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Poly[diaquabis[μ-1-hydroxy-2-(imidazol-3-ium-1-yl)ethane-1,1-diyldiphosphonato]tricopper(II)]

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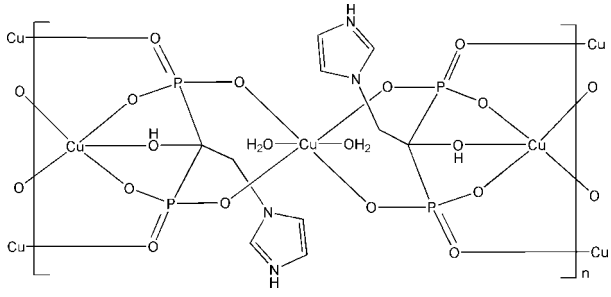
Key indicators: single-crystal X-ray study; T = 293 K; mean σ(C–C) = 0.005 Å; R factor = 0.027; wR factor = 0.073; data-to-parameter ratio = 11.3.

In the title coordination polymer, $[Cu_3(C_5H_7N_2O_7P_2)_2(H_2O)_2]_n$, one Cu^{II} atom is five-coordinated by five O atoms from three 1-hydroxy-2-(imidazol-3-ium-1-yl)ethane-1,1-diyldiphosphonate (L) ligands in a distorted square-pyramidal geometry. The other Cu^{II} atom, lying on an inversion center, is six-coordinated in a distorted octahedral geometry by four O atoms from two L ligands and two O atoms from two water molecules. The five-coordinated Cu^{II} atoms are linked by phosphonate O atoms of the L ligands, forming a polymeric chain. These chains are further linked by the six-coordinated Cu atoms into a layer parallel to $(\bar{1}01)$. N—H···O and O—H···O hydrogen bonds connect the layers into a three-dimensional supramolecular structure.

Related literature

For general background to the applications of metal phosphonates, see: Katz *et al.* (1994).



Experimental

Crystal data

$[Cu_3(C_5H_7N_2O_7P_2)_2(H_2O)_2]$
M_r = 764.81
Triclinic, $P\bar{1}$
a = 7.4167 (9) Å
b = 8.1502 (10) Å
c = 9.5228 (12) Å
α = 104.747 (2)°
β = 107.658 (2)°
γ = 101.484 (2)°
V = 506.03 (11) Å^3
Z = 1
Mo $K\alpha$ radiation
μ = 3.54 mm^{-1}
T = 293 K
0.30 × 0.28 × 0.21 mm

Data collection

Bruker APEX CCD diffractometer
Absorption correction: multi-scan (*SADABS*; Sheldrick, 1996)
T_{min} = 0.58, T_{max} = 0.75
2771 measured reflections
1973 independent reflections
1729 reflections with $I > 2\sigma(I)$
R_{int} = 0.012

Refinement

$R[F^2 > 2\sigma(F^2)]$ = 0.027
$wR(F^2)$ = 0.073
S = 1.05
1973 reflections
175 parameters
2 restraints
H atoms treated by a mixture of independent and constrained refinement
$\Delta\rho_{max}$ = 0.55 e Å^{-3}
$\Delta\rho_{min}$ = −0.68 e Å^{-3}

Table 1
Hydrogen-bond geometry (Å, °).

D—H···A	D—H	H···A	D···A	D—H···A
N2—H2A···O6^{i}	0.86	1.94	2.771 (4)	163
O7—H7···O4	0.82	2.16	2.724 (3)	126
O1W—H1A···O3^{ii}	0.88 (5)	2.09 (3)	2.921 (4)	157 (5)
O1W—H1B···O2^{iii}	0.87 (2)	2.13 (4)	2.851 (4)	140 (4)

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

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) and *DIAMOND* (Brandenburg, 1999); 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: HY2377).

