



Acta Crystallographica Section E

Structure Reports

Online

ISSN 1600-5368

[Cu₂(HF₂)(H₂O)₈][FeF₆]·2H₂O

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Received 19 November 2008; accepted 24 February 2009

Key indicators: single-crystal X-ray study; $T = 293$ K; mean $\sigma(\text{Cu}-\text{O}) = 0.003$ Å; R factor = 0.040; wR factor = 0.080; data-to-parameter ratio = 24.8.

The title compound, octaaqua(hydrogenfluorido)dicopper(II) hexafluoridoferrate(III) dihydrate, was synthesized under hydrothermal conditions. The Cu atom is coordinated by one F and five O atoms within a highly distorted octahedron, forming dimeric $[\text{Cu}_2(\text{H}_2\text{O})_8\text{HF}_2]^{3+}$ units by edge sharing. These units are hydrogen bonded to $[\text{FeF}_6]^{3-}$ anions and to an interstitial water molecule. The former feature Fe^{3+} on a special position ($\bar{1}$). The dimeric copper units are linked to adjacent dimers by $\text{F}-\text{H}\cdots\text{F}$ hydrogen bonds. Additional $\text{O}-\text{H}\cdots\text{O}$ and $\text{O}-\text{H}\cdots\text{F}$ hydrogen bonds help to consolidate the crystal packing.

Related literature

For other hydrated copper-iron fluorides, see: Kummer & Babel (1987); Leblanc & Ferey (1990). For $\text{F}\cdots\text{F}$ distances, see: Frevel & Rinn (1962); Massa & Herdtweck (1983). For asymmetrical $\text{F}-\text{H}\cdots\text{F}$ hydrogen bonding, see: Roesky *et al.* (1990); Gerasimenko *et al.* (2007); Gerken *et al.* (2002). For $\text{Pb}_8\text{MnFe}_2\text{F}_{24}$, see: Le Bail & Mercier (1992). For valence-bond analysis, see: Brown & Altermatt (1985); Brese & O'Keeffe (1991).

Experimental

Crystal data

 $[\text{Cu}_2(\text{HF}_2)(\text{H}_2\text{O})_8][\text{FeF}_6]\cdot 2\text{H}_2\text{O}$ $M_r = 516.12$ Triclinic, $P\bar{1}$ $a = 6.659$ (2) Å $b = 7.450$ (3) Å $c = 8.377$ (5) Å $\alpha = 107.37$ (4)° $\beta = 106.89$ (5)° $\gamma = 94.26$ (3)° $V = 373.6$ (3) Å³ $Z = 1$ Mo $K\alpha$ radiation $\mu = 3.91$ mm⁻¹ $T = 293$ K $0.11 \times 0.10 \times 0.05$ mm

Data collection

Siemens AED2 diffractometer

Absorption correction: gaussian

(SHELX76; Sheldrick, 2008)

 $T_{\min} = 0.698$, $T_{\max} = 0.845$

3303 measured reflections

3303 independent reflections

2044 reflections with $I > 2\sigma(I)$

3 standard reflections

frequency: 120 min

intensity decay: 15%

Refinement

 $R[F^2 > 2\sigma(F^2)] = 0.040$ $wR(F^2) = 0.080$ $S = 1.00$

3303 reflections

133 parameters

15 restraints

H atoms treated by a mixture of independent and constrained refinement

 $\Delta\rho_{\max} = 0.62$ e Å⁻³ $\Delta\rho_{\min} = -0.63$ e Å⁻³

Table 1

Selected bond lengths (Å).

Cu—F4	1.9058 (19)	Fe—F1	1.9204 (16)
Cu—O1	1.939 (2)	Fe—F2 ⁱ	1.930 (2)
Cu—O2	1.958 (2)	Fe—F3	1.9402 (17)
Cu—O3	1.977 (2)	Cu—O3 ⁱⁱ	2.715 (3)
Cu—O4	2.349 (3)	Cu···Cu ⁱⁱ	3.575 (3)

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

Table 2

Hydrogen-bond geometry (Å, °).

