Di-μ-acetato-κ^4O:O'-bis[(1,10-phenanthroline-κ^2N,N')(trifluoromethanesulfonato-κO)copper(II)]

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The complete molecule of the title compound, $[Cu_2(C_2H_3O_2)_2(CF_3O_3S)_2(C_{12}H_8N_2)_2]$, is completed by the application of a twofold rotation and comprises two Cu^{II} ions, each of which is pentacoordinated by two N atoms from a bidentate 1,10-phenanthroline (phen) ligand, two O atoms from acetate ligands and an O atom from a trifluoromethanesulfonate anion, forming a (4 + 1) distorted square-pyramidal coordination geometry. The Cu^{II} ions are connected by two acetate bridges in a *syn–syn* configuration. The F atoms of the trifluoromethanesulfonate ligands are disordered, with site-occupation factors of 70 and 30. The molecular structure is stabilized by intramolecular face-to-face π–π interactions with centroid–centroid distances in the range 3.5654 (12)–3.8775(12) Å. The crystal structure is stabilized by C—H···O interactions, leading to a three-dimensional lattice structure.

Related literature

For general background to this work, see: Moreira *et al.* (2007); Calvo *et al.* (2011); Reinoso *et al.* (2005, 2007); Ritchie *et al.* (2006); Wang *et al.* (2006). For literature used in the synthetic procedures, see: Youngme *et al.* (2008). For a related crystal structure, see: Tokii *et al.* (1990). For potential applications, see: Hill & Brown (1986); Mansuy *et al.* (1991); Hill & Zhang (1995). For an explanation of the τ parameter, see: Addison *et al.* (1984). For spectroscopic properties, see: Castro *et al.* (1992); Sletten & Julve (1999).
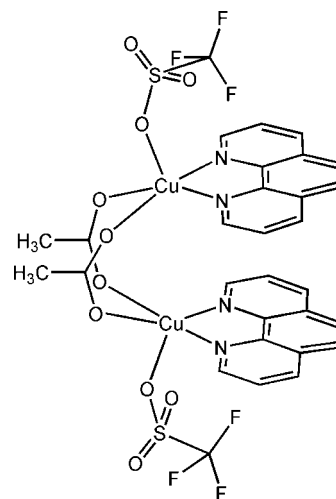
Experimental

Crystal data

$[Cu_2(C_2H_3O_2)_2(CF_3O_3S)_2(C_{12}H_8N_2)_2]$	β = 95.507 (1)°
M_r = 903.72	V = 3447.0 (2) Å^3
Monoclinic, $C2/c$	Z = 4
a = 13.1198 (5) Å	Mo $K\alpha$ radiation
b = 16.1282 (6) Å	μ = 1.45 mm^{-1}
c = 16.3659 (6) Å	T = 293 K
	0.24 × 0.21 × 0.18 mm

Data collection

Bruker SMART APEX CCD diffractometer	23313 measured reflections
Absorption correction: multi-scan (*SADABS*; Sheldrick, 2000)	4178 independent reflections
	3491 reflections with $I > 2\sigma(I)$
T_{min} = 0.872, T_{max} = 1.000	R_{int} = 0.022

Refinement

$R[F^2 > 2\sigma(F^2)]$ = 0.034	272 parameters
$wR(F^2)$ = 0.102	H-atom parameters constrained
S = 1.04	$\Delta\rho_{max}$ = 0.37 e Å^{-3}
4178 reflections	$\Delta\rho_{min}$ = −0.32 e Å^{-3}

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

D—H···A	D—H	H···A	D···A	D—H···A
C1—H1···O5^i	0.93	2.43	3.229 (3)	144
C3—H3···O4^{ii}	0.93	2.34	3.213 (3)	157
C8—H8···O3^{iii}	0.93	2.56	3.421 (3)	154

Symmetry codes: (i) $-x+1, -y+2, -z+1$; (ii) $x-\frac{1}{2}, y-\frac{1}{2}, z$; (iii) $-x+\frac{3}{2}, y-\frac{1}{2}, -z+\frac{3}{2}$.

Data collection: *SMART* (Bruker, 2000); cell refinement: *SMART*; data reduction: *SAINT* (Bruker, 2000) and *SHELXTL* (Sheldrick, 2008); program(s) used to solve structure: *SHELXTL*; program(s) used to refine structure: *SHELXL97* (Sheldrick, 2008); molecular graphics: *ORTEP-3 for Windows* (Farrugia, 2012) and *Mercury* (Macrae *et al.*, 2006); software used to prepare material for publication: *PLATON* (Spek, 2009) and *pubCIF* (Westrip, 2010).

metal-organic compounds

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supporting information

Di-*µ*-acetato-*κ⁴O:O′*-bis[(1,10-phenanthroline-*κ²N,N′*)(trifluoromethane-sulfonato-*κO*)copper(II)]

Nanthawat Wannarit, Chaveng Pakawatchai and Sujittra Youngme

S1. Comment

The synthesis and characterization of polycarboxylato-bridged dinuclear copper(II) compounds namely dinuclear tetra-carboxylato-bridged Cu^{II} compounds (paddlewheel-like structure) (*e.g.* Moreira *et al.*, 2007; Youngme *et al.*, 2008) and also dinuclear Cu^{II} compounds containing dicarboxylato-bridges (Tokii *et al.*, 1990; Reinoso *et al.*, 2005; Ritchie *et al.*, 2006) have attacted much attention in several years. These compounds have been prepared with the aim of studying their intramolecular magnetic properties which are determined predominantly by strong antiferromagnetic interactions. In addition, the dicarboxylato-bridged dinuclear Cu^{II} compounds have been frequently used as the models for the basic understanding of their magneto-structural correlations in theoretical studies (Moreira *et al.*, 2007; Calvo *et al.*, 2011). Copper(II) compounds containing doubly acetato-bridged dinuclear units, [Cu(phen)(*µ*-OOCCH$_3$)$_2$Cu(phen)]$^{2+}$ (where phen = 1,10-phenanthroline), have also been shown to exhibit antiferromagnetic behavior (Tokii *et al.*, 1990). Furthermore, this type of dinuclear unit was used as the secondary building block in functionalized polyoxometalate (POMs) materials (Wang *et al.*, 2006; Reinoso *et al.*, 2007; Calvo *et al.*, 2011) to extend the dimensionality of structures leading to new hybrid materials and more selective applications, for example catalytic properties in organic oxidations (Hill & Brown, 1986; Mansuy *et al.*, 1991; Hill, & Zhang, 1995).

