

catena-Poly[diammonium [diaquabis-(pyridine-2,4-dicarboxylato- $\kappa^2 N, O^2$)-cuprate(II)] [[diaquacopper(II)]- μ -pyridine-2,4-dicarboxylato- $\kappa^3 N, O^2:O^2'$ -[tetraaquacadmium(II)]- μ -pyridine-2,4-dicarboxylato- $\kappa^3 O^2:N, O^2'$] hexahydrate]

Guan-Hua Wang,^{a,b} Zhi-Gang Li,^{a,b} Heng-Qing Jia,^a Ning-Hai Hu^{a*} and Jing-Wei Xu^a

^aThe State Key Laboratory of Electroanalytical Chemistry, Changchun Institute of Applied Chemistry, Chinese Academy of Sciences, Changchun 130022, People's Republic of China, and ^bGraduate School, Chinese Academy of Sciences, Beijing 100039, People's Republic of China
Correspondence e-mail: hunh@ciac.jl.cn

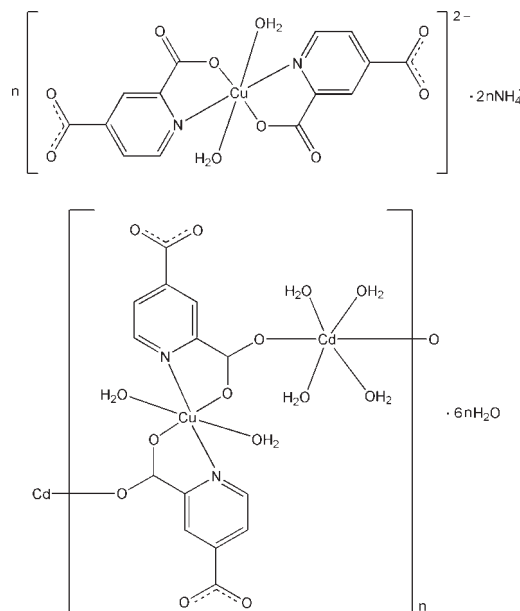
Received 21 October 2009; accepted 6 November 2009

Key indicators: single-crystal X-ray study; $T = 293$ K; mean $\sigma(C-C) = 0.003$ Å; R factor = 0.026; wR factor = 0.073; data-to-parameter ratio = 11.7.

The title mixed-metal complex, $\{(NH_4)_2[Cu(C_7H_3NO_4)_2(H_2O)_2] \cdot (H_2O)_2\} [CdCu(C_7H_3NO_4)_2(H_2O)_6] \cdot 6H_2O\}_n$, contains one octahedrally coordinated Cd^{II} center and two octahedrally coordinated Cu^{II} centers, each lying on an inversion center. The two Cu^{II} atoms are each coordinated by two O atoms and two N atoms from two 2,4-pydc (2,4- H_2 pydc = pyridine-2,4-dicarboxylic acid) ligands in the equatorial plane and two water molecules at the axial sites, thus producing two crystallographically independent $[Cu(2,4-pydc)_2(H_2O)_2]^{2-}$ metalloligands. One metalloligand exists as a discrete anion and the other connects the $Cd(H_2O)_4$ units, forming a neutral chain. $O-H \cdots O$ and $N-H \cdots O$ hydrogen bonds connects the polymeric chains, complex anions, ammonium cations and uncoordinated water molecules into a three-dimensional supramolecular network.

Related literature

For general background to coordination polymers, see: Caneschi *et al.* (2001); Dong *et al.* (2000); Evans & Lin (2002); Kitagawa *et al.* (1999, 2004, 2006). For related structures, see: Li *et al.* (2008); Noro *et al.* (2002a,b); Wang *et al.* (2009).



Experimental

Crystal data

$(NH_4)_2[Cu(C_7H_3NO_4)_2(H_2O)_2] \cdot [CdCu(C_7H_3NO_4)_2(H_2O)_6] \cdot 6H_2O$
 $M_r = 1188.20$
Triclinic, $P\bar{1}$
 $a = 10.4520$ (19) Å
 $b = 10.5252$ (19) Å
 $c = 10.6733$ (19) Å
 $\alpha = 102.869$ (2)°

$\beta = 103.536$ (2)°
 $\gamma = 94.834$ (2)°
 $V = 1101.3$ (3) Å³
 $Z = 1$
Mo $K\alpha$ radiation
 $\mu = 1.54$ mm⁻¹
 $T = 293$ K
 $0.22 \times 0.20 \times 0.16$ mm

Data collection

Bruker SMART APEX CCD diffractometer
Absorption correction: multi-scan (SADABS; Sheldrick, 1996)
 $T_{min} = 0.720$, $T_{max} = 0.785$

6205 measured reflections
4212 independent reflections
3869 reflections with $I > 2\sigma(I)$
 $R_{int} = 0.011$

Refinement

$R[F^2 > 2\sigma(F^2)] = 0.026$
 $wR(F^2) = 0.073$
 $S = 1.05$
4212 reflections
361 parameters
31 restraints

H atoms treated by a mixture of independent and constrained refinement
 $\Delta\rho_{max} = 0.67$ e Å⁻³
 $\Delta\rho_{min} = -0.41$ e Å⁻³

Table 1

Hydrogen-bond geometry (Å, °).