$D-H\cdots A$	$D-H$	$H\cdots A$	$D\cdots A$	$D-H\cdots A$
O1—H11···F1 ⁱⁱⁱ	1.00 (4)	1.59 (4)	2.590 (3)	173 (4)
O1—H12···F2	0.99 (4)	1.71 (4)	2.679 (3)	167 (4)
O2—H21···F1 ⁱⁱ	1.02 (5)	1.61 (5)	2.595 (3)	161 (4)
O2—H22···O5 ^{iv}	0.98 (4)	1.82 (4)	2.699 (3)	147 (5)
O3—H31···F3 ⁱⁱ	1.07 (5)	1.52 (5)	2.577 (2)	167 (5)
O3—H32···F3 ^v	1.04 (5)	1.64 (5)	2.668 (3)	171 (5)
O4—H41···F2 ^{vi}	1.02 (4)	2.07 (9)	2.766 (3)	123 (8)
O4—H42···O5 ^v	1.01 (9)	1.81 (9)	2.793 (3)	164 (8)
O5—H51···F2 ^{vii}	1.01 (6)	2.05 (7)	2.888 (3)	139 (5)
O5—H51···F3 ^{viii}	1.01 (6)	2.23 (5)	3.072 (4)	140 (6)
O5—H52···F4	1.05 (7)	1.63 (7)	2.676 (3)	175 (6)
F4—H6···F4 ^{vi}	0.91	1.82	2.597 (4)	142

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

Table 3

Valence-bond analysis according to the empirical expression from Brown & Altermatt (1985), using parameters for solids from Brese & O'Keeffe (1991).

	O1	O2	O3	O4	F4	F1	F2	F3	O5	Σ	Σ_{expected}
Cu	0.49	0.47	0.45	0.16	0.44					2.01	2
Fe						0.51 ($\times 2$)	0.49 ($\times 2$)	0.48 ($\times 2$)		2.96	3
H11	0.80					0.20				1	1
H12	0.80						0.20			1	1
H21		0.80				0.20				1	1
H22		0.80							0.20	1	1
H31			0.80					0.20		1	1
H41				0.80			0.20			1	1
H42				0.80					0.20	1	1
H51							0.10	0.10		0.80	1
H52					0.20					0.80	1
H6						0.80 ($\times 0.5$)				1	1
					0.20 ($\times 0.5$)						
Σ	2.09	2.07	2.05	1.76	1.14	0.91	0.99	0.98		2.00	
Σ_{expected}	2	2	2	2	1	1	1	1		2	

Data collection: *STADIA* (Stoe & Cie, 1998); cell refinement: *STADIA*; data reduction: *X-RED* (Stoe & Cie, 1998); program(s) used to solve structure: *SHELXS97* (Sheldrick, 2008); program(s) used to refine structure: *SHELXL97* (Sheldrick, 2008); molecular graphics: *DIAMOND* (Brandenburg, 1999) and *ORTEP-3* (Farrugia,

1997); software used to prepare material for publication: *publCIF* (Westrip, 2009).

Thanks are due to M. Leblanc for the single-crystal data collection.

Supplementary data and figures for this paper are available from the IUCr electronic archives (Reference: FI2069).

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supporting information

Acta Cryst. (2009). E65, i23–i24 [doi:10.1107/S1600536809006874]

