	147.01 (15)	C13—C14—C9	117.6 (4)
O1W—Cd1—O2	85.21 (10)	C13—C14—C14 ⁱ	114.8 (4)
N2—Cd1—O2	85.37 (12)	C9—C14—C14 ⁱ	127.5 (3)
N1—Cd1—O2	116.74 (12)	N2—C15—C16	123.9 (4)
O1—Cd1—O2	54.80 (9)	N2—C15—H15	118.0
C20—N1—C24	117.8 (4)	C16—C15—H15	118.0
C20—N1—Cd1	118.7 (3)	C15—C16—C17	119.3 (4)
C24—N1—Cd1	123.5 (4)	C15—C16—H16	120.4
C19—N2—C15	117.1 (4)	C17—C16—H16	120.4
C19—N2—Cd1	118.3 (2)	C16—C17—C18	117.0 (4)
C15—N2—Cd1	124.0 (3)	C16—C17—C25	120.9 (4)
C1—O1—Cd1	92.7 (2)	C18—C17—C25	122.1 (4)
C1—O2—Cd1	90.8 (2)	C19—C18—C17	121.2 (4)
C8—O3—Cd1	141.8 (3)	C19—C18—H18	119.4
Cd1—O1W—H1WA	117.7	C17—C18—H18	119.4
Cd1—O1W—H1WB	112.5	N2—C19—C18	121.5 (4)
H1WA—O1W—H1WB	107.7	N2—C19—C20	116.1 (4)
O2—C1—O1	121.7 (4)	C18—C19—C20	122.4 (4)
O2—C1—C2	120.2 (4)	N1—C20—C21	121.1 (4)
O1—C1—C2	118.1 (4)	N1—C20—C19	115.9 (4)
C3—C2—C7	118.8 (4)	C21—C20—C19	123.0 (4)
C3—C2—C1	118.0 (4)	C22—C21—C20	121.1 (5)
C7—C2—C1	123.1 (4)	C22—C21—H21	119.5
C4—C3—C2	121.1 (5)	C20—C21—H21	119.5
C4—C3—H3	119.5	C23—C22—C21	116.9 (4)
C2—C3—H3	119.5	C23—C22—C26	121.2 (4)
C3—C4—C5	120.7 (5)	C21—C22—C26	121.9 (5)
C3—C4—H4	119.7	C24—C23—C22	119.7 (4)
C5—C4—H4	119.7	C24—C23—H23	120.2
C4—C5—C6	119.0 (5)	C22—C23—H23	120.2
C4—C5—H5	120.5	N1—C24—C23	123.4 (5)
C6—C5—H5	120.5	N1—C24—H24	118.3
C5—C6—C7	121.6 (5)	C23—C24—H24	118.3
C5—C6—H6	119.2	C17—C25—H25A	109.5
C7—C6—H6	119.2	C17—C25—H25B	109.5
C6—C7—C2	118.8 (4)	H25A—C25—H25B	109.5
C6—C7—C7 ⁱ	116.4 (3)	C17—C25—H25C	109.5
C2—C7—C7 ⁱ	124.6 (3)	H25A—C25—H25C	109.5
O3—C8—O4	125.9 (6)	H25B—C25—H25C	109.5
O3—C8—C9	115.5 (6)	C22—C26—H26A	109.5
O4—C8—C9	118.7 (4)	C22—C26—H26B	109.5
C10—C9—C14	118.7 (5)	H26A—C26—H26B	109.5
C10—C9—C8	117.9 (5)	C22—C26—H26C	109.5
C14—C9—C8	123.5 (4)	H26A—C26—H26C	109.5
C11—C10—C9	120.3 (6)	H26B—C26—H26C	109.5
C11—C10—H10	119.9		
O3—Cd1—N1—C20	113.7 (3)	C1—C2—C7—C6	-179.5 (4)

