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A second polymorph of aqua{4-chloro-2-[(pyridin-2-ylmethyl)iminomethyl]-phenolato}copper(II) nitrate monohydrate

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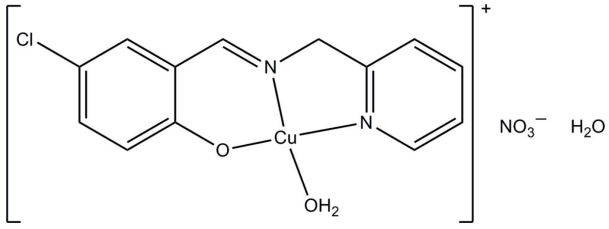
Key indicators: single-crystal X-ray study; T = 298 K; mean σ(C–C) = 0.004 Å; R factor = 0.037; wR factor = 0.089; data-to-parameter ratio = 14.6.

The title complex, $[Cu(C_{13}H_{10}ClN_2O)(H_2O)]NO_3\cdot H_2O$, was obtained by the reaction of 5-chlorosalicylaldehyde, 2-(aminomethyl)pyridine and copper nitrate in methanol. The first reported polymorph of this complex was triclinic [Liang *et al.* (2010). *Acta Cryst.* E**66**, m40]. The present polymorph crystallized in the monoclinic space group $P2_1/c$. The Cu^{II} ion is in a square planar environment and is coordinated by one phenolate O, one imine N and one pyridine N atom of the tridentate Schiff base ligand and by one water O atom. In the crystal, molecules are linked through intermolecular O—H···O hydrogen bonds to form chains along the *a* axis.

Related literature

For the structures and properties of Schiff base copper(II) complexes, see: Patel *et al.* (2011); Creaven *et al.* (2010); Osowole *et al.* (2008). For the complex with triclinic space group $P\overline{1}$, see: Liang *et al.* (2010).



Experimental

Crystal data

$[Cu(C_{13}H_{10}ClN_2O)(H_2O)]NO_3\cdot H_2O$
M_r = 407.26
Monoclinic, $P2_1/c$
a = 7.840 (2) Å
b = 8.815 (3) Å
c = 23.079 (3) Å
β = 99.680 (2)°
V = 1572.4 (7) Å^3
Z = 4
Mo $K\alpha$ radiation
μ = 1.60 mm^{-1}
T = 298 K
0.22 × 0.20 × 0.19 mm

Data collection

Bruker SMART 1K CCD area-detector diffractometer
Absorption correction: multi-scan (*SADABS*; Sheldrick, 2004)
T_{min} = 0.720, T_{max} = 0.752
12290 measured reflections
3410 independent reflections
2647 reflections with $I > 2\sigma(I)$
R_{int} = 0.046

Refinement

$R[F^2 > 2\sigma(F^2)]$ = 0.037
$wR(F^2)$ = 0.089
S = 1.06
3410 reflections
233 parameters
3 restraints
H atoms treated by a mixture of independent and constrained refinement
$\Delta\rho_{max}$ = 0.39 e Å^{-3}
$\Delta\rho_{min}$ = −0.48 e Å^{-3}

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

D—H···A	D—H	H···A	D···A	D—H···A
O6—H6B···O1^{i}	0.85 (1)	2.06 (1)	2.887 (3)	167 (3)
O2—H2B···O6	0.71 (4)	1.98 (4)	2.681 (4)	172 (4)
O2—H2A···O5	0.81 (4)	2.63 (4)	3.078 (3)	116 (3)
O2—H2A···O3	0.81 (4)	1.85 (4)	2.652 (4)	170 (4)
O6—H6A···O4^{ii}	0.84 (1)	2.02 (1)	2.831 (3)	162 (3)

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

Data collection: *SMART* (Bruker, 2001); cell refinement: *SAINT* (Bruker, 2001); data reduction: *SAINT*; program(s) used to solve structure: *SHELXTL* (Sheldrick, 2008); program(s) used to refine structure: *SHELXTL*; molecular graphics: *SHELXTL*; software used to prepare material for publication: *SHELXTL* and local programs.