[Cu₂(HF₂)(H₂O)₈][FeF₆]·2H₂O

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S1. Comment

[Cu(H₂O)₄H_{0.5}F]₂(FeF₆)(H₂O)₂ is the third hydrated copper–iron fluoride, the two previous ones being Cu₃Fe₂F₁₂(H₂O)₁₂ (Kummer & Babel, 1987) and CuFe₂F₈(H₂O)₂ (Leblanc & Ferey, 1990). In this new compound, an almost perfect (FeF₆)³⁻ octahedron, placed on an inversion center, is connected exclusively by hydrogen bonding to edge-sharing bi-octahedral [Cu₂(H₂O)₈HF₂]³⁺ units and to an interstitial water molecule. The copper atom can be considered to be five-coordinated by four water molecules and one F atom in the fashion of a square pyramid (Figure 1). The square is constituted by the F atom and three water molecules at distances in the 1.906–1.977 Å range (Table 1), whereas Ow(4), at the top of the pyramid is at 2.349 Å (Jahn-Teller distortion). However, if one considers the next neighbour Ow(3) at 2.715 Å as sixth ligand, the coordination can be described as very distorted octahedral, yielding centrosymmetric, dimeric units with a Cu—Cu distance of 3.575 Å. The dimers are bonded to adjacent dimers through F(4) by a HF₂⁻ ion (figure 2), however the F—F distance (2.597 Å) is much larger (Table 2) than usually observed (2.27 Å in LiHF₂ (Frevel & Rinn, 1962), 2.28 in BaHF₃ (Massa & Herdtweck, 1983). Moreover, the hydrogen atom H(6) which would have been expected at the 1/2, 1/2, 1/2 special position, exactly at the middle of the F(4)—F(4) atoms, in order to form a linear (F—H—F)⁻ ion, was seen on the Fourier difference map as occupying more probably half a general position. It is then more a F—H···F bridge than a F—H—F one. Such asymmetrical F—H···F hydrogen bonding was observed in many cases for F—F distances going up to 2.686 Å in [(η⁵-C₅Me₅)NbF₄(HF)AsF₃]₂ (Roesky *et al.*, 1990), 2.429 to 2.512 Å in [OsO₃F](HF)₂[AsF₆] (Gerken *et al.*, 2002), 2.326 to 2.402 Å in Rb_{2-x}K_xZrF₆(HF)₂ (Gerasimenko *et al.*, 2007).

The charge of the cations is balanced by the centrosymmetric anion [FeF₆]³⁻. There are 14 water molecules around the (FeF₆)³⁻ octahedron, the H(51) atom corresponds to a bifurcated hydrogen bond towards F(2) and F(3) (figure 3). This helps the bond valence calculations to provide relatively satisfying results (Table 3), the largest disagreements being on O(4) and F(4). The contribution from the long Cu—Ow(3) distance is negligible.

Comparing with the Cu₃Fe₂F₁₂(H₂O)₁₂ crystal structure (Kummer & Babel, 1987), also triclinic, there are strong differences. The three copper atoms and two Fe atoms are all placed on inversion centers. One (FeF₆)³⁻ octahedron is isolated, and the other forms chiolite-like square meshes by sharing F corners with tetra-hydrated Cu(H₂O)₄F₂ elongated octahedra (the long distances are the two Cu—F ones, close to 2.32 Å for the three independent copper sites). In CuFe₂F₈(H₂O)₂ (Leblanc & Ferey, 1990), the CuF₄(H₂O)₂ octahedra show two long Cu—F distances (2.451 Å). So, in both cases, the long distances are Cu—F ones, which is not the case of the title compound.

A study of the magnetic properties is currently in progress.

S2. Experimental

Hydrothermal growth at 493 K from (2PbF₂/2CuF₂/FeF₃) in HF 5M or 1M solutions, produced large crystals which could be identified as corresponding to Cu₃Fe₂F₁₂(H₂O)₁₂ (Kummer & Babel, 1987) (5M solution) or to the title compound (1M). Both compounds are occurring with the same intense blue color and could have been confused if no powder pattern

had been recorded. At other starting compositions, mixtures of both compounds could be observed, and also together especially with $\text{Pb}_8\text{CuFe}_2\text{F}_{24}$ (starting from $2\text{PbF}_2/\text{CuF}_2/2\text{FeF}_3$ for instance), isostructural with $\text{Pb}_8\text{MnFe}_2\text{F}_{24}$ (Le Bail & Mercier, 1992).

S3. Refinement

The hydrogen atoms were all located on the difference Fourier map. Those of the water molecules were restrained to have Ow—H and H—H distances respectively close to 1.0 and 1.59 Å, their U_{iso} values were refined by groups of two. The hydrogen atom of the HF group was let at its difference Fourier map position with a fixed U_{iso} .