O1W—Cd1—N1—C20	-154.3 (3)	C3—C2—C7—C7 ⁱ	-177.3 (5)
N2—Cd1—N1—C20	4.0 (3)	C1—C2—C7—C7 ⁱ	3.9 (8)
O1—Cd1—N1—C20	-10.8 (6)	Cd1—O3—C8—O4	69.7 (8)
O2—Cd1—N1—C20	-69.9 (3)	Cd1—O3—C8—C9	-111.2 (6)
O3—Cd1—N1—C24	-65.9 (4)	O3—C8—C9—C10	-21.7 (6)
O1W—Cd1—N1—C24	26.1 (4)	O4—C8—C9—C10	157.6 (4)
N2—Cd1—N1—C24	-175.6 (4)	O3—C8—C9—C14	157.5 (4)
O1—Cd1—N1—C24	169.6 (3)	O4—C8—C9—C14	-23.3 (6)
O2—Cd1—N1—C24	110.5 (4)	C14—C9—C10—C11	1.4 (7)
O3—Cd1—N2—C19	-95.8 (3)	C8—C9—C10—C11	-179.4 (5)
O1W—Cd1—N2—C19	42.3 (4)	C9—C10—C11—C12	-3.8 (10)
N1—Cd1—N2—C19	-6.6 (3)	C10—C11—C12—C13	2.4 (10)
O1—Cd1—N2—C19	168.5 (3)	C11—C12—C13—C14	1.3 (9)
O2—Cd1—N2—C19	114.0 (3)	C12—C13—C14—C9	-3.6 (7)
O3—Cd1—N2—C15	93.2 (4)	C12—C13—C14—C14 ⁱ	173.3 (5)
O1W—Cd1—N2—C15	-128.7 (3)	C10—C9—C14—C13	2.1 (6)
N1—Cd1—N2—C15	-177.6 (4)	C8—C9—C14—C13	-177.0 (4)
O1—Cd1—N2—C15	-2.5 (3)	C10—C9—C14—C14 ⁱ	-174.3 (5)
O2—Cd1—N2—C15	-57.1 (3)	C8—C9—C14—C14 ⁱ	6.5 (7)
O3—Cd1—O1—C1	166.2 (3)	C19—N2—C15—C16	-1.0 (7)
O1W—Cd1—O1—C1	72.6 (3)	Cd1—N2—C15—C16	170.2 (4)
N2—Cd1—O1—C1	-83.1 (3)	N2—C15—C16—C17	1.1 (7)
N1—Cd1—O1—C1	-69.2 (4)	C15—C16—C17—C18	-0.7 (6)
O2—Cd1—O1—C1	0.6 (2)	C15—C16—C17—C25	179.5 (4)
O3—Cd1—O2—C1	-27.6 (4)	C16—C17—C18—C19	0.2 (6)
O1W—Cd1—O2—C1	-114.0 (3)	C25—C17—C18—C19	-180.0 (4)
N2—Cd1—O2—C1	93.8 (3)	C15—N2—C19—C18	0.4 (6)
N1—Cd1—O2—C1	159.0 (3)	Cd1—N2—C19—C18	-171.2 (3)
O1—Cd1—O2—C1	-0.6 (2)	C15—N2—C19—C20	179.9 (4)
O1W—Cd1—O3—C8	22.1 (7)	Cd1—N2—C19—C20	8.3 (4)
N2—Cd1—O3—C8	-177.0 (7)	C17—C18—C19—N2	-0.1 (6)
N1—Cd1—O3—C8	111.8 (7)	C17—C18—C19—C20	-179.5 (4)
O1—Cd1—O3—C8	-84.2 (7)	C24—N1—C20—C21	-1.3 (6)
O2—Cd1—O3—C8	-62.3 (8)	Cd1—N1—C20—C21	179.0 (3)
Cd1—O2—C1—O1	1.1 (4)	C24—N1—C20—C19	178.2 (4)
Cd1—O2—C1—C2	-177.5 (4)	Cd1—N1—C20—C19	-1.4 (5)
Cd1—O1—C1—O2	-1.1 (4)	N2—C19—C20—N1	-4.5 (5)
Cd1—O1—C1—C2	177.5 (3)	C18—C19—C20—N1	175.0 (4)
O2—C1—C2—C3	151.3 (4)	N2—C19—C20—C21	175.0 (4)
O1—C1—C2—C3	-27.4 (6)	C18—C19—C20—C21	-5.5 (6)
O2—C1—C2—C7	-29.9 (7)	N1—C20—C21—C22	0.8 (6)
O1—C1—C2—C7	151.5 (4)	C19—C20—C21—C22	-178.7 (4)
C7—C2—C3—C4	2.4 (8)	C20—C21—C22—C23	1.0 (7)
C1—C2—C3—C4	-178.7 (5)	C20—C21—C22—C26	-179.8 (4)
C2—C3—C4—C5	-3.3 (8)	C21—C22—C23—C24	-2.1 (7)
C3—C4—C5—C6	2.4 (9)	C26—C22—C23—C24	178.6 (5)
C4—C5—C6—C7	-0.7 (9)	C20—N1—C24—C23	0.1 (7)
C5—C6—C7—C2	-0.1 (8)	Cd1—N1—C24—C23	179.7 (4)

supplementary materials

C5—C6—C7—C7 ⁱ	176.7 (5)	C22—C23—C24—N1	1.7 (8)
C3—C2—C7—C6	-0.7 (7)		

Symmetry codes: (i) $-x, y, -z+1/2$.

Hydrogen-bond geometry (Å, °)

<i>D</i> —H \cdots <i>A</i>	<i>D</i> —H	H \cdots <i>A</i>	<i>D</i> \cdots <i>A</i>	<i>D</i> —H \cdots <i>A</i>
O1W—H1WB \cdots O4 ⁱ	0.85	1.78	2.632 (4)	174
O1W—H1WA \cdots O1 ⁱ	0.85	1.95	2.782 (4)	164

Symmetry codes: (i) $-x, y, -z+1/2$.

Fig. 1