$D-H \cdots A$	$D-H$	$H \cdots A$	$D \cdots A$	$D-H \cdots A$
$O1W-H1A \cdots O7^i$	0.94 (2)	1.80 (1)	2.739 (2)	171 (2)
$O1W-H1B \cdots O4^{ii}$	0.95 (2)	1.82 (1)	2.769 (2)	177 (3)
$O2W-H2A \cdots O3^{ii}$	0.95 (1)	1.72 (1)	2.657 (3)	168 (2)
$O2W-H2B \cdots O8^{iii}$	0.95 (1)	1.77 (1)	2.722 (2)	177 (2)
$O3W-H3A \cdots O6W^{iv}$	0.94 (1)	1.84 (1)	2.781 (3)	172 (3)
$O3W-H3B \cdots O7^{iv}$	0.94 (1)	1.84 (1)	2.776 (3)	172 (3)
$O4W-H4A \cdots O3W^v$	0.95 (1)	1.89 (1)	2.827 (3)	169 (2)
$O4W-H4B \cdots O4^i$	0.95 (1)	1.80 (1)	2.752 (3)	176 (2)
$O5W-H5A \cdots O4W^{vi}$	0.95 (1)	2.10 (1)	3.048 (3)	170 (3)
$O5W-H5B \cdots O3$	0.96 (1)	1.93 (1)	2.882 (3)	171 (3)
$O6W-H6A \cdots O6$	0.95 (1)	1.79 (1)	2.742 (3)	173 (3)

$D-H\cdots A$	$D-H$	$H\cdots A$	$D\cdots A$	$D-H\cdots A$
$O6W-H6B\cdots O2W^{vii}$	0.96 (3)	2.14 (2)	2.991 (3)	147 (2)
$O7W-H7A\cdots O2$	0.95 (3)	2.09 (3)	3.009 (3)	163 (3)
$O7W-H7B\cdots O5W$	0.95 (3)	1.93 (3)	2.861 (3)	165 (4)
$N3-H31\cdots O7W^{viii}$	0.98 (2)	1.90 (2)	2.867 (3)	174 (2)
$N3-H32\cdots O8$	0.99 (2)	1.92 (1)	2.886 (3)	164 (2)
$N3-H33\cdots O5W^{ix}$	0.99 (1)	2.53 (2)	3.208 (4)	126 (2)
$N3-H33\cdots O6^x$	0.99 (1)	2.30 (2)	3.131 (3)	140 (2)
$N3-H33\cdots O6^x$	0.99 (1)	2.19 (2)	2.888 (3)	126 (2)
$N3-H34\cdots O8^{xi}$	0.99 (2)	2.34 (1)	3.277 (3)	157 (2)

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

Data collection: *SMART* (Bruker, 2007); cell refinement: *SAINT* (Bruker, 2007); 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*.

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Supplementary data and figures for this paper are available from the IUCr electronic archives (Reference: BG2303).

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supporting information

Acta Cryst. (2009). E65, m1568–m1569 [doi:10.1107/S1600536809046911]

***catena*-Poly[diammonium [diaquabis(pyridine-2,4-dicarboxylato- κ^2N,O^2)cuprate(II)] [[diaquacopper(II)]- μ -pyridine-2,4-dicarboxylato- $\kappa^3N,O^2:O^{2'}$]-[tetraaquacadmium(II)]- μ -pyridine-2,4-dicarboxylato- $\kappa^3O^2:N,O^{2'}$] hexahydrate]**

Guan-Hua Wang, Zhi-Gang Li, Heng-Qing Jia, Ning-Hai Hu and Jing-Wei Xu

S1. Comment

Coordination polymers constructed from metal ions and bridging ligands have been of great interest owing to their structural diversities and fascinating properties (Caneschi *et al.*, 2001; Evans & Lin, 2002; Kitagawa *et al.*, 1999, 2004). In recent years, the design and synthesis of mixed-metal coordination polymers have received much attention because such heterometallic materials might exhibit interesting physical properties, resulting from interactions between two neighboring metal centers connected by a suitable linker (Dong *et al.*, 2000; Kitagawa *et al.*, 2006). Noro *et al.* (2002a, b) have prepared mixed-metal coordination polymers by using the Et_3NH salt of a metalloligand, $[Cu(2,4-pydc)_2]^{2-}$ (2,4- H_2pydc = pyridine-2,4-dicarboxylic acid). We prepared recently a mixed-metal complex with a metalloligand $[Cu(2,5-pydc)_2]^{2-}$ by a simplified synthetic method (Wang *et al.*, 2009). As a continuation of our work, we report here the synthesis and structure of the title compound.

The asymmetric unit of the title compound contains one six-coordinated Cd^{II} atom and two six-coordinated Cu^{II} atoms, each lying on an inversion center, two 2,4- $pydc$ ligands, one ammonium ion, four coordinated water molecules and three uncoordinated water molecules (Fig. 1). Both $Cu1$ and $Cu2$ atoms have an axially elongated octahedral coordination geometry, defined by two O atoms and two N atoms from two 2,4- $pydc$ ligands in the equatorial plane and two water molecules at the axial sites, thus producing two crystallographically independent $[Cu(2,4-pydc)_2(H_2O)_2]^{2-}$ metalloligands. In each metalloligand, the equatorial plane consists of *trans* N donors and *trans* O donors. The Cd^{II} ions coordinated by four water molecules are linked by the $Cu1$ -metalloligands, *via* the bidentate-bridging 2-carboxylate groups, into a one-dimensional polymeric chain along the [100] direction (Fig. 2). The shortest $Cu\cdots Cd$ distance is 5.226 (1) Å. The 2,4- $pydc$ ligand binds $Cu1$ and $Cd1$ atoms in a $\mu_2-(\kappa^3N,O^2:O^{2'})$ mode with the 4-carboxylate group uncoordinated (Li *et al.*, 2008). The $Cu2$ -metalloligand acts as a discrete divalent anion and does not interact with a second metal ion. The 2,4- $pydc$ ligand in the $Cu2$ -metalloligand adopts a (κ^2N,O^2) chelating mode with the 4-carboxylate group remaining idle. Extensive O—H \cdots O and N—H \cdots O hydrogen bonds (Table 1) assemble the various components into a supramolecular network (Fig. 3).