The College of Biological and Chemical Sciences Engineering at Jiaxing University is acknowledged for the provision of facilities to prepare and characterize the compound.

Supplementary data and figures for this paper are available from the IUCr electronic archives (Reference: QM2051).

References

Bruker (2001). *SMART* and *SAINT*. Bruker AXS Inc., Madison, Wisconsin, USA.
Creaven, B. S., Czegledi, E., Devereux, M., Enyedy, E. A., Kia, A. F. A., Karcz, D., Kellett, A., McClean, S., Nagy, N. V., Noble, A., Rockenbauer, A., Szabo-Planka, T. & Walsh, M. (2010). *Dalton Trans.* **39**, 10854–10865.
Liang, Q., Chen, X., Zhang, H. & Zou, Z. (2010). *Acta Cryst.* E**66**, m40.
Osowole, A. A., Kolawole, G. A. & Fagade, O. E. (2008). *J. Coord. Chem.* **61**, 1046–1055.
Patel, R. N., Singh, A., Shukla, K. K., Patel, D. K. & Sondhiya, V. P. (2011). *J. Coord. Chem.* **64**, 902–919.
Sheldrick, G. M. (2004). *SADABS*. University of Göttingen, Germany.
Sheldrick, G. M. (2008). *Acta Cryst.* A**64**, 112–122.

supporting information

Acta Cryst. (2012). E**68**, m275 [doi:10.1107/S1600536812004564]

A second polymorph of aqua{4-chloro-2-[(pyridin-2-ylmethyl)iminomethyl]-phenolato}copper(II) nitrate monohydrate

Jing Yu

S1. Comment

Schiff base copper(II) complexes have been received much attention due to their interesting structures and biological properties (Patel *et al.*, 2011; Creaven *et al.*, 2010; Osowole *et al.*, 2008). The title complex was first reported as triclinic space group P-1 (Liang *et al.*, 2010). We report herein a monoclinic polymorph in space group P21/c.

The title complex, (I) (Fig. 1), contains a mononuclear copper complex cation, a nitrate anion, and a water molecule. The Cu atom is coordinated by one phenolate O, one imine N, and one pyridine N of the Schiff base ligand, and one water O atom, forming a square planar coordination. The bond lengths (Table 1) are within the normal range. In the crystal, molecules are linked through intermolecular O—H···O hydrogen bonds (Table 2), to form chains along the *a* axis, Fig. 2.

S2. Experimental

To a solution of 5-chlorosalicylaldehyde (0.156 g, 1.0 mmol), 2-aminomethylpyridine (0.108 g, 1.0 mmol) in 30 ml me thanol was added slowly a solution of copper nitrate (0.241 g, 1.0 mmol) in methanol. The mixture was stirred for 2 h at room temperature to give a blue solution, which was filtered and the filtrate was left to stand at room temperature. Blue block crystals suitable for X-ray diffraction were obtained by slow evaporation.

S3. Refinement

The water H atoms were located in a difference map and refined with distances restraint of O—H = 0.85 (1) Å and H···H = 1.37 (2) Å. Other H atoms were positioned geometrically and refined using a riding model, with C—H = 0.93–0.97 Å.