A new doubly acetato-bridged dinuclear Cu^{II} compound containing additional trifluoromethanesulfonate anions has been synthesized and its structural features are reported here. Compound **I**, bis((*µ*-acetato)(trifluoromethanesulfonato) (1,10-phenanthroline))dicopper(II) crystallized in the space group *C*2/*c* with an asymmetric unit containing one half of the dinuclear unit (Fig.1). This dinuclear unit has *C*$_2$ symmetry around the *b* axis with Cu···Cu distance of 3.0309 (4) Å. Structurally, compound **I** consists of two [Cu(phen)(OSO$_2$CF$_3$)]$^+$ cations connected together by two bridging acetato ligands in a *syn-syn* configuration. Both Cu^{II} atoms exhibit five coordination of CuN$_2$O$_2$O′ chromophore, with the basal plane consisting of two phen N atoms [Cu—N = 2.0153 (18) and 1.9980 (18) Å] and two O atoms from acetate ligands [Cu—O = 1.9387 (17) and 1.9377 (18)]. Due to symmetry both square planes are parallel to one another. The apical position at Cu^{II} is occupied by an O atom from trifluoromethanesulfonate anion [Cu—O = 2.261 (2)], leading to the (4 + 1) square-pyramidal geometry. The square base of Cu^{II} chromophore is not perfectly planar, with the tetrahedral twist of 16.52 (7)° and Cu^{II} is situated above the basal plane by 0.14 (1) Å pointing towards the O atom of the trifluoromethane-sulfonate anion. The distortion of a square pyramid can be best described by the structural parameter *τ* (*τ* = 0 for a square pyramid and *τ* =1 for a trigonal bipyramid (Addison *et al.*, 1984)), with *τ* = 0.23 for the title compound. The molecular structure of **I** reveals intramolecular face-to-face *π*-*π* interaction between aromatic rings of phen ligands (Fig. 1). Phenanthroline molecules are parallel with an average contact and angle of phen planes of 3.63 (3) Å and 5.96 (3)°, respectively. In general, the ligands are featureless: neither of phen group departs significantly from planarity [maximum deviations: 0.082 Å for C11 and 0.099 Å for C10 of *Cg*3 ring(N2, C6, C9, C10, C11, C12)] and the C—O bonds in the

acetato bridging ligands display an almost perfect resonance [C13⋯O1 = 1.257 (3) Å and C13⋯O2 = 1.250 (3) Å]. The crystal structure of compound **I** is determined by intermolecular hydrogen bonding interactions between methyl groups of acetato ligands (H14A) or phen ligands (H7 as hydrogen bond donor sites and H10) and oxygen/fluoride accepetors at trifluoromethanesulfonate anions (O3, O5 and F3) (see Table 1), generating two-dimensional layers parallel to the *ab* plane (Fig. 2). Moreover, these two-dimensional sheets are interconnected by hydrogen bond interactions between C—H of phen ligands and oxygen atoms of trifluoromethanesulfonate anions [C1—H1⋯O5^i; symetry code (i) = -*x*+1, -*y*+2, -*z*+1] (see Table 2) in direction of crystallographic *c* axis, leading to three-dimensional lattice structure (Fig. 3). Although containing the same [(phen)Cu(μ-OOCCH$_3$)$_2$Cu(phen)]$^{2+}$ unit, the structural topology of **I** is distinct from that of the related compound [Cu(phen)(μ-O$_2$CCH$_3$)(H$_2$O)]$_2$(NO$_3$)$_2$.4H$_2$O (Tokii *et al.*, 1990) in which the apical position is occupied by water molecule. The dinuclear unit of this related compound also crystallized in *C*2/*c* space group and has *C*$_2$ symmetry around the *b* axis with Cu⋯Cu distance of 3.063 Å, but its crystal lattice is mainly stabilized by intra- and intermolecular π-π interactions, generating a one-dimensional chain-like sructure. It is clear that the difference of the structural topology between compound **I** and the related compound caused by the effect of coordinated trifluoromethane-sulfonate anions whereas nitrate anions are not coordinated to Cu in the other structure.

The diffuse reflectance spectrum of **I** displays a broad band at 15400 cm^{-1} and a lower energy shoulder at 14300 cm^{-1}. This feature corresponds to a dominantly distorted square pyramidal geometry of Cu^{II} ions and is consistent with the observed structural parameters. The transitions may be assigned as d$_{xy}$, d$_{yz}$, d$_{xz}$→ d$_{x2-y2}$ and d$_{z2}$→ d$_{x2-y2}$. The IR spectrum of **I**, in addition to the phen vibrations shows the broad and intense bands of the stretching of the ionic CF$_3$SO$_3$$^-$ at 1276 v_{as}(S–O), 1158 v_{as}(C–F) and 1031 v_s(S–O) cm^{-1} (Castro *et al.*, 1992). The IR spectrum also shows two broad and intense bands at 1567 and 1385 cm^{-1}, corresponding to the v_{as}(COO$^-$) and v_s(COO$^-$) vibrations of acetate bridging ligands. The latter spectral properties completely disappear for related mononuclear compounds as [Cu(phen)$_3$](CF$_3$SO$_3$)$_2$.H$_2$O (Sletten & Julve, 1999).