S2. Experimental

An aqueous solution (20 ml) of $Cu(NO_3)_2 \cdot 3H_2O$ (0.125 g, 0.3 mmol) and a suspension of 2,4- H_2pydc (0.083 g, 0.3 mmol) in ethanol (10 ml) were mixed and refluxed for 24 h until a clear solution was obtained. To this solution, an aqueous solution (5 ml) of $CdCl_2$ (0.055 g, 0.5 mmol) was added. Aqueous NH_3 (25%, 0.06 ml) was then slowly added to the reaction mixture. The resulting solution was filtered off. Blue block crystals were obtained by allowing the filtrate to

stand at room temperature for several days.

S3. Refinement

H atoms on C atoms were positioned geometrically and refined using a riding model, with C—H = 0.93 Å and with $U_{\text{iso}}(\text{H}) = 1.2U_{\text{eq}}(\text{C})$. H atoms of water molecules and ammonium ion were located in a difference Fourier map and refined with distance restraints of O—H = 0.96 (1), H···H = 1.56 (1) Å, and N—H = 0.99 (1), H···H = 1.62 (1) Å, and with $U_{\text{iso}}(\text{H}) = 1.2U_{\text{eq}}(\text{O},\text{N})$.

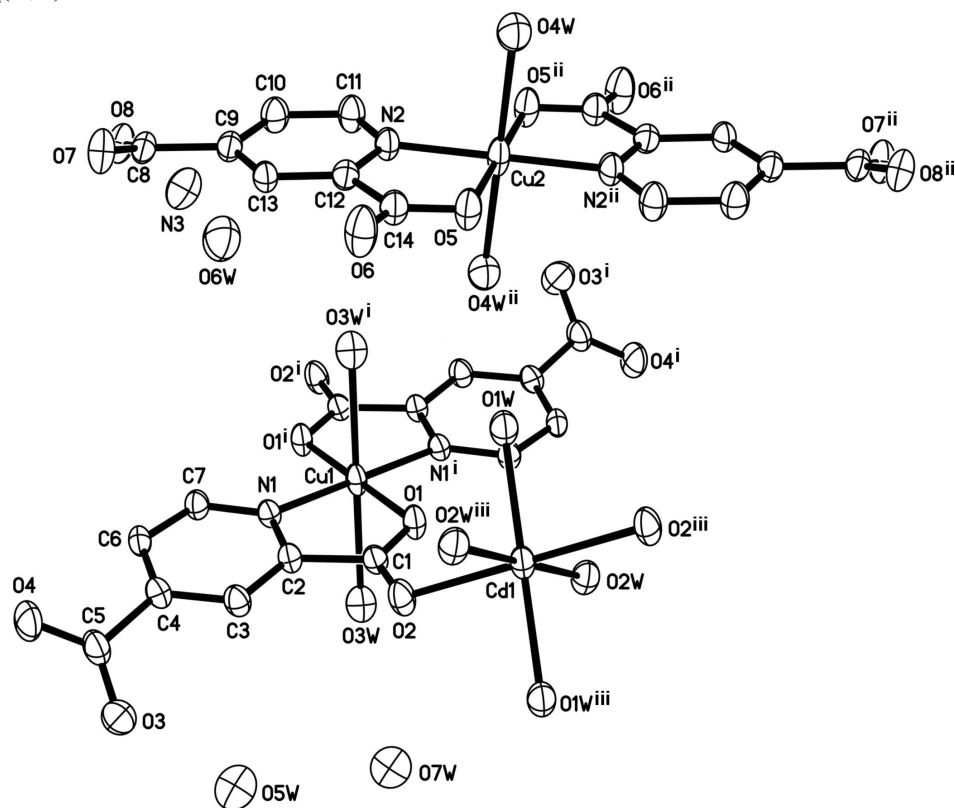


Figure 1

The asymmetric unit of the title compound, together with symmetry-related atoms to complete the Cd1, Cu1 and Cu2 coordination. Displacement ellipsoids are drawn at the 50% probability level. H atoms have been omitted for clarity.

[Symmetry codes: (i) $1 - x, -y, 1 - z$; (ii) $1 - x, 1 - y, 2 - z$; (iii) $-x, -y, 1 - z$.]

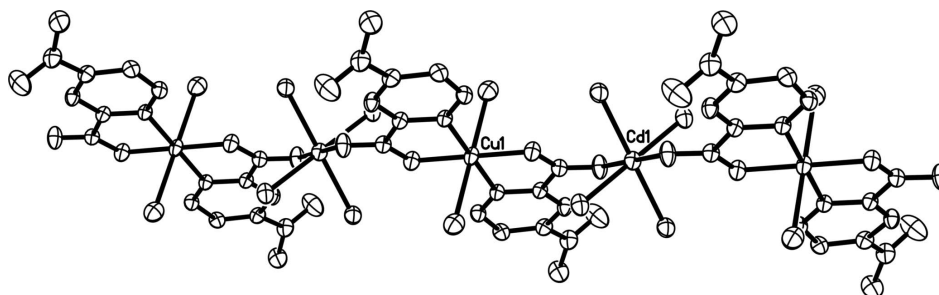


Figure 2

The one-dimensional chain in the title compound. H atoms have been omitted for clarity.

