

Acta Crystallographica Section E

Structure Reports

Online

ISSN 1600-5368

4,11-Diaza-1,8-diazoniacyclotetradecane bis(pyridine-2-carboxylate) dihydrate

Nam-Ho Kim and Kwang Ha*

School of Applied Chemical Engineering, The Research Institute of Catalysis, Chonnam National University, Gwangju 500-757, Republic of Korea
Correspondence e-mail: hakwang@chonnam.ac.kr

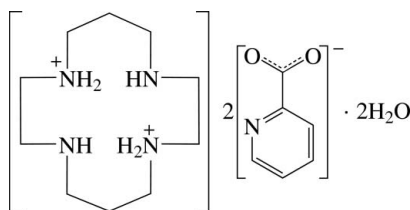
Received 10 September 2009; accepted 15 September 2009

Key indicators: single-crystal X-ray study; $T = 200$ K; mean $\sigma(\text{C}-\text{C}) = 0.003$ Å; R factor = 0.055; wR factor = 0.123; data-to-parameter ratio = 20.5.

The asymmetric unit of the title compound, $\text{C}_{10}\text{H}_{26}\text{N}_4^{2+} \cdot 2\text{C}_6\text{H}_4\text{NO}_2^- \cdot 2\text{H}_2\text{O}$, consists of half of a doubly protonated 1,4,8,11-tetraazacyclotetradecane (cyclam) dication, a pyridine-2-carboxylate anion and a solvent water molecule. The complete dication is generated by a crystallographic centre and adopts an endodentate conformation which may be influenced by intramolecular $\text{N}-\text{H} \cdots \text{N}$ hydrogen bonding. The carboxylate group of the anion appears to be delocalized on the basis of the $\text{C}-\text{O}$ bond lengths [1.257 (2) and 1.250 (2) Å]. In the crystal structure, the components are linked by intermolecular $\text{N}-\text{H} \cdots \text{O}$, $\text{N}-\text{H} \cdots \text{N}$ and $\text{O}-\text{H} \cdots \text{O}$ hydrogen bonds.

Related literature

For the crystal structures of $[\text{H}_2(\text{cyclam})]X$ [$X = (\text{ClO}_4)_2$ or Cl_2], see: Nave & Truter (1974); Kim *et al.* (2009). For the crystal structures of $[\text{H}_4(\text{cyclam})]X \cdot n\text{H}_2\text{O}$ [$X = \text{Cl}_4$, Br_4 , $(\text{ClO}_4)_4$, $(\text{SCN})_4$, $(\text{SO}_4)_2$ or $(p\text{-CH}_3\text{C}_6\text{H}_4\text{SO}_3)_4$], see: Robinson *et al.* (1989); Subramanian & Zaworotko (1995). For the structure of pyridine-2-carboxylic acid, see: Hamazaki *et al.* (1998). For the crystal structures of pyridine-2-carboxylate compounds, see: Kim & Ha (2009*a,b*).



Experimental

Crystal data

$\text{C}_{10}\text{H}_{26}\text{N}_4^{2+} \cdot 2\text{C}_6\text{H}_4\text{NO}_2^- \cdot 2\text{H}_2\text{O}$
 $M_r = 482.58$
Monoclinic, $P2_1/c$
 $a = 10.2746$ (8) Å
 $b = 12.0551$ (9) Å
 $c = 10.3244$ (8) Å
 $\beta = 93.104$ (2)°
 $V = 1276.92$ (17) Å³
 $Z = 2$
Mo $K\alpha$ radiation
 $\mu = 0.09$ mm⁻¹
 $T = 200$ K
 $0.22 \times 0.17 \times 0.11$ mm

Data collection

Bruker SMART 1000 CCD diffractometer
Absorption correction: multi-scan (SADABS; Sheldrick, 1996)
 $T_{\min} = 0.868$, $T_{\max} = 1.000$
9364 measured reflections
3152 independent reflections
1540 reflections with $I > 2\sigma(I)$
 $R_{\text{int}} = 0.070$

Refinement

$R[F^2 > 2\sigma(F^2)] = 0.055$
 $wR(F^2) = 0.123$
 $S = 1.02$
3152 reflections
154 parameters
H-atom parameters constrained
 $\Delta\rho_{\text{max}} = 0.22$ e Å⁻³
 $\Delta\rho_{\text{min}} = -0.25$ e Å⁻³

Table 1

Hydrogen-bond geometry (Å, °).