References

Brandenburg, K. (1999). *DIAMOND*. Crystal Impact GbR, Bonn, Germany.
Bruker (2007). *SMART* and *SAINT*. Bruker AXS Inc., Madison, Wisconsin, USA.
Katz, H. E., Wilson, W. L. & Scheller, G. (1994). *J. Am. Chem. Soc.* **116**, 6636–6640.
Sheldrick, G. M. (1996). *SADABS*. University of Göttingen, Germany.
Sheldrick, G. M. (2008). *Acta Cryst.* A**64**, 112–122.

supporting information

Acta Cryst. (2010). E**66**, m1576 [https://doi.org/10.1107/S1600536810046398]

Poly[diaquabis[μ-1-hydroxy-2-(imidazol-3-ium-1-yl)ethane-1,1-diyldiphosphonato]tricopper(II)]

Yaping Li, Dajun Sun, Hu Zang, Liying Han and Guanfang Su

S1. Comment

During the last two decades great research efforts have been devoted to the synthesis and design of metal phosphonates due to their potential applications in electrooptics, ion exchange, catalysis, and stent in intestinal or biliary (Katz *et al.*, 1994). Herein, we present a new copper(II)–phosphonate complex.

The structure analysis reveals that the title compound has a two-dimensional polymeric structure. As shown in Fig. 1, there exist two kinds of crystallographically unique Cu^{II} ions. Atom Cu1 is five-coordinated by four phosphonate O atoms and one hydroxy O atom from three 2-(imidazol-3-ium-1-yl)-1-hydroxy-1,1-ethylidenediphosphonate (*L*) ligands. Atom Cu2 is six-coordinated by four O atoms from two *L* ligands and two O atoms from two water molecules. The Cu1 atoms are linked by the phosphonate O atoms, resulting in a one-dimensional polymeric chain. These chains are further linked by the Cu2 atoms into a layer (Fig. 2). N—H···O and O—H···O hydrogen bonds involving the coordinated water molecules and *L* ligands (Table 1) lead to the formation of a three-dimensional supramolecular network.

S2. Experimental

The synthesis was performed under hydrothermal conditions. A mixture of $CuCl_2.2H_2O$ (0.034 g, 0.2 mmol), *L* ligand (0.070 g, 0.2 mmol) and H_2O (15 ml) in a 25 ml stainless steel reactor with a Teflon liner was heated from 293 to 423 K in 2 h and maintained at 423 K for 72 h. After the mixture was cooled to 298 K, green crystals of the title compound were obtained (yield: 56%).

S3. Refinement

H atoms bound to C, N and hydroxy O were positioned geometrically and refined using a riding model, with C—H = 0.93 and 0.97, N—H = 0.86 and O—H = 0.82 Å and with U_{iso}(H) = 1.2(1.5 for hydroxy)U_{eq}(C,*N*,*O*). H atoms of water molecules were located in a difference Fourier map and refined with U_{iso}(H) = 1.5U_{eq}(O).