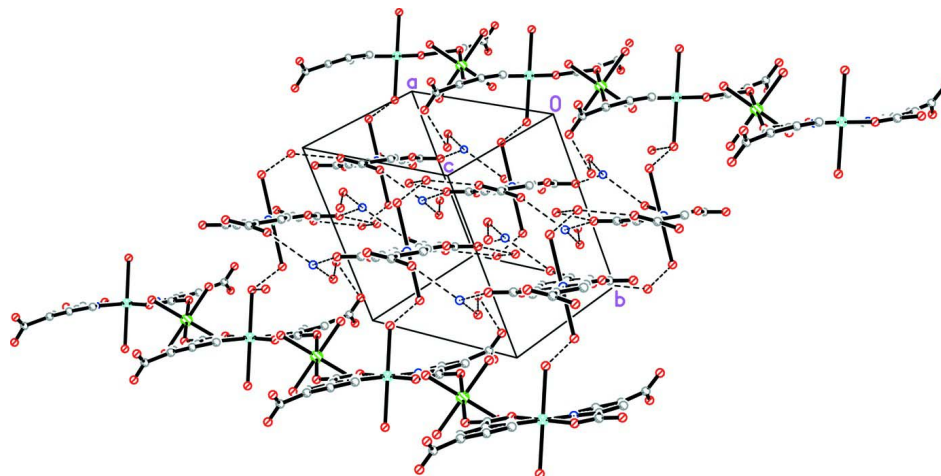


Figure 3

The crystal packing of the title compound. Dashed lines denote hydrogen bonds.

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Crystal data

$(NH_4)_2[Cu(C_7H_3NO_4)_2(H_2O)_2]$
 $[CdCu(C_7H_3NO_4)_2(H_2O)_6] \cdot 6H_2O$

$M_r = 1188.20$

Triclinic, $P\bar{1}$

Hall symbol: -P 1

$a = 10.4520$ (19) Å

$b = 10.5252$ (19) Å

$c = 10.6733$ (19) Å

$\alpha = 102.869$ (2)°

$\beta = 103.536$ (2)°

$\gamma = 94.834$ (2)°

$V = 1101.3$ (3) Å³

$Z = 1$

$F(000) = 604$

$D_x = 1.792$ Mg m⁻³

Mo $K\alpha$ radiation, $\lambda = 0.71073$ Å

Cell parameters from 4140 reflections

$\theta = 2.5$ – 26.1 °

$\mu = 1.54$ mm⁻¹

$T = 293$ K

Block, blue

$0.22 \times 0.20 \times 0.16$ mm

Data collection

Bruker SMART APEX CCD
 diffractometer

Radiation source: fine-focus sealed tube

Graphite monochromator

φ and ω scans

Absorption correction: multi-scan
 (SADABS; Sheldrick, 1996)

$T_{\min} = 0.720$, $T_{\max} = 0.785$

6205 measured reflections

4212 independent reflections

3869 reflections with $I > 2\sigma(I)$

$R_{\text{int}} = 0.011$

$\theta_{\max} = 26.1$ °, $\theta_{\min} = 2.0$ °

$h = -12 \rightarrow 8$

$k = -12 \rightarrow 12$

$l = -11 \rightarrow 13$

Refinement

Refinement on F^2

Least-squares matrix: full

$R[F^2 > 2\sigma(F^2)] = 0.026$

$wR(F^2) = 0.073$

$S = 1.05$

4212 reflections

361 parameters

31 restraints

Primary atom site location: structure-invariant
 direct methods

Secondary atom site location: difference Fourier
 map

Hydrogen site location: inferred from
 neighbouring sites

H atoms treated by a mixture of independent
and constrained refinement
 $w = 1/[\sigma^2(F_o^2) + (0.0412P)^2 + 0.7186P]$
where $P = (F_o^2 + 2F_c^2)/3$

$(\Delta/\sigma)_{\max} = 0.004$
 $\Delta\rho_{\max} = 0.67 \text{ e } \text{\AA}^{-3}$
 $\Delta\rho_{\min} = -0.41 \text{ e } \text{\AA}^{-3}$