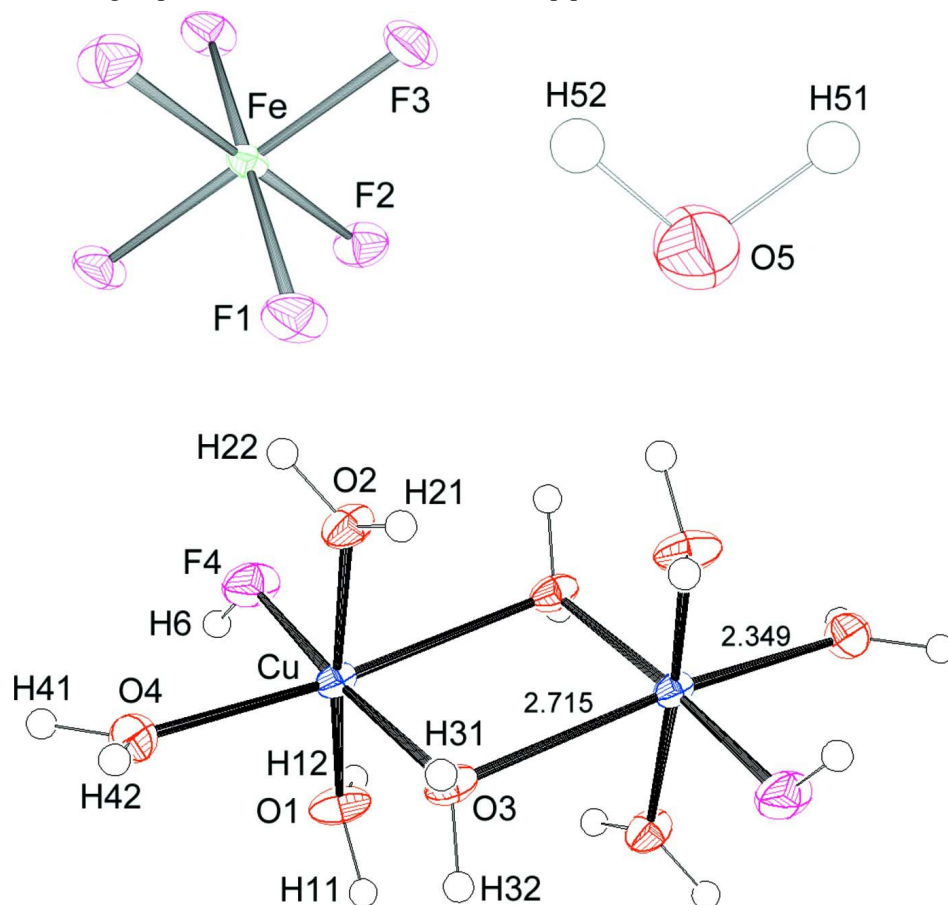
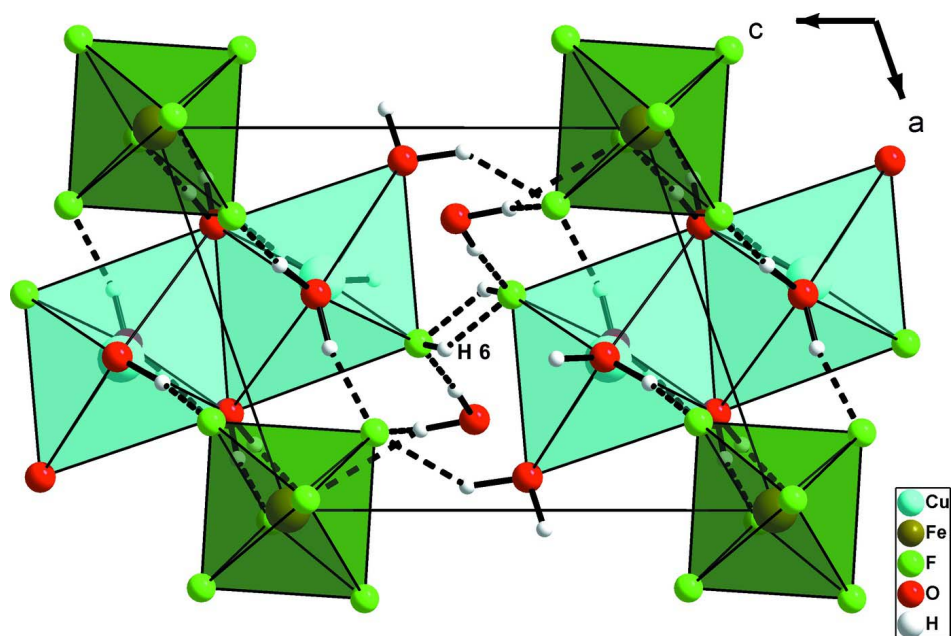