S2. Experimental

A warm ethanolic solution (25 ml) of phen (0.198 g, 1.0 mmol) was added to a warm aqueous solution (15 ml) of Cu(CF$_3$SO$_3$)$_2$ (0.370 g, 1.0 mmol). Then NaO$_2$CCH$_3$ solid (0.124 g, 1.0 mmol) was added to the mixture, yielding a clear dark blue solution. After a week, the blue rectangle-shaped crystals of compound **I** were obtained. The crystals were filtered off, washed with mother liquor and air-dried. Yield: *ca* 45%. Anal. Calc. for Cu$_2$C$_{30}$H$_{24}$N$_4$O$_{10}$F$_6$S$_2$: C, 39.78; H, 2.67; N, 6.19%. Found: C, 39.12; H, 2.51; N, 6.36%.

S3. Refinement

All H atoms were constrained to ideal positions, with C—H = 0.93 Å and U_{iso}(H) =1.2U_{eq}(C) for H atoms at phen and C—H = 0.96 Å and U_{iso}(H) =1.2U_{eq}(C) for H atoms of acetate groups. Fluorine atoms of the trifluoromethanesulfonato ligands are disordered with site occupation factors of 70:30%.
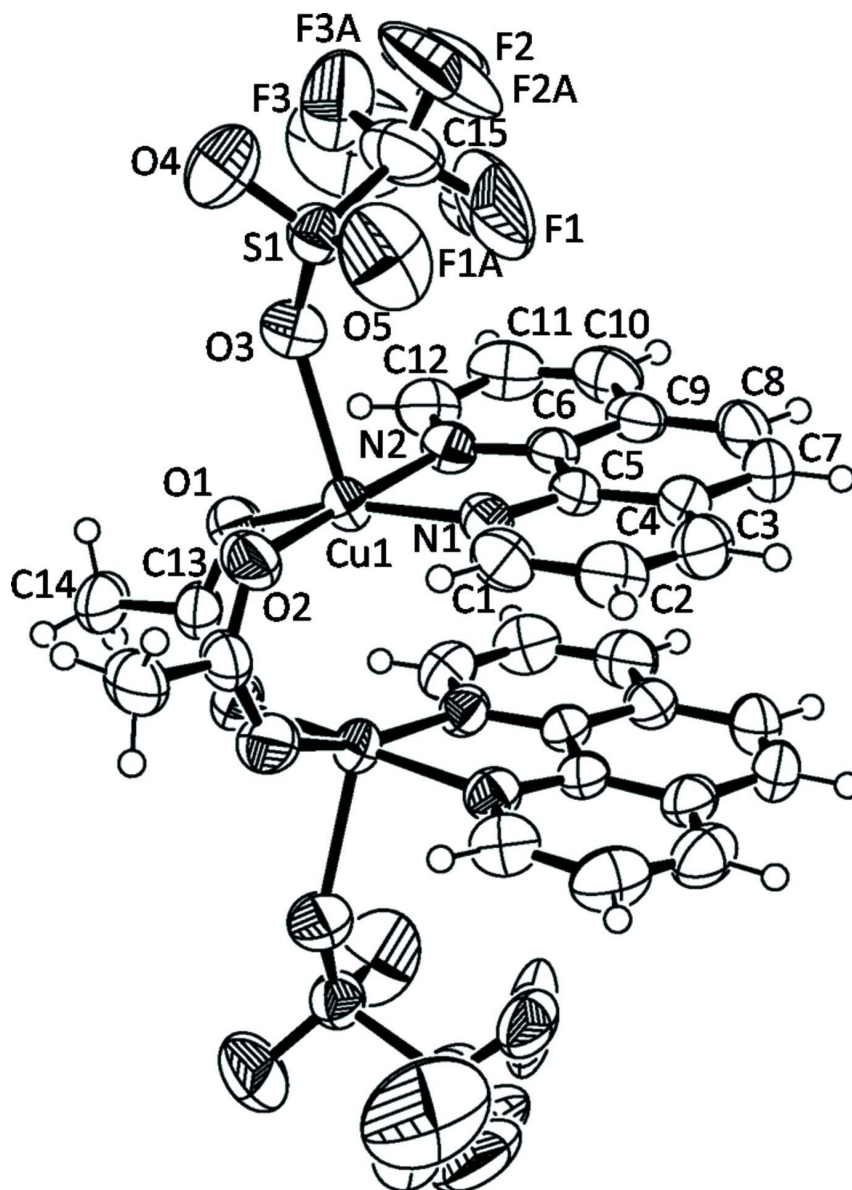
Figure 1