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

	x	y	z	$U_{\text{iso}}^*/U_{\text{eq}}$
Cd1	0.0000	0.0000	0.5000	0.02604 (8)
Cu1	0.5000	0.0000	0.5000	0.02460 (10)
Cu2	0.5000	0.5000	1.0000	0.03268 (11)
N1	0.44406 (17)	0.04440 (17)	0.32615 (17)	0.0221 (4)
N2	0.60608 (19)	0.56353 (19)	0.88835 (18)	0.0282 (4)
O1	0.30942 (15)	-0.02363 (16)	0.48585 (15)	0.0281 (3)
O2	0.11497 (15)	-0.03483 (18)	0.34073 (16)	0.0342 (4)
O3	0.1713 (2)	-0.0467 (2)	-0.13119 (19)	0.0591 (6)
O4	0.30140 (19)	0.1332 (2)	-0.12763 (17)	0.0439 (5)
O5	0.35050 (16)	0.49779 (19)	0.84955 (17)	0.0369 (4)
O6	0.30210 (18)	0.5639 (2)	0.6633 (2)	0.0490 (5)
O7	0.71230 (17)	0.72200 (19)	0.51106 (18)	0.0391 (4)
O8	0.89977 (16)	0.65182 (18)	0.59992 (18)	0.0366 (4)
C1	0.2382 (2)	-0.0169 (2)	0.3746 (2)	0.0239 (4)
C2	0.3115 (2)	0.0180 (2)	0.2780 (2)	0.0227 (4)
C3	0.2475 (2)	0.0220 (2)	0.1505 (2)	0.0269 (5)
H3	0.1557	0.0001	0.1186	0.032*
C4	0.3247 (2)	0.0598 (2)	0.0707 (2)	0.0267 (5)
C5	0.2599 (2)	0.0495 (3)	-0.0752 (2)	0.0327 (5)
C6	0.4603 (2)	0.0970 (2)	0.1247 (2)	0.0262 (5)
H6	0.5124	0.1301	0.0763	0.031*
C7	0.5176 (2)	0.0845 (2)	0.2518 (2)	0.0255 (4)
H7	0.6094	0.1046	0.2858	0.031*
C8	0.7820 (2)	0.6710 (2)	0.5938 (2)	0.0280 (5)
C9	0.7192 (2)	0.6332 (2)	0.6975 (2)	0.0270 (5)
C10	0.7977 (2)	0.6251 (3)	0.8193 (2)	0.0337 (5)
H10	0.8898	0.6421	0.8379	0.040*
C11	0.7380 (2)	0.5914 (3)	0.9128 (2)	0.0340 (5)
H11	0.7913	0.5882	0.9948	0.041*
C12	0.5292 (2)	0.5726 (2)	0.7720 (2)	0.0261 (5)
C13	0.5820 (2)	0.6079 (2)	0.6751 (2)	0.0258 (4)
H13	0.5262	0.6147	0.5958	0.031*
C14	0.3820 (2)	0.5433 (2)	0.7580 (2)	0.0305 (5)
O1W	0.14804 (16)	0.17674 (17)	0.64065 (16)	0.0324 (4)
H1A	0.202 (2)	0.205 (3)	0.590 (2)	0.039*
H1B	0.201 (2)	0.159 (3)	0.7184 (16)	0.039*
O2W	0.08911 (16)	-0.15033 (17)	0.60774 (17)	0.0323 (4)
H2A	0.121 (2)	-0.102 (2)	0.6985 (13)	0.039*
H2B	0.023 (2)	-0.2214 (18)	0.603 (2)	0.039*
O3W	0.46396 (19)	-0.24026 (19)	0.37308 (18)	0.0397 (4)

H3A	0.3913 (16)	-0.275 (3)	0.399 (3)	0.048*
H3B	0.5449 (13)	-0.261 (3)	0.419 (3)	0.048*
O4W	0.4544 (2)	0.7280 (2)	1.10086 (19)	0.0445 (4)
H4A	0.447 (3)	0.731 (3)	1.1887 (15)	0.053*
H4B	0.5379 (17)	0.775 (3)	1.106 (3)	0.053*
O5W	0.1659 (2)	-0.2927 (2)	-0.0551 (2)	0.0585 (6)
H5A	0.2575 (14)	-0.294 (3)	-0.016 (3)	0.070*
H5B	0.158 (3)	-0.213 (2)	-0.084 (3)	0.070*
O6W	0.2430 (2)	0.6784 (2)	0.4538 (2)	0.0459 (5)
H6A	0.260 (3)	0.632 (2)	0.522 (2)	0.055*
H6B	0.210 (3)	0.7573 (18)	0.488 (3)	0.055*
O7W	0.0227 (3)	-0.2976 (3)	0.1422 (3)	0.0668 (7)
H7A	0.033 (3)	-0.2138 (19)	0.202 (3)	0.080*
H7B	0.083 (3)	-0.297 (3)	0.088 (3)	0.080*
N3	1.0503 (2)	0.4588 (2)	0.6944 (3)	0.0482 (6)
H31	1.020 (2)	0.406 (2)	0.749 (2)	0.058*
H32	0.9847 (18)	0.5165 (19)	0.666 (2)	0.058*
H33	1.1357 (14)	0.5156 (19)	0.745 (2)	0.058*
H34	1.065 (2)	0.4011 (19)	0.6132 (15)	0.058*

Atomic displacement parameters (\AA^2)