$D-\text{H} \cdots A$	$D-\text{H}$	$\text{H} \cdots A$	$D \cdots A$	$D-\text{H} \cdots A$
$\text{N}2-\text{H}21 \cdots \text{O}3$	0.92	2.02	2.932 (2)	170
$\text{N}3-\text{H}32 \cdots \text{N}2$	0.92	2.53	2.926 (3)	106
$\text{N}3-\text{H}32 \cdots \text{N}2^i$	0.92	2.08	2.846 (3)	139
$\text{N}3-\text{H}31 \cdots \text{O}1^{\text{ii}}$	0.92	1.86	2.749 (2)	161
$\text{N}3-\text{H}31 \cdots \text{N}1^{\text{ii}}$	0.92	2.46	3.039 (2)	121
$\text{O}3-\text{H}3A \cdots \text{O}1^{\text{iii}}$	0.84	1.99	2.808 (2)	165
$\text{O}3-\text{H}3B \cdots \text{O}2^{\text{iv}}$	0.84	1.91	2.739 (2)	168

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

Data collection: SMART (Bruker, 2007); cell refinement: SAINT (Bruker, 2007); data reduction: SAINT; program(s) used to solve structure: SHELXTL (Sheldrick, 2008); program(s) used to refine structure: SHELXTL; molecular graphics: ORTEP-3 for Windows (Farrugia, 1997) and PLATON (Spek, 2009); software used to prepare material for publication: SHELXTL.

This work was supported by a Korea Research Foundation Grant funded by the Korean Government (MOEHRD) (KRF-2007-412-J02001).

Supplementary data and figures for this paper are available from the IUCr electronic archives (Reference: NK2004).

References

- Bruker (2007). SMART and SAINT. Bruker AXS Inc., Madison, Wisconsin, USA.
Farrugia, L. J. (1997). *J. Appl. Cryst.* **30**, 565.
Hamazaki, H., Hosomi, H., Takeda, S., Kataoka, H. & Ohba, S. (1998). *Acta Cryst.* **C54**, IUC9800049.
Kim, N.-H. & Ha, K. (2009*a*). *Acta Cryst.* **E65**, o1415.
Kim, N.-H. & Ha, K. (2009*b*). *Acta Cryst.* **E65**, o2151.
Kim, N.-H., Hwang, I.-C. & Ha, K. (2009). *Acta Cryst.* **E65**, o2128.
Nave, C. & Truter, M. R. (1974). *J. Chem. Soc. Dalton Trans.* pp 2351–2354.
Robinson, G. H., Sangokoya, S. A., Pennington, W. T., Self, M. F. & Rogers, R. D. (1989). *J. Coord. Chem.* **19**, 287–294.
Sheldrick, G. M. (1996). SADABS. University of Göttingen, Germany.
Sheldrick, G. M. (2008). *Acta Cryst.* **A64**, 112–122.
Spek, A. L. (2009). *Acta Cryst.* **D65**, 148–155.
Subramanian, S. & Zaworotko, M. J. (1995). *Can. J. Chem.* **73**, 414–424.

supporting information

Acta Cryst. (2009). E65, o2504 [doi:10.1107/S1600536809037258]

4,11-Diaza-1,8-diazoniacyclotetradecane bis(pyridine-2-carboxylate) dihydrate**Nam-Ho Kim and Kwang Ha****S1. Comment**