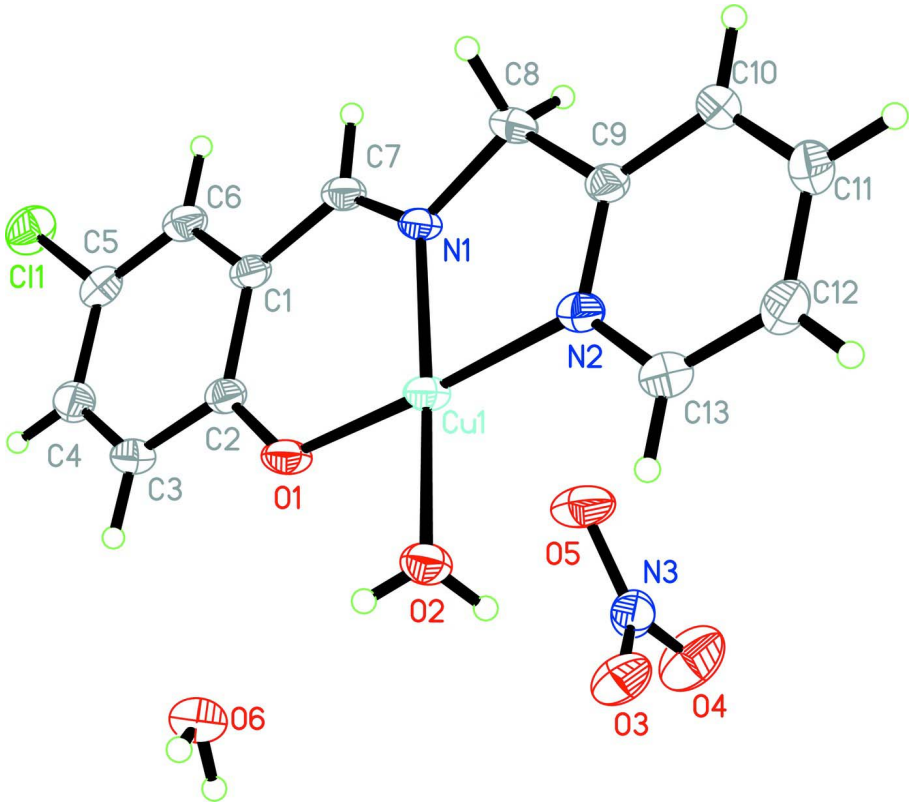


Figure 1

The molecular structure of (I), with atom labels and 30% probability displacement ellipsoids for non-H atoms.

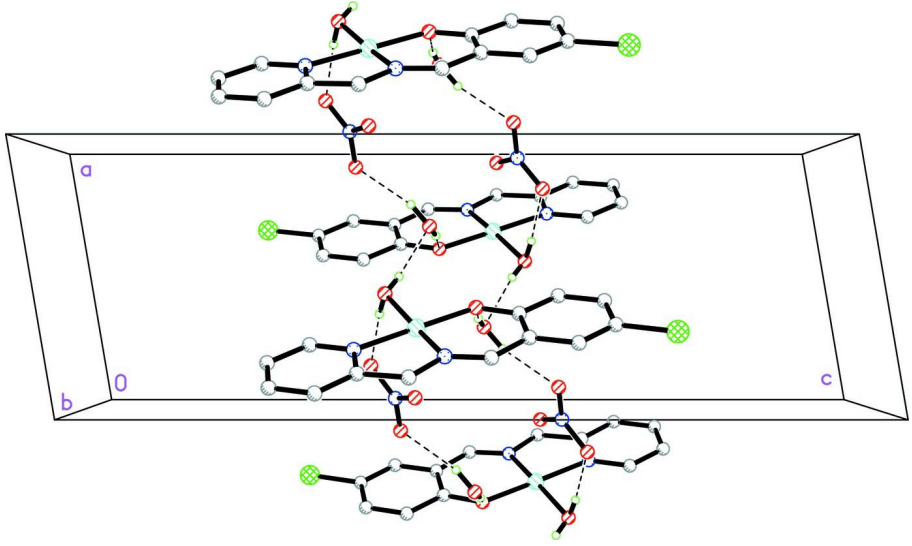


Figure 2

The packing of (I), viewed down the *b* axis. Hydrogen bonds are drawn as dashed lines. H atoms not involved in hydrogen bonding have been omitted.

aqua{4-chloro-2-[(pyridin-2-ylmethyl)iminomethyl]phenolato}copper(II) nitrate monohydrate

Crystal data

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c = 23.079 (3) Å
β = 99.680 (2)°
V = 1572.4 (7) Å^3
Z = 4
$F(000)$ = 828
D_x = 1.720 Mg m^{-3}
Mo $K\alpha$ radiation, λ = 0.71073 Å
Cell parameters from 3984 reflections
θ = 2.5–26.9°
μ = 1.60 mm^{-1}
T = 298 K
Block, blue
0.22 × 0.20 × 0.19 mm