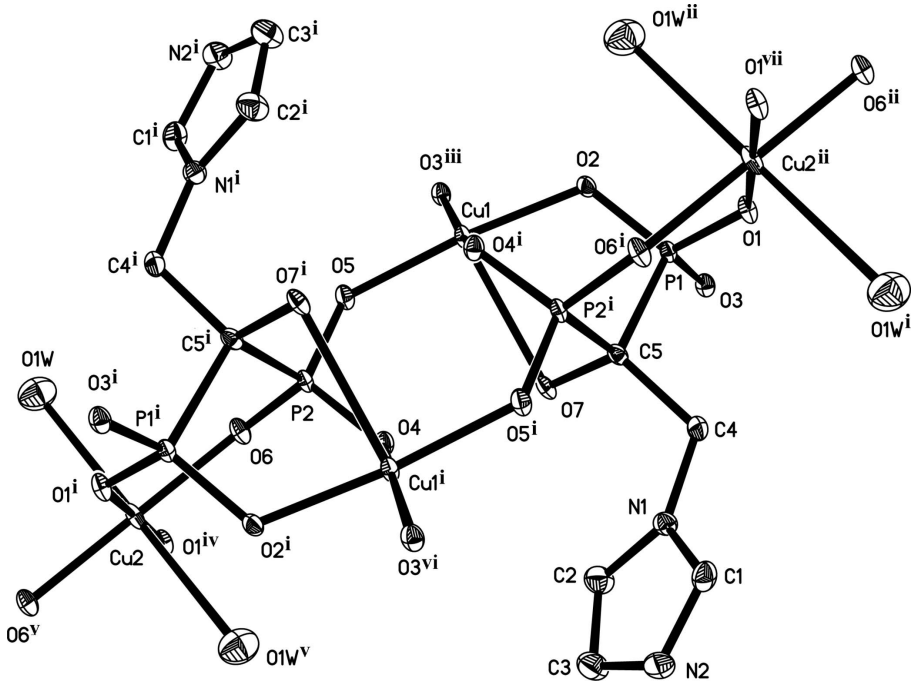

Figure 1

Structure of the title compound. Displacement ellipsoids are drawn at the 50% probability level. H atoms have been omitted for clarity. [Symmetry codes: (i) -*x*, -*y* - 1, -*z*; (ii) *x* - 1, *y* - 1, *z* - 1; (iii) -*x*, -*y* - 2, -*z*; (iv) *x* + 1, *y* + 1, *z* + 1; (v) -*x* + 1, -*y*, -*z* + 1; (vi) *x*, *y* + 1, *z*; (vii) -*x* - 1, -*y* - 2, -*z* - 1.]

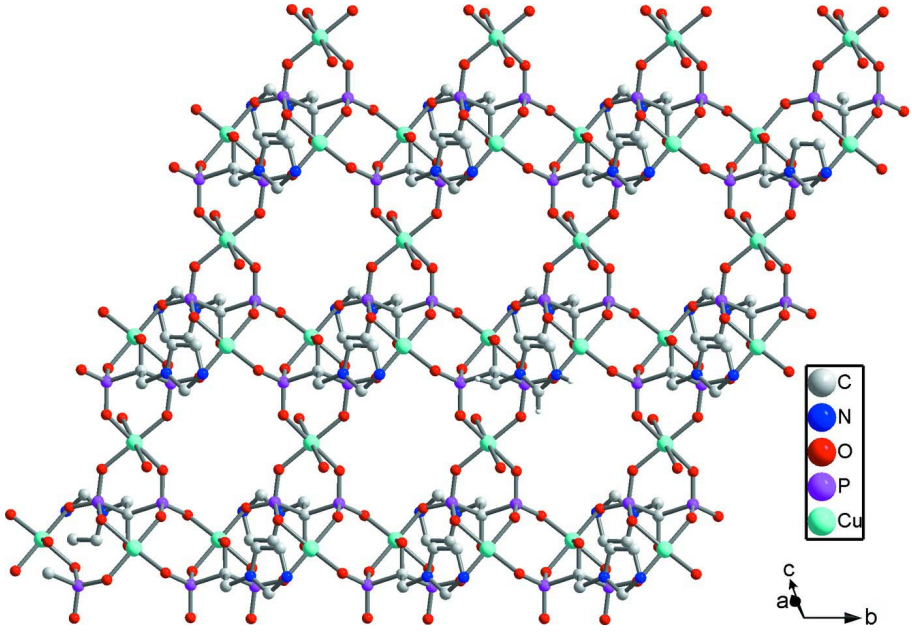

Figure 2

Two-dimensional layer structure in the title compound.