Molecular structure and atomic numbering scheme with thermal ellipsoids shown at 50% probability level.
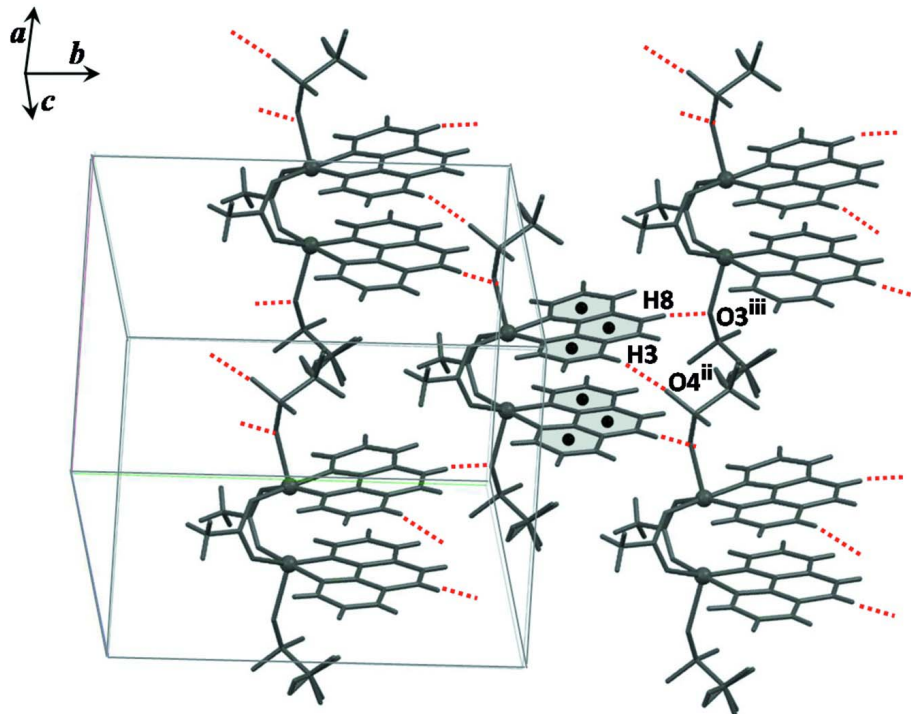
Figure 2

The crystal packing. View of two-dimensional layer constructed by intermolecular hydrogen bonding and view of the intramolecular face-to-face π-π interactions between aromatic rings of phen ligands.
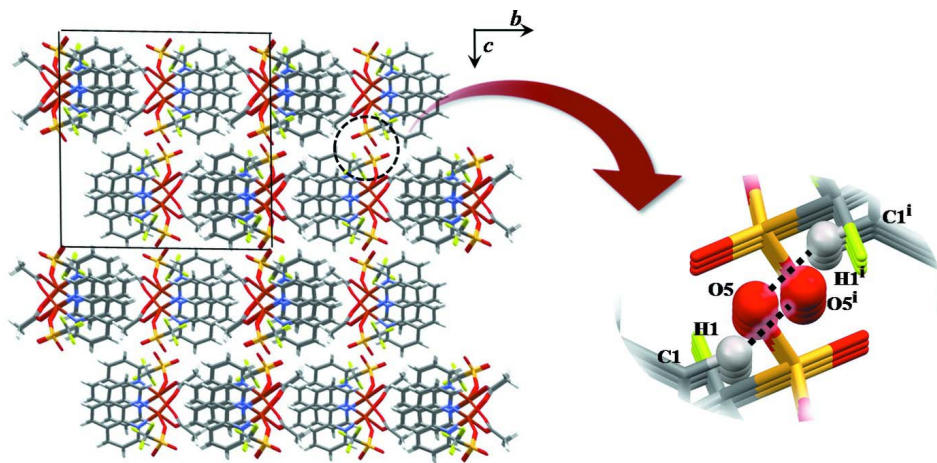
Figure 3

The crystal packing. View of three-dimensional framework (side view) constructued by intermolecular hydrogen bonding interactions between two-dimensional layers (C1—H1···O5[i], symmetry code: (i) = -x+1, -y+2, -z+1).

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Crystal data

[Cu₂(C₂H₃O₂)₂(CF₃O₃S)₂(C₁₂H₈N₂)₂]	$F(000) = 1816$
$M_r = 903.72$	$D_x = 1.741$ Mg m⁻³
Monoclinic, $C2/c$	Mo $K\alpha$ radiation, $\lambda = 0.71073$ Å
Hall symbol: -C 2yc	Cell parameters from 8485 reflections
$a = 13.1198$ (5) Å	$\theta = 2.3–26.4°$
$b = 16.1282$ (6) Å	$\mu = 1.45$ mm⁻¹
$c = 16.3659$ (6) Å	$T = 293$ K
$\beta = 95.507$ (1)°	Block, blue
$V = 3447.0$ (2) Å³	0.24 × 0.21 × 0.18 mm
$Z = 4$	

Data collection

Bruker SMART APEX CCD diffractometer	23313 measured reflections
Radiation source: fine-focus sealed tube	4178 independent reflections
Graphite monochromator	3491 reflections with $I > 2\sigma(I)$
phi and ω scans	$R_{int} = 0.022$
Absorption correction: multi-scan (*SADABS*; Sheldrick, 2000)	$\theta_{max} = 28.1°$, $\theta_{min} = 2.0°$
$T_{min} = 0.872$, $T_{max} = 1.000$	$h = -17\rightarrow17$
	$k = -21\rightarrow21$
	$l = -21\rightarrow21$

Refinement

Refinement on F^2	Secondary atom site location: difference Fourier map
Least-squares matrix: full	
$R[F^2 > 2\sigma(F^2)] = 0.034$	Hydrogen site location: inferred from neighbouring sites
$wR(F^2) = 0.102$	
$S = 1.04$	H-atom parameters constrained
4178 reflections	$w = 1/[\sigma^2(F_o^2) + (0.0624P)^2 + 1.5014P]$ where $P = (F_o^2 + 2F_c^2)/3$
272 parameters	
0 restraints	$(\Delta/\sigma)_{max} = 0.001$
Primary atom site location: structure-invariant direct methods	$\Delta\rho_{max} = 0.37$ e Å⁻³
	$\Delta\rho_{min} = -0.32$ e Å⁻³

Special details

Geometry. Bond distances, angles *etc*. have been calculated using the rounded fractional coordinates. All e.s.d.'s are estimated from the variances of the (full) variance-covariance matrix. The cell e.s.d.'s are taken into account in the estimation of distances, angles and torsion angles