Data collection

Bruker SMART 1K CCD area-detector diffractometer
Radiation source: fine-focus sealed tube
Graphite monochromator
ω scans
Absorption correction: multi-scan (*SADABS*; Sheldrick, 2004)
T_{min} = 0.720, T_{max} = 0.752
12290 measured reflections
3410 independent reflections
2647 reflections with $I > 2\sigma(I)$
R_{int} = 0.046
θ_{max} = 27.0°, θ_{min} = 2.5°
h = −9→9
k = −11→10
l = −29→29

Refinement

Refinement on F^2
Least-squares matrix: full
$R[F^2 > 2\sigma(F^2)]$ = 0.037
$wR(F^2)$ = 0.089
S = 1.06
3410 reflections
233 parameters
3 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
$w = 1/[\sigma^2(F_o^2) + (0.0362P)^2 + 0.669P]$ where $P = (F_o^2 + 2F_c^2)/3$
$(\Delta/\sigma)_{max} < 0.001$
$\Delta\rho_{max}$ = 0.39 e Å^{-3}
$\Delta\rho_{min}$ = −0.48 e Å^{-3}

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 (Å²)

	x	y	z	U_{iso}*/U_{eq}
Cu1	0.31780 (4)	0.86046 (4)	0.441029 (14)	0.03273 (12)
Cl1	0.30943 (12)	1.01565 (10)	0.74737 (3)	0.0561 (2)
N1	0.2244 (3)	1.0428 (2)	0.46999 (9)	0.0311 (5)
N2	0.2433 (3)	0.9578 (2)	0.36353 (9)	0.0341 (5)

N3	0.0449 (3)	0.5644 (3)	0.39668 (10)	0.0427 (6)
O1	0.3889 (3)	0.7778 (2)	0.51693 (8)	0.0387 (5)
O2	0.4383 (4)	0.6939 (3)	0.40783 (11)	0.0392 (5)
O3	0.1648 (3)	0.5200 (3)	0.37200 (10)	0.0539 (6)
O4	−0.0834 (3)	0.4839 (3)	0.39510 (12)	0.0764 (8)
O5	0.0556 (3)	0.6892 (3)	0.42285 (11)	0.0595 (6)
O6	0.6908 (3)	0.5408 (3)	0.47630 (10)	0.0499 (5)
C1	0.2878 (3)	0.9799 (3)	0.57295 (12)	0.0315 (6)
C2	0.3686 (3)	0.8378 (3)	0.56748 (11)	0.0317 (6)
C3	0.4316 (4)	0.7585 (3)	0.61943 (12)	0.0388 (7)
H3	0.4867	0.6658	0.6170	0.047*
C4	0.4142 (4)	0.8134 (3)	0.67321 (12)	0.0397 (7)
H4	0.4573	0.7583	0.7069	0.048*
C5	0.3323 (4)	0.9515 (3)	0.67803 (12)	0.0383 (7)
C6	0.2709 (4)	1.0331 (3)	0.62901 (12)	0.0380 (7)
H6	0.2169	1.1257	0.6327	0.046*
C7	0.2211 (3)	1.0744 (3)	0.52380 (12)	0.0339 (6)
H7	0.1715	1.1663	0.5317	0.041*
C8	0.1576 (4)	1.1552 (3)	0.42524 (12)	0.0383 (7)
H8A	0.2252	1.2476	0.4318	0.046*
H8B	0.0385	1.1793	0.4279	0.046*
C9	0.1674 (3)	1.0935 (3)	0.36544 (12)	0.0323 (6)
C10	0.1044 (4)	1.1749 (3)	0.31527 (13)	0.0443 (7)
H10	0.0529	1.2693	0.3177	0.053*
C11	0.1198 (4)	1.1132 (4)	0.26167 (13)	0.0486 (8)
H11	0.0773	1.1651	0.2272	0.058*
C12	0.1979 (4)	0.9748 (4)	0.25938 (13)	0.0470 (8)
H12	0.2096	0.9321	0.2234	0.056*
C13	0.2586 (4)	0.9002 (3)	0.31071 (13)	0.0420 (7)
H13	0.3122	0.8066	0.3090	0.050*
H6A	0.773 (3)	0.535 (3)	0.4574 (12)	0.045 (10)*
H2A	0.362 (5)	0.633 (4)	0.3958 (16)	0.067 (14)*
H2B	0.502 (5)	0.656 (4)	0.4284 (15)	0.047 (12)*
H6B	0.657 (5)	0.452 (2)	0.4820 (18)	0.100 (16)*