Poly[diaquabis[μ-1-hydroxy-2-(imidazol-3-ium-1-yl)ethane-1,1- diyldiphosphonato]tricopper(II)]

Crystal data

$[Cu_3(C_5H_7N_2O_7P_2)_2(H_2O)_2]$
$M_r = 764.81$
Triclinic, $P\overline{1}$
Hall symbol: -P 1
$a = 7.4167$ (9) Å
$b = 8.1502$ (10) Å
$c = 9.5228$ (12) Å
$\alpha = 104.747$ (2)°
$\beta = 107.658$ (2)°
$\gamma = 101.484$ (2)°
$V = 506.03$ (11) Å^3
$Z = 1$
$F(000) = 381$
$D_x = 2.510$ Mg m^{-3}
Mo $K\alpha$ radiation, $\lambda = 0.71073$ Å
Cell parameters from 1973 reflections
$\theta = 1.9$–$28.3°$
$\mu = 3.54$ mm^{-1}
$T = 293$ K
Block, blue
0.30 × 0.28 × 0.21 mm

Data collection

Bruker APEX CCD diffractometer
Radiation source: fine-focus sealed tube
Graphite monochromator
φ and ω scans
Absorption correction: multi-scan (*SADABS*; Sheldrick, 1996)
$T_{min} = 0.58$, $T_{max} = 0.75$
2771 measured reflections
1973 independent reflections
1729 reflections with $I > 2\sigma(I)$
$R_{int} = 0.012$
$\theta_{max} = 26.1°$, $\theta_{min} = 2.4°$
$h = -9 \rightarrow 8$
$k = -10 \rightarrow 6$
$l = -11 \rightarrow 11$

Refinement

Refinement on F^2
Least-squares matrix: full
$R[F^2 > 2\sigma(F^2)] = 0.027$
$wR(F^2) = 0.073$
$S = 1.05$
1973 reflections
175 parameters
2 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
$w = 1/[\sigma^2(F_o^2) + (0.039P)^2 + 0.8352P]$ where $P = (F_o^2 + 2F_c^2)/3$
$(\Delta/\sigma)_{max} = 0.001$
$\Delta\rho_{max} = 0.55$ e Å^{-3}
$\Delta\rho_{min} = -0.68$ e Å^{-3}

Fractional atomic coordinates and isotropic or equivalent isotropic displacement parameters (Å²)

	x	*y*	*z*	U_{iso}*/U_{eq}
C1	0.2068 (5)	−0.5648 (4)	−0.4517 (4)	0.0140 (7)
H1	0.1047	−0.6119	−0.5497	0.017*
C2	0.3998 (5)	−0.5281 (5)	−0.2131 (4)	0.0173 (7)
H2	0.4524	−0.5475	−0.1190	0.021*
C3	0.4782 (5)	−0.3932 (5)	−0.2542 (4)	0.0186 (7)
H3	0.5935	−0.2995	−0.1928	0.022*
C4	0.0950 (5)	−0.7988 (4)	−0.3483 (4)	0.0122 (7)
H4A	0.1673	−0.8856	−0.3460	0.015*
H4B	−0.0147	−0.8435	−0.4487	0.015*
C5	0.0110 (5)	−0.7833 (4)	−0.2191 (4)	0.0090 (6)
N1	0.2266 (4)	−0.6324 (4)	−0.3368 (3)	0.0110 (6)
N2	0.3567 (4)	−0.4195 (4)	−0.4031 (3)	0.0159 (6)