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

	x	*y*	*z*	U_{iso}*/U_{eq}	Occ. (<1)
Cu1	0.59608 (2)	0.99743 (1)	0.70547 (2)	0.0458 (1)	
S1	0.78035 (4)	1.01647 (3)	0.56674 (3)	0.0539 (2)	
F1	0.8335 (6)	0.8733 (4)	0.6161 (4)	0.176 (4)	0.700
F2	0.9340 (5)	0.9174 (4)	0.5429 (4)	0.137 (2)	0.700
F3	0.9359 (6)	0.9687 (6)	0.6618 (6)	0.199 (4)	0.700
O1	0.63242 (11)	1.07473 (9)	0.79413 (10)	0.0626 (5)	
O2	0.51246 (12)	1.08054 (9)	0.64547 (10)	0.0608 (5)	
O3	0.73435 (13)	1.03226 (11)	0.64052 (11)	0.0712 (6)	
O4	0.8331 (2)	1.08491 (14)	0.53822 (17)	0.1123 (10)	

O5	0.7149 (2)	0.97542 (18)	0.50650 (17)	0.1224 (11)	
N1	0.53445 (12)	0.90229 (9)	0.63975 (10)	0.0449 (5)	
N2	0.68014 (12)	0.90588 (10)	0.76211 (9)	0.0472 (5)	
C1	0.46043 (16)	0.90316 (14)	0.57797 (12)	0.0556 (7)	
C2	0.41990 (18)	0.83081 (17)	0.54255 (14)	0.0640 (8)	
C3	0.45639 (18)	0.75590 (15)	0.56951 (15)	0.0633 (8)	
C4	0.53672 (15)	0.75244 (12)	0.63350 (13)	0.0509 (6)	
C5	0.57234 (13)	0.82808 (11)	0.66659 (11)	0.0423 (5)	
C6	0.65295 (13)	0.82997 (11)	0.73215 (11)	0.0430 (5)	
C7	0.58190 (18)	0.67773 (13)	0.66651 (17)	0.0649 (8)	
C8	0.65946 (18)	0.67936 (13)	0.72685 (16)	0.0642 (8)	
C9	0.69878 (15)	0.75606 (13)	0.76071 (13)	0.0526 (6)	
C10	0.78116 (18)	0.76280 (16)	0.82166 (15)	0.0664 (8)	
C11	0.81020 (18)	0.83952 (18)	0.85040 (15)	0.0701 (8)	
C12	0.75733 (17)	0.91011 (15)	0.82085 (13)	0.0605 (7)	
C13	0.57629 (16)	1.10508 (11)	0.84421 (13)	0.0514 (6)	
C14	0.6188 (2)	1.17618 (14)	0.89565 (17)	0.0706 (8)	
C15	0.8783 (3)	0.9414 (2)	0.5977 (2)	0.0953 (14)	
F3A	0.9596 (9)	0.9817 (7)	0.6337 (13)	0.153 (7)	0.300
F1A	0.8515 (8)	0.8809 (8)	0.6463 (7)	0.083 (3)	0.300
F2A	0.9048 (15)	0.9173 (13)	0.5189 (12)	0.193 (8)	0.300
H1	0.43520	0.95390	0.55800	0.0670*	
H2	0.36750	0.83360	0.50010	0.0770*	
H3	0.42880	0.70730	0.54610	0.0760*	
H7	0.55740	0.62700	0.64600	0.0780*	
H12	0.77660	0.96160	0.84290	0.0730*	
H14A	0.68970	1.18380	0.88760	0.1060*	
H14B	0.61240	1.16460	0.95250	0.1060*	
H14C	0.58140	1.22570	0.87990	0.1060*	
H8	0.68800	0.62970	0.74680	0.0770*	
H10	0.81550	0.71570	0.84220	0.0800*	
H11	0.86560	0.84480	0.89000	0.0840*	

Atomic displacement parameters (Å²)

	U^{11}	U^{22}	U^{33}	U^{12}	U^{13}	U^{23}
Cu1	0.0501 (2)	0.0364 (1)	0.0522 (2)	0.0032 (1)	0.0111 (1)	−0.0044 (1)
S1	0.0583 (3)	0.0503 (3)	0.0537 (3)	0.0031 (2)	0.0083 (2)	0.0036 (2)
F1	0.280 (8)	0.069 (3)	0.194 (7)	0.081 (4)	0.099 (6)	0.051 (4)
F2	0.115 (3)	0.157 (5)	0.152 (4)	0.072 (3)	0.077 (4)	0.039 (4)
F3	0.128 (6)	0.280 (9)	0.172 (5)	0.051 (5)	−0.073 (5)	0.031 (5)
O1	0.0620 (9)	0.0553 (8)	0.0732 (10)	−0.0075 (7)	0.0200 (7)	−0.0240 (7)
O2	0.0659 (9)	0.0482 (7)	0.0716 (9)	0.0163 (7)	0.0235 (7)	0.0118 (7)
O3	0.0704 (10)	0.0658 (10)	0.0820 (11)	0.0007 (8)	0.0305 (9)	−0.0070 (9)
O4	0.135 (2)	0.0787 (14)	0.1311 (19)	−0.0137 (13)	0.0530 (16)	0.0345 (13)
O5	0.127 (2)	0.131 (2)	0.0992 (18)	0.0116 (17)	−0.0404 (16)	−0.0369 (16)
N1	0.0464 (8)	0.0425 (8)	0.0464 (8)	0.0047 (6)	0.0076 (6)	−0.0021 (6)
N2	0.0475 (8)	0.0505 (9)	0.0444 (8)	0.0024 (7)	0.0084 (6)	−0.0017 (6)