	U^{11}	U^{22}	U^{33}	U^{12}	U^{13}	U^{23}
Cd1	0.02452 (13)	0.03503 (14)	0.02213 (13)	0.00417 (9)	0.00922 (9)	0.01115 (9)
Cu1	0.02026 (19)	0.0393 (2)	0.01908 (19)	0.00659 (15)	0.00640 (15)	0.01509 (16)
Cu2	0.0253 (2)	0.0519 (3)	0.0262 (2)	-0.00001 (18)	0.00792 (17)	0.02147 (19)
N1	0.0217 (9)	0.0281 (9)	0.0180 (8)	0.0046 (7)	0.0060 (7)	0.0077 (7)
N2	0.0284 (10)	0.0354 (10)	0.0239 (9)	0.0021 (8)	0.0081 (8)	0.0132 (8)
O1	0.0248 (8)	0.0418 (9)	0.0230 (8)	0.0056 (7)	0.0091 (6)	0.0158 (7)
O2	0.0214 (8)	0.0579 (11)	0.0241 (8)	0.0019 (7)	0.0082 (7)	0.0113 (8)
O3	0.0686 (15)	0.0667 (14)	0.0275 (10)	-0.0263 (12)	-0.0090 (10)	0.0183 (9)
O4	0.0468 (11)	0.0582 (12)	0.0280 (9)	-0.0027 (9)	0.0038 (8)	0.0235 (9)
O5	0.0264 (9)	0.0565 (11)	0.0323 (9)	-0.0003 (8)	0.0089 (7)	0.0214 (8)
O6	0.0295 (9)	0.0816 (15)	0.0422 (11)	0.0023 (9)	0.0037 (8)	0.0367 (11)
O7	0.0360 (9)	0.0528 (11)	0.0423 (10)	0.0131 (8)	0.0178 (8)	0.0298 (9)
O8	0.0295 (9)	0.0450 (10)	0.0460 (10)	0.0076 (7)	0.0177 (8)	0.0241 (8)
C1	0.0238 (11)	0.0291 (11)	0.0201 (10)	0.0040 (8)	0.0076 (8)	0.0068 (8)
C2	0.0219 (10)	0.0292 (11)	0.0199 (10)	0.0055 (8)	0.0077 (8)	0.0087 (8)
C3	0.0220 (11)	0.0375 (12)	0.0216 (11)	0.0040 (9)	0.0047 (9)	0.0096 (9)
C4	0.0305 (12)	0.0319 (12)	0.0191 (10)	0.0054 (9)	0.0063 (9)	0.0089 (9)
C5	0.0345 (13)	0.0440 (14)	0.0214 (11)	0.0053 (11)	0.0069 (10)	0.0124 (10)
C6	0.0290 (11)	0.0314 (11)	0.0213 (10)	0.0021 (9)	0.0110 (9)	0.0092 (9)
C7	0.0231 (11)	0.0322 (12)	0.0236 (11)	0.0030 (9)	0.0088 (9)	0.0095 (9)
C8	0.0308 (12)	0.0278 (11)	0.0310 (12)	0.0035 (9)	0.0139 (10)	0.0129 (9)
C9	0.0305 (12)	0.0263 (11)	0.0285 (11)	0.0041 (9)	0.0127 (9)	0.0104 (9)
C10	0.0257 (12)	0.0453 (14)	0.0328 (12)	0.0026 (10)	0.0090 (10)	0.0149 (11)
C11	0.0263 (12)	0.0498 (15)	0.0291 (12)	0.0031 (10)	0.0058 (10)	0.0190 (11)
C12	0.0277 (11)	0.0267 (11)	0.0263 (11)	0.0037 (9)	0.0092 (9)	0.0091 (9)

C13	0.0280 (11)	0.0299 (11)	0.0223 (10)	0.0035 (9)	0.0085 (9)	0.0104 (9)
C14	0.0280 (12)	0.0377 (13)	0.0274 (12)	0.0017 (10)	0.0076 (10)	0.0123 (10)
O1W	0.0271 (8)	0.0448 (10)	0.0260 (8)	-0.0001 (7)	0.0065 (7)	0.0126 (7)
O2W	0.0289 (9)	0.0393 (9)	0.0310 (9)	0.0019 (7)	0.0074 (7)	0.0150 (7)
O3W	0.0398 (10)	0.0460 (11)	0.0390 (10)	0.0101 (8)	0.0139 (8)	0.0168 (8)
O4W	0.0488 (12)	0.0475 (11)	0.0367 (10)	-0.0018 (9)	0.0118 (9)	0.0120 (9)
O5W	0.0555 (13)	0.0568 (13)	0.0681 (15)	0.0083 (11)	0.0205 (12)	0.0208 (11)
O6W	0.0532 (12)	0.0450 (11)	0.0466 (11)	0.0098 (9)	0.0174 (9)	0.0205 (9)
O7W	0.0744 (17)	0.0650 (15)	0.0564 (15)	-0.0103 (13)	0.0218 (13)	0.0085 (12)
N3	0.0302 (12)	0.0545 (15)	0.0682 (17)	0.0102 (10)	0.0157 (11)	0.0277 (13)

Geometric parameters (Å, °)