Atomic displacement parameters (Å²)

	U^{11}	U^{22}	U^{33}	U^{12}	U^{13}	U^{23}
Cu1	0.0389 (2)	0.02223 (18)	0.03751 (19)	0.00418 (15)	0.00787 (14)	−0.00095 (14)
Cl1	0.0715 (6)	0.0575 (5)	0.0400 (4)	−0.0038 (4)	0.0120 (4)	−0.0117 (4)
N1	0.0343 (12)	0.0188 (11)	0.0396 (12)	0.0034 (9)	0.0045 (10)	0.0015 (9)
N2	0.0384 (13)	0.0257 (12)	0.0381 (12)	−0.0011 (10)	0.0063 (10)	0.0001 (10)
N3	0.0444 (16)	0.0418 (16)	0.0400 (13)	−0.0038 (13)	0.0014 (11)	0.0011 (12)
O1	0.0533 (12)	0.0250 (10)	0.0385 (10)	0.0094 (9)	0.0100 (9)	−0.0021 (8)
O2	0.0399 (14)	0.0312 (12)	0.0468 (13)	0.0071 (11)	0.0079 (11)	−0.0014 (11)
O3	0.0568 (14)	0.0522 (14)	0.0567 (13)	−0.0071 (11)	0.0214 (11)	−0.0186 (11)
O4	0.0560 (16)	0.082 (2)	0.095 (2)	−0.0297 (15)	0.0250 (14)	−0.0259 (16)
O5	0.0533 (14)	0.0371 (13)	0.0913 (17)	0.0026 (11)	0.0211 (13)	−0.0148 (13)

O6	0.0554 (15)	0.0354 (13)	0.0579 (14)	0.0075 (11)	0.0067 (12)	−0.0017 (11)
C1	0.0319 (15)	0.0227 (13)	0.0401 (14)	−0.0018 (11)	0.0063 (12)	−0.0035 (11)
C2	0.0319 (15)	0.0252 (14)	0.0382 (14)	−0.0022 (11)	0.0064 (11)	−0.0022 (11)
C3	0.0438 (17)	0.0255 (15)	0.0464 (16)	0.0020 (13)	0.0057 (13)	0.0011 (12)
C4	0.0437 (18)	0.0359 (16)	0.0385 (15)	−0.0040 (13)	0.0042 (13)	0.0042 (13)
C5	0.0402 (17)	0.0370 (17)	0.0390 (15)	−0.0085 (13)	0.0099 (12)	−0.0094 (13)
C6	0.0402 (17)	0.0297 (15)	0.0451 (16)	0.0001 (13)	0.0099 (13)	−0.0076 (13)
C7	0.0329 (15)	0.0230 (13)	0.0460 (16)	0.0026 (12)	0.0073 (12)	−0.0037 (12)
C8	0.0460 (17)	0.0240 (14)	0.0438 (15)	0.0078 (13)	0.0040 (13)	0.0021 (12)
C9	0.0283 (14)	0.0245 (13)	0.0429 (15)	−0.0029 (11)	0.0021 (12)	−0.0002 (12)
C10	0.0478 (19)	0.0321 (16)	0.0468 (17)	0.0032 (14)	−0.0100 (14)	0.0017 (13)
C11	0.056 (2)	0.0434 (19)	0.0400 (16)	−0.0029 (16)	−0.0088 (14)	0.0048 (14)
C12	0.053 (2)	0.0446 (19)	0.0403 (16)	−0.0076 (16)	−0.0013 (14)	−0.0066 (14)
C13	0.0471 (18)	0.0327 (16)	0.0459 (17)	0.0002 (13)	0.0074 (14)	−0.0042 (13)

Geometric parameters (Å, º)