C1	0.0570 (11)	0.0594 (12)	0.0500 (11)	0.0093 (9)	0.0027 (9)	−0.0019 (9)
C2	0.0589 (12)	0.0782 (16)	0.0539 (12)	−0.0017 (10)	−0.0005 (9)	−0.0116 (10)
C3	0.0647 (13)	0.0629 (13)	0.0634 (13)	−0.0117 (10)	0.0125 (10)	−0.0195 (10)
C4	0.0527 (10)	0.0442 (9)	0.0586 (11)	−0.0023 (8)	0.0194 (9)	−0.0061 (8)
C5	0.0433 (8)	0.0400 (8)	0.0458 (9)	0.0023 (7)	0.0154 (7)	−0.0022 (7)
C6	0.0423 (8)	0.0433 (9)	0.0456 (9)	0.0042 (7)	0.0154 (7)	0.0017 (7)
C7	0.0725 (14)	0.0384 (10)	0.0869 (16)	−0.0017 (9)	0.0240 (13)	−0.0038 (10)
C8	0.0719 (14)	0.0399 (10)	0.0845 (16)	0.0131 (9)	0.0271 (12)	0.0100 (10)
C9	0.0505 (10)	0.0547 (11)	0.0554 (11)	0.0112 (8)	0.0189 (9)	0.0101 (9)
C10	0.0628 (13)	0.0766 (15)	0.0608 (13)	0.0211 (11)	0.0104 (10)	0.0149 (11)
C11	0.0553 (12)	0.1009 (19)	0.0528 (12)	0.0110 (12)	−0.0016 (10)	0.0062 (12)
C12	0.0567 (11)	0.0728 (14)	0.0515 (11)	−0.0018 (10)	0.0028 (9)	−0.0072 (10)
C13	0.0623 (11)	0.0364 (8)	0.0575 (11)	−0.0067 (8)	0.0154 (9)	−0.0053 (8)
C14	0.0791 (15)	0.0531 (12)	0.0838 (16)	−0.0205 (11)	0.0291 (13)	−0.0256 (11)
C15	0.083 (2)	0.098 (2)	0.109 (3)	0.0357 (18)	0.0305 (19)	0.0181 (19)
F3A	0.046 (3)	0.082 (5)	0.32 (2)	0.000 (3)	−0.039 (7)	0.004 (8)
F1A	0.093 (4)	0.075 (5)	0.087 (5)	0.038 (3)	0.042 (3)	0.038 (4)
F2A	0.165 (14)	0.234 (16)	0.185 (14)	0.086 (10)	0.037 (10)	−0.120 (12)

Geometric parameters (Å, º)

Cu1—O1	1.9376 (16)	C3—C4	1.414 (3)
Cu1—O2	1.9385 (16)	C4—C5	1.397 (3)
Cu1—O3	2.2597 (18)	C4—C7	1.426 (3)
Cu1—N1	1.9995 (15)	C5—C6	1.433 (2)
Cu1—N2	2.0148 (16)	C6—C9	1.395 (3)
S1—O3	1.4237 (18)	C7—C8	1.349 (4)
S1—O4	1.406 (2)	C8—C9	1.431 (3)
S1—O5	1.409 (3)	C9—C10	1.403 (3)
S1—C15	1.803 (4)	C10—C11	1.365 (4)
F1—C15	1.295 (8)	C11—C12	1.395 (4)
F1A—C15	1.327 (13)	C13—C14	1.498 (3)
F2—C15	1.270 (7)	C1—H1	0.9302
F2A—C15	1.42 (2)	C2—H2	0.9304
F3—C15	1.309 (10)	C3—H3	0.9304
F3A—C15	1.337 (15)	C7—H7	0.9299
O1—C13	1.253 (3)	C8—H8	0.9300
O2—C13ⁱ	1.256 (3)	C10—H10	0.9295
N1—C1	1.333 (3)	C11—H11	0.9301
N1—C5	1.353 (2)	C12—H12	0.9307
N2—C12	1.329 (3)	C14—H14A	0.9598
N2—C6	1.354 (2)	C14—H14B	0.9605
C1—C2	1.386 (3)	C14—H14C	0.9596
C2—C3	1.358 (4)		
Cu1···O5	3.760 (3)	O4···H3ⁱᵛ	2.3363
Cu1···O1ⁱ	3.2474 (15)	O4···H11ᵛ	2.7477
Cu1···O2ⁱ	3.2315 (16)	O5···H14Bᵛ	2.7315