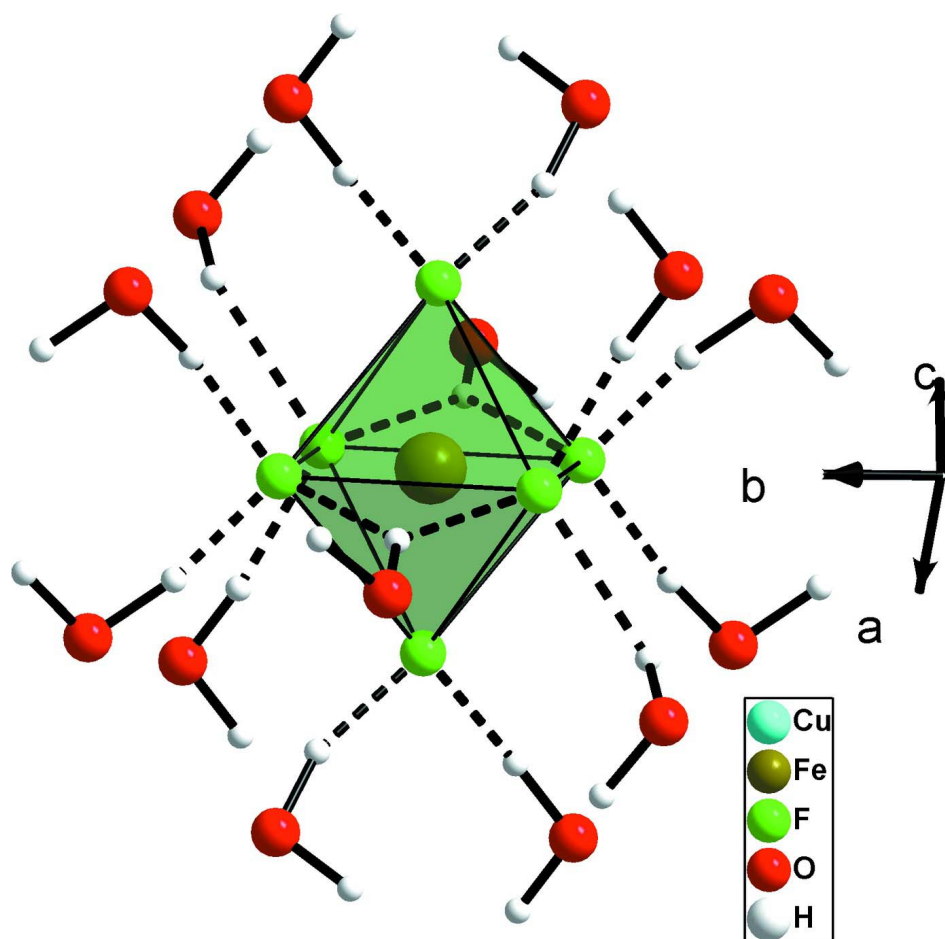
Figure 1

ORTEP-3 view (Farrugia, 1997) of the FeF_6^{3-} octahedron, the isolated water molecule and the $\text{Cu}(\text{H}_2\text{O})_4\text{F}^+$ square-based pyramid which is forming octahedra dimeric units when considering the very long Cu—O3 distance (2.715 Å).