Cd1—O2	2.2850 (16)	C3—H3	0.9300
Cd1—O2 ⁱ	2.2850 (16)	C4—C6	1.384 (3)
Cd1—O1W	2.3004 (17)	C4—C5	1.521 (3)
Cd1—O2W ⁱ	2.2915 (17)	C6—C7	1.389 (3)
Cd1—O2W	2.2915 (17)	C6—H6	0.9300
Cd1—O1W ⁱ	2.3004 (17)	C7—H7	0.9300
Cu1—O1 ⁱⁱ	1.9523 (15)	C8—C9	1.519 (3)
Cu1—O1	1.9523 (15)	C9—C13	1.390 (3)
Cu1—N1	1.9819 (17)	C9—C10	1.389 (3)
Cu1—N1 ⁱⁱ	1.9819 (17)	C10—C11	1.386 (3)
Cu1—O3W ⁱⁱ	2.539 (2)	C10—H10	0.9300
Cu1—O3W	2.539 (2)	C11—H11	0.9300
Cu2—O5	1.9553 (17)	C12—C13	1.385 (3)
Cu2—O5 ⁱⁱⁱ	1.9553 (17)	C12—C14	1.509 (3)
Cu2—N2	1.9862 (18)	C13—H13	0.9300
Cu2—N2 ⁱⁱⁱ	1.9862 (18)	O1W—H1A	0.94 (2)
Cu2—O4W ⁱⁱⁱ	2.535 (2)	O1W—H1B	0.95 (2)
Cu2—O4W	2.535 (2)	O2W—H2A	0.95 (1)
N1—C7	1.334 (3)	O2W—H2B	0.95 (1)
N1—C2	1.342 (3)	O3W—H3A	0.94 (1)
N2—C11	1.336 (3)	O3W—H3B	0.94 (1)
N2—C12	1.340 (3)	O4W—H4A	0.95 (1)
O1—C1	1.266 (3)	O4W—H4B	0.95 (1)
O2—C1	1.239 (3)	O5W—H5A	0.95 (1)
O3—C5	1.246 (3)	O5W—H5B	0.96 (1)
O4—C5	1.244 (3)	O6W—H6A	0.95 (1)
O5—C14	1.275 (3)	O6W—H6B	0.96 (3)
O6—C14	1.224 (3)	O7W—H7A	0.95 (3)
O7—C8	1.255 (3)	O7W—H7B	0.95 (3)
O8—C8	1.253 (3)	N3—H31	0.98 (2)
C1—C2	1.509 (3)	N3—H32	0.99 (2)
C2—C3	1.379 (3)	N3—H33	0.99 (1)
C3—C4	1.397 (3)	N3—H34	0.99 (2)
O2—Cd1—O2 ⁱ	180.00 (6)	N1—C2—C3	122.80 (19)

O2—Cd1—O2W ⁱ	84.25 (6)	N1—C2—C1	114.40 (18)
O2 ⁱ —Cd1—O2W ⁱ	95.75 (6)	C3—C2—C1	122.80 (19)
O2—Cd1—O2W	95.75 (6)	C2—C3—C4	118.1 (2)
O2 ⁱ —Cd1—O2W	84.25 (6)	C2—C3—H3	121.0
O2W ⁱ —Cd1—O2W	180.0	C4—C3—H3	121.0
O2—Cd1—O1W	95.40 (6)	C6—C4—C3	118.78 (19)
O2 ⁱ —Cd1—O1W	84.60 (6)	C6—C4—C5	121.7 (2)
O2W ⁱ —Cd1—O1W	85.58 (6)	C3—C4—C5	119.4 (2)
O2W—Cd1—O1W	94.42 (6)	O4—C5—O3	126.5 (2)
O2—Cd1—O1W ⁱ	84.60 (6)	O4—C5—C4	118.3 (2)
O2 ⁱ —Cd1—O1W ⁱ	95.40 (6)	O3—C5—C4	115.1 (2)
O2W ⁱ —Cd1—O1W ⁱ	94.43 (6)	C4—C6—C7	119.3 (2)
O2W—Cd1—O1W ⁱ	85.57 (6)	C4—C6—H6	120.3
O1W—Cd1—O1W ⁱ	179.999 (1)	C7—C6—H6	120.3
O1 ⁱⁱ —Cu1—O1	180.0	N1—C7—C6	121.5 (2)
O1 ⁱⁱ —Cu1—N1	96.26 (7)	N1—C7—H7	119.2
O1—Cu1—N1	83.74 (7)	C6—C7—H7	119.2
O1 ⁱⁱ —Cu1—N1 ⁱⁱ	83.74 (7)	O8—C8—O7	125.5 (2)
O1—Cu1—N1 ⁱⁱ	96.26 (7)	O8—C8—C9	117.5 (2)
N1—Cu1—N1 ⁱⁱ	180.0	O7—C8—C9	117.0 (2)
O1 ⁱⁱ —Cu1—O3W ⁱⁱ	85.99 (6)	C13—C9—C10	117.8 (2)
O1—Cu1—O3W ⁱⁱ	94.01 (6)	C13—C9—C8	121.3 (2)
N1—Cu1—O3W ⁱⁱ	92.15 (7)	C10—C9—C8	120.8 (2)
N1 ⁱⁱ —Cu1—O3W ⁱⁱ	87.85 (7)	C11—C10—C9	119.8 (2)
O1 ⁱⁱ —Cu1—O3W	94.01 (6)	C11—C10—H10	120.1
O1—Cu1—O3W	85.99 (6)	C9—C10—H10	120.1
N1—Cu1—O3W	87.85 (7)	N2—C11—C10	121.8 (2)
N1 ⁱⁱ —Cu1—O3W	92.15 (7)	N2—C11—H11	119.1
O3W ⁱⁱ —Cu1—O3W	180.00 (4)	C10—C11—H11	119.1
O5—Cu2—O5 ⁱⁱⁱ	180.0	N2—C12—C13	122.2 (2)
O5—Cu2—N2	83.06 (7)	N2—C12—C14	114.18 (19)
O5 ⁱⁱⁱ —Cu2—N2	96.94 (7)	C13—C12—C14	123.7 (2)
O5—Cu2—N2 ⁱⁱⁱ	96.95 (7)	C12—C13—C9	119.4 (2)
O5 ⁱⁱⁱ —Cu2—N2 ⁱⁱⁱ	83.05 (7)	C12—C13—H13	120.3
N2—Cu2—N2 ⁱⁱⁱ	179.998 (1)	C9—C13—H13	120.3
O5—Cu2—O4W ⁱⁱⁱ	94.09 (7)	O6—C14—O5	124.6 (2)
O5 ⁱⁱⁱ —Cu2—O4W ⁱⁱⁱ	85.91 (7)	O6—C14—C12	119.9 (2)
N2—Cu2—O4W ⁱⁱⁱ	86.13 (7)	O5—C14—C12	115.49 (19)
N2 ⁱⁱⁱ —Cu2—O4W ⁱⁱⁱ	93.87 (7)	Cd1—O1W—H1A	106.3 (16)
O5—Cu2—O4W	85.91 (7)	Cd1—O1W—H1B	114.8 (16)
O5 ⁱⁱⁱ —Cu2—O4W	94.09 (7)	H1A—O1W—H1B	110 (1)
N2—Cu2—O4W	93.87 (7)	Cd1—O2W—H2A	104.0 (16)
N2 ⁱⁱⁱ —Cu2—O4W	86.13 (7)	Cd1—O2W—H2B	112.0 (16)
O4W ⁱⁱⁱ —Cu2—O4W	180.0	H2A—O2W—H2B	108 (1)
C7—N1—C2	119.16 (18)	H3A—O3W—H3B	112 (1)
C7—N1—Cu1	129.85 (15)	H4A—O4W—H4B	109 (3)
C2—N1—Cu1	110.76 (13)	H5A—O5W—H5B	108 (3)
C11—N2—C12	119.09 (19)	H6A—O6W—H6B	108 (1)