H2A	0.3750	−0.3518	−0.4568	0.019*
O1	−0.2890 (3)	−1.0888 (3)	−0.3981 (2)	0.0104 (5)
O2	−0.2018 (3)	−0.9799 (3)	−0.1041 (2)	0.0098 (5)
O3	0.0217 (3)	−1.1193 (3)	−0.2195 (2)	0.0099 (5)
O4	0.2060 (3)	−0.3636 (3)	0.0771 (2)	0.0100 (5)
O5	0.0648 (3)	−0.5288 (3)	0.2312 (2)	0.0105 (5)
O6	0.3438 (3)	−0.2484 (3)	0.3787 (3)	0.0115 (5)
O7	0.1644 (3)	−0.7118 (3)	−0.0644 (2)	0.0105 (5)
H7	0.2315	−0.6121	−0.0511	0.016*
P1	−0.12407 (12)	−1.01022 (10)	−0.23716 (9)	0.00780 (18)
P2	0.16460 (12)	−0.34839 (10)	0.22756 (9)	0.00806 (18)
Cu1	−0.07744 (6)	−0.75596 (5)	0.06756 (4)	0.00871 (12)
Cu2	0.5000	0.0000	0.5000	0.01057 (15)
O1W	0.3909 (4)	−0.0271 (4)	0.7275 (3)	0.0282 (6)
H1A	0.302 (6)	−0.054 (7)	0.768 (5)	0.042*
H1B	0.491 (5)	−0.001 (6)	0.815 (4)	0.042*

Atomic displacement parameters (Å²)

	U^{11}	U^{22}	U^{33}	U^{12}	U^{13}	U^{23}
C1	0.0184 (18)	0.0150 (17)	0.0093 (15)	0.0056 (14)	0.0050 (13)	0.0048 (13)
C2	0.0147 (17)	0.0198 (18)	0.0129 (16)	0.0011 (14)	0.0017 (14)	0.0058 (14)
C3	0.0171 (18)	0.0175 (18)	0.0163 (17)	−0.0001 (14)	0.0032 (14)	0.0056 (14)
C4	0.0152 (17)	0.0084 (15)	0.0117 (16)	0.0021 (13)	0.0061 (13)	0.0011 (13)
C5	0.0099 (15)	0.0080 (15)	0.0064 (14)	−0.0002 (12)	0.0010 (12)	0.0026 (12)
N1	0.0119 (13)	0.0090 (13)	0.0128 (13)	0.0032 (11)	0.0050 (11)	0.0040 (11)
N2	0.0209 (16)	0.0133 (15)	0.0156 (14)	0.0029 (12)	0.0077 (12)	0.0089 (12)
O1	0.0133 (12)	0.0068 (11)	0.0075 (11)	0.0025 (9)	0.0004 (9)	0.0009 (9)
O2	0.0126 (11)	0.0060 (11)	0.0093 (10)	0.0004 (9)	0.0045 (9)	0.0019 (9)
O3	0.0137 (11)	0.0072 (11)	0.0089 (11)	0.0032 (9)	0.0036 (9)	0.0032 (9)
O4	0.0134 (11)	0.0080 (11)	0.0091 (11)	0.0042 (9)	0.0044 (9)	0.0029 (9)
O5	0.0149 (12)	0.0075 (11)	0.0072 (11)	0.0021 (9)	0.0028 (9)	0.0020 (9)
O6	0.0127 (11)	0.0082 (11)	0.0086 (11)	0.0016 (9)	−0.0003 (9)	0.0013 (9)
O7	0.0107 (11)	0.0075 (11)	0.0077 (11)	−0.0004 (9)	−0.0008 (9)	0.0008 (9)
P1	0.0101 (4)	0.0055 (4)	0.0064 (4)	0.0017 (3)	0.0022 (3)	0.0014 (3)
P2	0.0104 (4)	0.0049 (4)	0.0062 (4)	0.0012 (3)	0.0014 (3)	0.0006 (3)
Cu1	0.0123 (2)	0.0056 (2)	0.0063 (2)	0.00188 (15)	0.00214 (15)	0.00108 (15)
Cu2	0.0109 (3)	0.0053 (3)	0.0098 (3)	0.0012 (2)	−0.0012 (2)	0.0005 (2)
O1W	0.0233 (15)	0.0366 (17)	0.0267 (15)	0.0095 (13)	0.0093 (12)	0.0132 (13)

Geometric parameters (Å, º)