Displacement ellipsoids are shown at the 50% probability level. Symmetry-equivalent F atoms in the FeF_6 group are generated by (i) $-x, -y, -z$. Symmetry-equivalent atoms in the $[\text{Cu}_2(\text{H}_2\text{O})_8\text{HF}_2]^{3+}$ unit are generated by (iii) $1-x, 1-y, -z$.

**Figure 2**

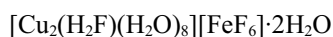
Crystal packing with view along [010]. Hydrogen bonding is shown as dashed lines. Copper coordination shown as distorted octahedra (adding O(3) at 2.715 Å from Cu) sharing an edge.

**Figure 3**

The 14 water molecules, from the Cu coordination sphere or isolated, involved in hydrogen bonding with the FeF_6^{3-} octahedron.

Octaaqua(hydrogenfluorido)dicopper(II) hexafluoridoferrate(III) dihydrate

Crystal data



$M_r = 516.12$

Triclinic, $P\bar{1}$

Hall symbol: $-\text{P } 1$

$a = 6.659 (2) \text{ \AA}$

$b = 7.450 (3) \text{ \AA}$

$c = 8.377 (5) \text{ \AA}$

$\alpha = 107.37 (4)^\circ$

$\beta = 106.89 (5)^\circ$

$\gamma = 94.26 (3)^\circ$

$V = 373.6 (3) \text{ \AA}^3$

$Z = 1$

$F(000) = 257$

$D_x = 2.294 \text{ Mg m}^{-3}$

Mo $K\alpha$ radiation, $\lambda = 0.71069 \text{ \AA}$

Cell parameters from 28 reflections

$\theta = 2.7\text{--}35^\circ$

$\mu = 3.91 \text{ mm}^{-1}$

$T = 293 \text{ K}$

Platelet, blue

$0.11 \times 0.10 \times 0.05 \text{ mm}$

Data collection

Siemens AED2
diffractometer

Radiation source: fine-focus sealed tube

Graphite monochromator
 $2\theta/\omega$ scans

Absorption correction: gaussian
(*SHELX76*; Sheldrick, 2008)
 $T_{\min} = 0.698$, $T_{\max} = 0.845$
3303 measured reflections
3303 independent reflections
2044 reflections with $I > 2\sigma(I)$
 $R_{\text{int}} = 0.000$

$\theta_{\max} = 35.0^\circ$, $\theta_{\min} = 2.7^\circ$
 $h = -10 \rightarrow 10$
 $k = -12 \rightarrow 11$
 $l = 0 \rightarrow 13$
3 standard reflections every 120 min
intensity decay: 15%

Refinement

Refinement on F^2
Least-squares matrix: full
 $R[F^2 > 2\sigma(F^2)] = 0.040$
 $wR(F^2) = 0.080$
 $S = 1.00$
3303 reflections
133 parameters
15 restraints
Primary atom site location: structure-invariant
direct methods
Secondary atom site location: difference Fourier
map

Hydrogen site location: difference Fourier map
H atoms treated by a mixture of independent
and constrained refinement
 $w = 1/[\sigma^2(F_o^2) + (0.03P)^2]$
where $P = (F_o^2 + 2F_c^2)/3$
 $(\Delta/\sigma)_{\max} = 0.004$
 $\Delta\rho_{\max} = 0.62 \text{ e } \text{\AA}^{-3}$
 $\Delta\rho_{\min} = -0.63 \text{ e } \text{\AA}^{-3}$
Extinction correction: *SHELXL97* (Sheldrick,
2008), $F_c^* = kFc[1 + 0.001xFe^2\lambda^3/\sin(2\theta)]^{-1/4}$
Extinction coefficient: 0.0072 (17)