C11—N2—Cu2	128.85 (15)	H7A—O7W—H7B	110 (3)
C12—N2—Cu2	112.00 (15)	H31—N3—H32	112 (1)
C1—O1—Cu1	113.79 (13)	H31—N3—H33	110 (1)
C1—O2—Cd1	119.57 (14)	H32—N3—H33	108 (1)
C14—O5—Cu2	114.67 (15)	H31—N3—H34	111 (1)
O2—C1—O1	125.21 (19)	H32—N3—H34	108 (1)
O2—C1—C2	118.48 (18)	H33—N3—H34	108 (1)
O1—C1—C2	116.30 (18)		

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

Hydrogen-bond geometry ($\text{\AA}, ^\circ$)

<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>
O1 <i>W</i> —H1 <i>A</i> \cdots O7 ^{iv}	0.94 (2)	1.80 (1)	2.739 (2)	171 (2)
O1 <i>W</i> —H1 <i>B</i> \cdots O4 ^v	0.95 (2)	1.82 (1)	2.769 (2)	177 (3)
O2 <i>W</i> —H2 <i>A</i> \cdots O3 ^v	0.95 (1)	1.72 (1)	2.657 (3)	168 (2)
O2 <i>W</i> —H2 <i>B</i> \cdots O8 ^{vi}	0.95 (1)	1.77 (1)	2.722 (2)	177 (2)
O3 <i>W</i> —H3 <i>A</i> \cdots O6 <i>W</i> ^{vii}	0.94 (1)	1.84 (1)	2.781 (3)	172 (3)
O3 <i>W</i> —H3 <i>B</i> \cdots O7 ^{vii}	0.94 (1)	1.84 (1)	2.776 (3)	172 (3)
O4 <i>W</i> —H4 <i>A</i> \cdots O3 <i>W</i> ^{viii}	0.95 (1)	1.89 (1)	2.827 (3)	169 (2)
O4 <i>W</i> —H4 <i>B</i> \cdots O4 ^{iv}	0.95 (1)	1.80 (1)	2.752 (3)	176 (2)
O5 <i>W</i> —H5 <i>A</i> \cdots O4 <i>W</i> ^{ix}	0.95 (1)	2.10 (1)	3.048 (3)	170 (3)
O5 <i>W</i> —H5 <i>B</i> \cdots O3	0.96 (1)	1.93 (1)	2.882 (3)	171 (3)
O6 <i>W</i> —H6 <i>A</i> \cdots O6	0.95 (1)	1.79 (1)	2.742 (3)	173 (3)
O6 <i>W</i> —H6 <i>B</i> \cdots O2 <i>W</i> ^x	0.96 (3)	2.14 (2)	2.991 (3)	147 (2)
O7 <i>W</i> —H7 <i>A</i> \cdots O2	0.95 (3)	2.09 (3)	3.009 (3)	163 (3)
O7 <i>W</i> —H7 <i>B</i> \cdots O5 <i>W</i>	0.95 (3)	1.93 (3)	2.861 (3)	165 (4)
N3—H31 \cdots O7 <i>W</i> ^{xi}	0.98 (2)	1.90 (2)	2.867 (3)	174 (2)
N3—H32 \cdots O8	0.99 (2)	1.92 (1)	2.886 (3)	164 (2)
N3—H33 \cdots O5 <i>W</i> ^{xii}	0.99 (1)	2.53 (2)	3.208 (4)	126 (2)
N3—H33 \cdots O5 ^{xii}	0.99 (1)	2.30 (2)	3.131 (3)	140 (2)
N3—H33 \cdots O6 ^{xii}	0.99 (1)	2.19 (2)	2.888 (3)	126 (2)
N3—H34 \cdots O8 ^{xiii}	0.99 (2)	2.34 (1)	3.277 (3)	157 (2)

Symmetry codes: (ii) $-x+1, -y, -z+1$; (iv) $-x+1, -y+1, -z+1$; (v) $x, y, z+1$; (vi) $x-1, y-1, z$; (vii) $x, y-1, z$; (viii) $x, y+1, z+1$; (ix) $x, y-1, z-1$; (x) $x, y+1, z$; (xi) $x+1, y+1, z+1$; (xii) $x+1, y, z$; (xiii) $-x+2, -y+1, -z+1$.