C1—N2	1.318 (4)	O2—Cu1	1.936 (2)
C1—N1	1.329 (4)	O3—P1	1.529 (2)
C1—H1	0.9300	O3—Cu1^{iii}	1.962 (2)
C2—C3	1.343 (5)	O4—P2	1.534 (2)
C2—N1	1.377 (4)	O4—Cu1^{i}	2.003 (2)
C2—H2	0.9300	O5—P2	1.523 (2)

C3—N2	1.364 (4)	O5—Cu1	1.930 (2)
C3—H3	0.9300	O6—P2	1.521 (2)
C4—N1	1.462 (4)	O6—Cu2	1.959 (2)
C4—C5	1.528 (4)	O7—H7	0.8200
C4—H4A	0.9700	P2—C5^{i}	1.842 (3)
C4—H4B	0.9700	Cu1—O3^{iii}	1.962 (2)
C5—O7	1.444 (4)	Cu1—O4^{i}	2.003 (2)
C5—P2^{i}	1.842 (3)	Cu2—O1^{i}	1.950 (2)
C5—P1	1.857 (3)	Cu2—O1^{iv}	1.950 (2)
N2—H2A	0.8600	Cu2—O6^{v}	1.959 (2)
O1—P1	1.519 (2)	O1W—H1A	0.88 (5)
O1—Cu2^{ii}	1.950 (2)	O1W—H1B	0.87 (2)
O2—P1	1.530 (2)		

N2—C1—N1	108.3 (3)	P2—O4—Cu1^{i}	119.28 (13)
N2—C1—H1	125.8	P2—O5—Cu1	131.88 (14)
N1—C1—H1	125.8	P2—O6—Cu2	136.89 (14)
C3—C2—N1	107.0 (3)	C5—O7—H7	109.5
C3—C2—H2	126.5	O1—P1—O3	111.33 (12)
N1—C2—H2	126.5	O1—P1—O2	112.88 (13)
C2—C3—N2	107.0 (3)	O3—P1—O2	112.51 (12)
C2—C3—H3	126.5	O1—P1—C5	106.82 (13)
N2—C3—H3	126.5	O3—P1—C5	108.50 (14)
N1—C4—C5	114.6 (3)	O2—P1—C5	104.30 (13)
N1—C4—H4A	108.6	O6—P2—O5	109.75 (13)
C5—C4—H4A	108.6	O6—P2—O4	114.99 (13)
N1—C4—H4B	108.6	O5—P2—O4	112.22 (12)
C5—C4—H4B	108.6	O6—P2—C5^{i}	107.03 (13)
H4A—C4—H4B	107.6	O5—P2—C5^{i}	108.36 (14)
O7—C5—C4	112.5 (3)	O4—P2—C5^{i}	104.02 (13)
O7—C5—P2^{i}	108.6 (2)	O5—Cu1—O2	174.89 (9)
C4—C5—P2^{i}	114.1 (2)	O5—Cu1—O3^{iii}	91.03 (9)
O7—C5—P1	105.3 (2)	O2—Cu1—O3^{iii}	91.03 (9)
C4—C5—P1	108.5 (2)	O5—Cu1—O4^{i}	90.70 (9)
P2^{i}—C5—P1	107.27 (16)	O2—Cu1—O4^{i}	88.63 (9)
C1—N1—C2	108.3 (3)	O3^{iii}—Cu1—O4^{i}	163.80 (9)
C1—N1—C4	124.8 (3)	O1^{i}—Cu2—O1^{iv}	180.00 (13)
C2—N1—C4	126.7 (3)	O1^{i}—Cu2—O6	92.58 (9)
C1—N2—C3	109.3 (3)	O1^{iv}—Cu2—O6	87.42 (9)
C1—N2—H2A	125.3	O1^{i}—Cu2—O6^{v}	87.42 (9)
C3—N2—H2A	125.3	O1^{iv}—Cu2—O6^{v}	92.58 (9)
P1—O1—Cu2^{ii}	131.22 (13)	O6—Cu2—O6^{v}	180.00 (19)
P1—O2—Cu1	118.08 (13)	H1A—O1W—H1B	93 (4)
P1—O3—Cu1^{iii}	125.96 (13)		