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>	$U_{\text{iso}}^*/U_{\text{eq}}$	Occ. (<1)
Cu	0.60473 (5)	0.55727 (4)	0.23479 (4)	0.01662 (8)	
Fe	0.0000	0.0000	0.0000	0.01433 (10)	
F1	0.2310 (3)	-0.0246 (2)	-0.0944 (2)	0.0270 (3)	
F2	0.2029 (3)	0.1318 (2)	0.2331 (2)	0.0275 (3)	
F3	-0.0278 (3)	0.2400 (2)	-0.0437 (2)	0.0272 (4)	
F4	0.4365 (3)	0.5885 (3)	0.3872 (2)	0.0338 (4)	
O1	0.5648 (3)	0.2890 (3)	0.2113 (3)	0.0283 (4)	
O2	0.5992 (3)	0.8217 (3)	0.2411 (3)	0.0244 (4)	
O3	0.7470 (3)	0.5094 (3)	0.0530 (3)	0.0219 (4)	
O4	0.9139 (3)	0.6601 (3)	0.4849 (3)	0.0262 (4)	
O5	0.2376 (4)	0.8884 (3)	0.4476 (3)	0.0291 (4)	
H11	0.641 (6)	0.188 (5)	0.158 (6)	0.064 (10)*	
H12	0.424 (5)	0.226 (6)	0.201 (6)	0.064 (10)*	
H21	0.667 (7)	0.876 (7)	0.167 (5)	0.084 (12)*	
H22	0.604 (8)	0.935 (5)	0.341 (5)	0.084 (12)*	
H31	0.866 (7)	0.623 (6)	0.067 (8)	0.117 (16)*	
H32	0.829 (8)	0.398 (6)	0.022 (8)	0.117 (16)*	

H41	0.937 (15)	0.679 (15)	0.615 (5)	0.24 (3)*	
H42	1.049 (10)	0.734 (13)	0.487 (12)	0.24 (3)*	
H51	0.211 (11)	0.909 (10)	0.330 (6)	0.16 (2)*	
H52	0.315 (12)	0.770 (8)	0.430 (10)	0.16 (2)*	
H6	0.4219	0.4979	0.4375	0.080*	0.50

Atomic displacement parameters (Å²)

	U^{11}	U^{22}	U^{33}	U^{12}	U^{13}	U^{23}
Cu	0.02080 (15)	0.01342 (14)	0.01901 (15)	0.00339 (11)	0.01121 (12)	0.00561 (11)
Fe	0.0157 (2)	0.0129 (2)	0.0174 (2)	0.00267 (17)	0.00901 (19)	0.00591 (18)
F1	0.0276 (8)	0.0268 (8)	0.0391 (9)	0.0100 (6)	0.0248 (7)	0.0139 (7)
F2	0.0255 (8)	0.0296 (8)	0.0217 (8)	−0.0025 (6)	0.0064 (6)	0.0038 (6)
F3	0.0295 (8)	0.0181 (7)	0.0442 (10)	0.0077 (6)	0.0185 (8)	0.0176 (7)
F4	0.0475 (11)	0.0338 (9)	0.0375 (10)	0.0184 (8)	0.0290 (9)	0.0192 (8)
O1	0.0285 (10)	0.0151 (8)	0.0470 (13)	0.0038 (7)	0.0227 (9)	0.0086 (8)
O2	0.0386 (11)	0.0138 (8)	0.0261 (10)	0.0040 (7)	0.0177 (8)	0.0074 (7)
O3	0.0270 (9)	0.0179 (8)	0.0281 (10)	0.0056 (7)	0.0191 (8)	0.0079 (7)
O4	0.0243 (9)	0.0267 (10)	0.0261 (10)	0.0042 (8)	0.0076 (8)	0.0074 (8)
O5	0.0359 (11)	0.0279 (10)	0.0264 (10)	0.0109 (9)	0.0128 (9)	0.0093 (8)

Geometric parameters (Å, °)