Cu1···N1[i]	3.5407 (16)	O5···H1[vi]	2.4275
Cu1···N2[i]	3.9955 (16)	N1···Cu1[i]	3.5407 (16)
Cu1···C1[i]	3.991 (2)	N2···F1A	3.103 (11)
Cu1···C5[i]	4.1970 (18)	N2···Cu1[i]	3.9955 (16)
Cu1···H8[ii]	3.5723	N2···C1[i]	3.344 (3)
F1···O3	2.920 (7)	C1···C12[i]	3.439 (3)
F1···O5	2.796 (7)	C1···O5[vi]	3.229 (3)
F1···C6	3.251 (8)	C1···N2[i]	3.344 (3)
F1···C14[iii]	3.249 (7)	C1···C13[i]	3.546 (3)
F1A···N2	3.103 (11)	C1···Cu1[i]	3.991 (2)
F1A···C6	3.184 (11)	C3···O4[vii]	3.213 (3)
F1A···C12	3.254 (11)	C3···C9[i]	3.599 (3)
F1A···O5	3.160 (12)	C5···C6[i]	3.525 (2)
F1A···O3	2.881 (12)	C5···Cu1[i]	4.1970 (18)
F2···O5	3.027 (7)	C5···C5[i]	3.472 (2)
F2···O4	3.006 (7)	C6···C5[i]	3.525 (2)
F2A···O4	2.89 (2)	C6···F1A	3.184 (11)
F2A···O5	2.65 (2)	C6···F1	3.251 (8)
F3···O4	2.983 (10)	C8···O1[iii]	3.256 (3)
F3···O3	2.826 (8)	C9···C3[i]	3.599 (3)
F3A···O4	2.733 (16)	C11···O4[viii]	3.293 (4)
F3A···O3	3.078 (12)	C12···C1[i]	3.439 (3)
F1···H14C[iii]	2.6275	C12···F1A	3.254 (11)
F1A···H14C[iii]	2.7013	C13···C13[i]	3.512 (3)
F3A···H7[iv]	2.6700	C13···C1[i]	3.546 (3)
O1···C8[ii]	3.256 (3)	C13···O5[viii]	3.335 (3)
O1···Cu1[i]	3.2474 (15)	C14···O5[viii]	3.228 (4)
O2···Cu1[i]	3.2315 (16)	C14···F1[ii]	3.249 (7)
O3···F3	2.826 (8)	C2···H14B[v]	3.0447
O3···F1	2.920 (7)	C8···H14A[iii]	2.8533
O3···F3A	3.078 (12)	C13···H1[i]	2.9284
O3···F1A	2.881 (12)	H1···C13[i]	2.9284
O4···F3	2.983 (10)	H1···O5[vi]	2.4275
O4···C3[iv]	3.213 (3)	H1···O2	2.6401
O4···F2	3.006 (7)	H3···H7	2.5822
O4···C11[v]	3.293 (4)	H3···O4[vii]	2.3363
O4···F2A	2.89 (2)	H7···F3A[vii]	2.6700
O4···F3A	2.733 (16)	H7···H3	2.5822
O5···Cu1	3.760 (3)	H8···H10	2.5804
O5···F2A	2.65 (2)	H8···Cu1[iii]	3.5723
O5···C13[v]	3.335 (3)	H8···O1[iii]	2.6624
O5···C14[v]	3.228 (4)	H8···O3[iii]	2.5589
O5···F2	3.027 (7)	H10···H8	2.5804
O5···F1A	3.160 (12)	H11···O4[viii]	2.7477
O5···F1	2.796 (7)	H12···O1	2.6934
O5···C1[vi]	3.229 (3)	H14A···C8[ii]	2.8533
O1···H12	2.6934	H14B···C2[viii]	3.0447
O1···H8[ii]	2.6624	H14B···O5[viii]	2.7315

O2···H1	2.6401	H14C···F1ⁱⁱ	2.6275
O3···H8ⁱⁱ	2.5589	H14C···F1Aⁱⁱ	2.7013
O1—Cu1—O2	91.21 (6)	C8—C9—C10	124.5 (2)
O1—Cu1—O3	92.37 (6)	C9—C10—C11	119.1 (2)
O1—Cu1—N1	162.62 (7)	C10—C11—C12	120.5 (2)
O1—Cu1—N2	92.44 (6)	N2—C12—C11	121.8 (2)
O2—Cu1—O3	91.76 (7)	O1—C13—C14	117.14 (19)
O2—Cu1—N1	94.48 (6)	O1—C13—O2ⁱ	125.13 (18)
O2—Cu1—N2	176.35 (6)	O2ⁱ—C13—C14	117.73 (19)
O3—Cu1—N1	103.84 (7)	S1—C15—F1	107.9 (4)
O3—Cu1—N2	88.28 (6)	S1—C15—F2	116.7 (4)
N1—Cu1—N2	81.97 (6)	S1—C15—F3	109.9 (5)
O3—S1—O4	113.79 (13)	S1—C15—F1A	116.1 (5)
O3—S1—O5	113.45 (13)	S1—C15—F2A	99.1 (8)
O3—S1—C15	103.33 (13)	S1—C15—F3A	108.3 (5)
O4—S1—O5	115.02 (16)	F1—C15—F2	102.1 (5)
O4—S1—C15	105.09 (16)	F1—C15—F3	109.8 (6)
O5—S1—C15	104.55 (16)	F2—C15—F3	110.0 (6)
Cu1—O1—C13	128.57 (14)	F1A—C15—F2A	116.4 (10)
Cu1—O2—C13ⁱ	129.53 (14)	F1A—C15—F3A	109.8 (9)
Cu1—O3—S1	141.19 (11)	F2A—C15—F3A	106.2 (12)
Cu1—N1—C1	128.86 (14)	N1—C1—H1	118.98
Cu1—N1—C5	112.78 (12)	C2—C1—H1	118.97
C1—N1—C5	118.28 (16)	C1—C2—H2	119.86
Cu1—N2—C6	112.49 (12)	C3—C2—H2	119.87
Cu1—N2—C12	129.82 (15)	C2—C3—H3	120.32
C6—N2—C12	117.66 (17)	C4—C3—H3	120.31
N1—C1—C2	122.0 (2)	C4—C7—H7	119.34
C1—C2—C3	120.3 (2)	C8—C7—H7	119.47
C2—C3—C4	119.4 (2)	C7—C8—H8	119.37
C3—C4—C5	116.78 (18)	C9—C8—H8	119.37
C3—C4—C7	124.6 (2)	C9—C10—H10	120.49
C5—C4—C7	118.66 (19)	C11—C10—H10	120.44
N1—C5—C4	123.22 (17)	C10—C11—H11	119.78
N1—C5—C6	116.49 (16)	C12—C11—H11	119.74
C4—C5—C6	120.29 (17)	N2—C12—H12	119.07
N2—C6—C5	116.13 (16)	C11—C12—H12	119.09
N2—C6—C9	124.13 (17)	C13—C14—H14A	109.53
C5—C6—C9	119.74 (17)	C13—C14—H14B	109.45
C4—C7—C8	121.2 (2)	C13—C14—H14C	109.45
C7—C8—C9	121.3 (2)	H14A—C14—H14B	109.50
C6—C9—C8	118.75 (19)	H14A—C14—H14C	109.50
C6—C9—C10	116.73 (19)	H14B—C14—H14C	109.39
O2—Cu1—O1—C13	68.10 (18)	Cu1—N1—C1—C2	−174.67 (16)
O3—Cu1—O1—C13	159.91 (17)	C5—N1—C1—C2	1.8 (3)
N2—Cu1—O1—C13	−111.71 (17)	Cu1—N1—C5—C4	176.12 (15)