N1—C2—C3—N2	−2.0 (4)	P2^{i}—C5—P1—O1	63.64 (18)
N1—C4—C5—O7	−56.4 (4)	O7—C5—P1—O3	−60.7 (2)
N1—C4—C5—P2^{i}	67.9 (3)	C4—C5—P1—O3	60.0 (2)

N1—C4—C5—P1	−172.6 (2)	P2^{i}—C5—P1—O3	−176.24 (13)
N2—C1—N1—C2	−2.1 (4)	O7—C5—P1—O2	59.4 (2)
N2—C1—N1—C4	−176.7 (3)	C4—C5—P1—O2	−179.9 (2)
C3—C2—N1—C1	2.5 (4)	P2^{i}—C5—P1—O2	−56.12 (17)
C3—C2—N1—C4	177.0 (3)	Cu2—O6—P2—O5	−156.45 (19)
C5—C4—N1—C1	−127.4 (3)	Cu2—O6—P2—O4	75.9 (2)
C5—C4—N1—C2	58.9 (4)	Cu2—O6—P2—C5^{i}	−39.1 (2)
N1—C1—N2—C3	0.9 (4)	Cu1—O5—P2—O6	−147.75 (17)
C2—C3—N2—C1	0.7 (4)	Cu1—O5—P2—O4	−18.6 (2)
Cu2^{ii}—O1—P1—O3	−172.39 (16)	Cu1—O5—P2—C5^{i}	95.7 (2)
Cu2^{ii}—O1—P1—O2	60.0 (2)	Cu1^{i}—O4—P2—O6	−115.31 (15)
Cu2^{ii}—O1—P1—C5	−54.1 (2)	Cu1^{i}—O4—P2—O5	118.31 (14)
Cu1^{iii}—O3—P1—O1	−118.20 (16)	Cu1^{i}—O4—P2—C5^{i}	1.40 (18)
Cu1^{iii}—O3—P1—O2	9.7 (2)	P2—O5—Cu1—O3^{iii}	156.80 (19)
Cu1^{iii}—O3—P1—C5	124.53 (16)	P2—O5—Cu1—O4^{i}	−39.31 (19)
Cu1—O2—P1—O1	−134.92 (14)	P1—O2—Cu1—O3^{iii}	−124.84 (14)
Cu1—O2—P1—O3	98.05 (15)	P1—O2—Cu1—O4^{i}	71.36 (15)
Cu1—O2—P1—C5	−19.34 (18)	P2—O6—Cu2—O1^{i}	19.3 (2)
O7—C5—P1—O1	179.20 (18)	P2—O6—Cu2—O1^{iv}	−160.7 (2)
C4—C5—P1—O1	−60.1 (2)		

Symmetry codes: (i) −*x*, −*y*−1, −*z*; (ii) *x*−1, *y*−1, *z*−1; (iii) −*x*, −*y*−2, −*z*; (iv) *x*+1, *y*+1, *z*+1; (v) −*x*+1, −*y*, −*z*+1.

Hydrogen-bond geometry (Å, º)

D—H···*A*	*D*—H	H···*A*	*D*···*A*	*D*—H···*A*
N2—H2*A*···O6^{vi}	0.86	1.94	2.771 (4)	163
O7—H7···O4	0.82	2.16	2.724 (3)	126
O1*W*—H1*A*···O3^{vii}	0.88 (5)	2.09 (3)	2.921 (4)	157 (5)
O1*W*—H1*B*···O2^{iv}	0.87 (2)	2.13 (4)	2.851 (4)	140 (4)

Symmetry codes: (iv) *x*+1, *y*+1, *z*+1; (vi) *x*, *y*, *z*−1; (vii) *x*, *y*+1, *z*+1.