Cu—F4	1.9058 (19)	F2—O5 ^v	2.888 (3)
Cu—O1	1.939 (2)	F3—O3 ⁱⁱⁱ	2.577 (2)
Cu—O2	1.958 (2)	F3—O3 ^{vi}	2.668 (3)
Cu—O3	1.977 (2)	F3—O5 ^{vii}	3.072 (4)
Cu—O4	2.349 (3)	F4—F4 ^{iv}	2.597 (4)
Fe—F1	1.9204 (16)	F4—O1	2.632 (3)
Fe—F2 ⁱ	1.930 (2)	F4—O5	2.676 (3)
Fe—F3	1.9402 (17)	F4—O2	2.730 (3)
F1—O1 ⁱⁱ	2.590 (3)	F4—O4	3.006 (3)
F1—O2 ⁱⁱⁱ	2.595 (3)	O1—O3	2.804 (3)
F1—F2	2.709 (3)	O2—O5 ^{viii}	2.699 (3)
F1—F3 ⁱ	2.728 (2)	O2—O3	2.814 (3)
F1—F3	2.732 (2)	O2—O4	3.061 (3)
F1—F2 ⁱ	2.736 (3)	O2—O3 ⁱⁱⁱ	3.113 (4)
F1—O1	3.051 (4)	O3—O3 ⁱⁱⁱ	3.128 (4)
F2—O1	2.679 (3)	O4—O4 ^{ix}	2.768 (4)
F2—F3 ⁱ	2.714 (3)	O4—O5 ^x	2.793 (3)
F2—F3	2.759 (3)	Cu—O3 ⁱⁱⁱ	2.715 (3)
F2—O4 ^{iv}	2.766 (3)	Cu—Cu ⁱⁱⁱ	3.575 (3)
F4—Cu—O1	86.42 (9)	F1 ⁱ —Fe—F2 ⁱ	89.43 (9)
F4—Cu—O2	89.89 (8)	F1—Fe—F3	90.08 (7)
O1—Cu—O2	171.49 (9)	F1 ⁱ —Fe—F3	89.92 (7)
F4—Cu—O3	173.09 (9)	F2 ⁱ —Fe—F3	89.06 (9)
O1—Cu—O3	91.47 (9)	F2—Fe—F3	90.94 (9)

O2—Cu—O3	91.30 (8)	H11—O1—H12	109 (3)
F4—Cu—O4	89.27 (9)	H21—O2—H22	104 (3)
O1—Cu—O4	97.50 (10)	H31—O3—H32	98 (3)
O2—Cu—O4	90.11 (9)	H41—O4—H42	103 (4)
O3—Cu—O4	97.53 (9)	H51—O5—H52	102 (3)
F1—Fe—F2 ⁱ	90.57 (9)		

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

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

$D-H\cdots A$	$D-H$	$H\cdots A$	$D\cdots A$	$D-H\cdots A$
O1—H11 \cdots F1 ⁱⁱ	1.00 (4)	1.59 (4)	2.590 (3)	173 (4)
O1—H12 \cdots F2	0.99 (4)	1.71 (4)	2.679 (3)	167 (4)
O2—H21 \cdots F1 ⁱⁱⁱ	1.02 (5)	1.61 (5)	2.595 (3)	161 (4)
O2—H22 \cdots O5 ^{viii}	0.98 (4)	1.82 (4)	2.699 (3)	147 (5)
O3—H31 \cdots F3 ⁱⁱⁱ	1.07 (5)	1.52 (5)	2.577 (2)	167 (5)
O3—H32 \cdots F3 ^x	1.04 (5)	1.64 (5)	2.668 (3)	171 (5)
O4—H41 \cdots F2 ^{iv}	1.02 (4)	2.07 (9)	2.766 (3)	123 (8)
O4—H42 \cdots O5 ^x	1.01 (9)	1.81 (9)	2.793 (3)	164 (8)
O5—H51 \cdots F2 ^{xi}	1.01 (6)	2.05 (7)	2.888 (3)	139 (5)
O5—H51 \cdots F3 ^{vii}	1.01 (6)	2.23 (5)	3.072 (4)	140 (6)
O5—H52 \cdots F4	1.05 (7)	1.63 (7)	2.676 (3)	175 (6)
F4—H6 \cdots F4 ^{iv}	0.91	1.82	2.597 (4)	142

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