Cu1—O1	1.8925 (18)	C2—C3	1.404 (4)
Cu1—N1	1.932 (2)	C3—C4	1.360 (4)
Cu1—O2	1.970 (2)	C3—H3	0.9300
Cu1—N2	1.981 (2)	C4—C5	1.389 (4)
Cl1—C5	1.735 (3)	C4—H4	0.9300
N1—C7	1.277 (3)	C5—C6	1.359 (4)
N1—C8	1.463 (3)	C6—H6	0.9300
N2—C9	1.340 (3)	C7—H7	0.9300
N2—C13	1.345 (3)	C8—C9	1.497 (4)
N3—O4	1.227 (3)	C8—H8A	0.9700
N3—O3	1.241 (3)	C8—H8B	0.9700
N3—O5	1.251 (3)	C9—C10	1.381 (4)
O1—C2	1.314 (3)	C10—C11	1.375 (4)
O2—H2A	0.81 (4)	C10—H10	0.9300
O2—H2B	0.71 (4)	C11—C12	1.371 (4)
O6—H6A	0.839 (10)	C11—H11	0.9300
O6—H6B	0.845 (10)	C12—C13	1.368 (4)
C1—C6	1.403 (4)	C12—H12	0.9300
C1—C2	1.419 (3)	C13—H13	0.9300
C1—C7	1.434 (4)		
O1—Cu1—N1	94.00 (8)	C5—C4—H4	119.9
O1—Cu1—O2	89.27 (9)	C6—C5—C4	120.1 (3)
N1—Cu1—O2	171.71 (10)	C6—C5—Cl1	121.2 (2)
O1—Cu1—N2	176.94 (8)	C4—C5—Cl1	118.7 (2)
N1—Cu1—N2	83.13 (9)	C5—C6—C1	121.0 (3)
O2—Cu1—N2	93.43 (10)	C5—C6—H6	119.5
C7—N1—C8	118.4 (2)	C1—C6—H6	119.5
C7—N1—Cu1	126.06 (19)	N1—C7—C1	125.3 (2)
C8—N1—Cu1	115.50 (16)	N1—C7—H7	117.3
C9—N2—C13	118.3 (2)	C1—C7—H7	117.3

C9—N2—Cu1	114.98 (18)	N1—C8—C9	109.7 (2)
C13—N2—Cu1	126.67 (19)	N1—C8—H8A	109.7
O4—N3—O3	118.9 (3)	C9—C8—H8A	109.7
O4—N3—O5	120.7 (3)	N1—C8—H8B	109.7
O3—N3—O5	120.3 (3)	C9—C8—H8B	109.7
C2—O1—Cu1	127.32 (17)	H8A—C8—H8B	108.2
Cu1—O2—H2A	105 (3)	N2—C9—C10	122.3 (3)
Cu1—O2—H2B	115 (3)	N2—C9—C8	116.5 (2)
H2A—O2—H2B	108 (4)	C10—C9—C8	121.2 (2)
H6A—O6—H6B	109 (2)	C11—C10—C9	118.4 (3)
C6—C1—C2	119.3 (2)	C11—C10—H10	120.8
C6—C1—C7	117.2 (2)	C9—C10—H10	120.8
C2—C1—C7	123.5 (2)	C12—C11—C10	119.6 (3)
O1—C2—C3	118.7 (2)	C12—C11—H11	120.2
O1—C2—C1	123.8 (2)	C10—C11—H11	120.2
C3—C2—C1	117.5 (2)	C13—C12—C11	119.1 (3)
C4—C3—C2	121.8 (3)	C13—C12—H12	120.4
C4—C3—H3	119.1	C11—C12—H12	120.4
C2—C3—H3	119.1	N2—C13—C12	122.2 (3)
C3—C4—C5	120.2 (3)	N2—C13—H13	118.9
C3—C4—H4	119.9	C12—C13—H13	118.9

Hydrogen-bond geometry (Å, º)

D—H···*A*	*D*—H	H···*A*	*D*···*A*	*D*—H···*A*
O6—H6*B*···O1^{i}	0.85 (1)	2.06 (1)	2.887 (3)	167 (3)
O2—H2*B*···O6	0.71 (4)	1.98 (4)	2.681 (4)	172 (4)
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Symmetry codes: (i) −*x*+1, −*y*+1, −*z*+1; (ii) *x*+1, *y*, *z*.