O1—Cu1—O2—C13[i]	−78.62 (18)	Cu1—N1—C5—C6	−4.0 (2)
O3—Cu1—O2—C13[i]	−171.03 (18)	C1—N1—C5—C4	−0.9 (3)
N1—Cu1—O2—C13[i]	84.94 (18)	C1—N1—C5—C6	178.99 (17)
O1—Cu1—O3—S1	−175.37 (18)	Cu1—N2—C6—C5	0.7 (2)
O2—Cu1—O3—S1	−84.09 (18)	Cu1—N2—C6—C9	−179.80 (15)
N1—Cu1—O3—S1	10.96 (19)	C12—N2—C6—C5	−177.49 (17)
N2—Cu1—O3—S1	92.26 (18)	C12—N2—C6—C9	2.0 (3)
O2—Cu1—N1—C1	−0.85 (18)	Cu1—N2—C12—C11	−176.94 (16)
O2—Cu1—N1—C5	−177.49 (13)	C6—N2—C12—C11	0.9 (3)
O3—Cu1—N1—C1	−93.78 (17)	N1—C1—C2—C3	−1.1 (3)
O3—Cu1—N1—C5	89.58 (13)	C1—C2—C3—C4	−0.6 (3)
N2—Cu1—N1—C1	−180.00 (18)	C2—C3—C4—C5	1.4 (3)
N2—Cu1—N1—C5	3.37 (13)	C2—C3—C4—C7	−179.5 (2)
O1—Cu1—N2—C6	161.25 (13)	C3—C4—C7—C8	179.0 (2)
O1—Cu1—N2—C12	−20.80 (18)	C5—C4—C7—C8	−1.9 (3)
O3—Cu1—N2—C6	−106.45 (13)	C7—C4—C5—C6	0.2 (3)
O3—Cu1—N2—C12	71.50 (18)	C3—C4—C5—N1	−0.7 (3)
N1—Cu1—N2—C6	−2.21 (12)	C3—C4—C5—C6	179.43 (18)
N1—Cu1—N2—C12	175.74 (18)	C7—C4—C5—N1	−179.86 (19)
O4—S1—O3—Cu1	146.81 (18)	C4—C5—C6—C9	2.6 (3)
O5—S1—O3—Cu1	12.8 (2)	N1—C5—C6—N2	2.2 (2)
C15—S1—O3—Cu1	−99.8 (2)	N1—C5—C6—C9	−177.30 (17)
O3—S1—C15—F1	67.4 (4)	C4—C5—C6—N2	−177.90 (17)
O3—S1—C15—F2	−178.4 (4)	N2—C6—C9—C10	−3.1 (3)
O3—S1—C15—F3	−52.3 (5)	C5—C6—C9—C8	−3.8 (3)
O4—S1—C15—F1	−173.0 (4)	C5—C6—C9—C10	176.37 (18)
O4—S1—C15—F2	−58.8 (4)	N2—C6—C9—C8	176.76 (19)
O4—S1—C15—F3	67.2 (5)	C4—C7—C8—C9	0.6 (4)
O5—S1—C15—F1	−51.5 (4)	C7—C8—C9—C6	2.2 (3)
O5—S1—C15—F2	62.7 (4)	C7—C8—C9—C10	−178.0 (2)
O5—S1—C15—F3	−171.3 (5)	C8—C9—C10—C11	−178.5 (2)
Cu1—O1—C13—C14	−167.96 (15)	C6—C9—C10—C11	1.3 (3)
Cu1—O1—C13—O2[i]	12.1 (3)	C9—C10—C11—C12	1.4 (4)
Cu1—O2—C13[i]—O1[i]	−5.3 (3)	C10—C11—C12—N2	−2.6 (4)
Cu1—O2—C13[i]—C14[i]	174.68 (15)		

Symmetry codes: (i) −*x*+1, *y*, −*z*+3/2; (ii) −*x*+3/2, *y*+1/2, −*z*+3/2; (iii) −*x*+3/2, *y*−1/2, −*z*+3/2; (iv) *x*+1/2, *y*+1/2, *z*; (v) *x*, −*y*+2, *z*−1/2; (vi) −*x*+1, −*y*+2, −*z*+1; (vii) *x*−1/2, *y*−1/2, *z*; (viii) *x*, −*y*+2, *z*+1/2.

Hydrogen-bond geometry (Å, °)

D—H···*A*	*D*—H	H···*A*	*D*···*A*	*D*—H···*A*
C1—H1···O5[vi]	0.93	2.43	3.229 (3)	144
C3—H3···O4[vii]	0.93	2.34	3.213 (3)	157
C8—H8···O3[iii]	0.93	2.56	3.421 (3)	154

Symmetry codes: (iii) −*x*+3/2, *y*−1/2, −*z*+3/2; (vi) −*x*+1, −*y*+2, −*z*+1; (vii) *x*−1/2, *y*−1/2, *z*.