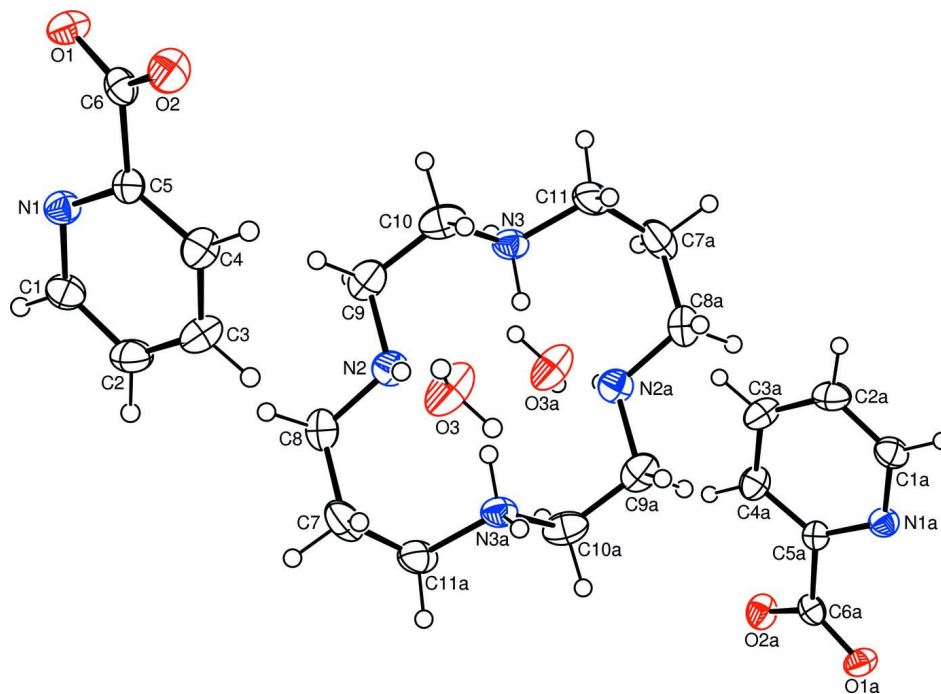
The asymmetric unit of the title compound, $C_{10}H_{26}N_4^{2+} \cdot 2C_6H_4NO_2^- \cdot 2H_2O$, (I), consists of half of a doubly protonated 1,4,8,11-tetraazacyclotetradecane (cyclam) dication, a pyridine-2-carboxylate anion and a solvent water molecule (Fig. 1). The macrocyclic dication contains two protonated N atoms and two secondary amine N atoms, and is located on a centre of inversion. The dication adopts an endodentate conformation with the N atoms oriented towards the centre of the macrocyclic cavity. The conformation may be stabilized by intramolecular N—H \cdots N hydrogen bonding (Table 1 and Fig. 2). The N2—C9—C10—N3 torsion angle of 66.5 (2) $^\circ$ displays the *gauche* conformation for the group within the dication. A similar conformation is also observed in the structures cyclam (Robinson *et al.*, 1989) and $[H_2(\text{cyclam})]X$ [$X = (\text{ClO}_4)_2$ or Cl_2] (Nave & Truter, 1974; Kim *et al.*, 2009). Unlike cyclam and the dication, the tetracation, $[H_4(\text{cyclam})]^{4+}$, adopts an exodentate conformation, in which all four N atoms are oriented away from the ring cavity, occupying positions as far away as possible from each other on the ring periphery (Robinson *et al.*, 1989; Subramanian & Zaworotko, 1995). The protonated N—C bond lengths (N3—C10/C11: 1.482 (3)/1.487 (3) Å) are longer than unprotonated N—C bond lengths (N2—C8/C9: 1.468 (3)/1.459 (3) Å). The carboxylate group of the anion appears to be delocalized on the basis of the C—O bond lengths (C—O: 1.250 (2) and 1.257 (2) Å). The components of the crystal structure are linked by intermolecular N—H \cdots O, N—H \cdots N and O—H \cdots O hydrogen bonds into one-dimensional chains along [001] (Table 1 and Fig. 2).

S2. Experimental

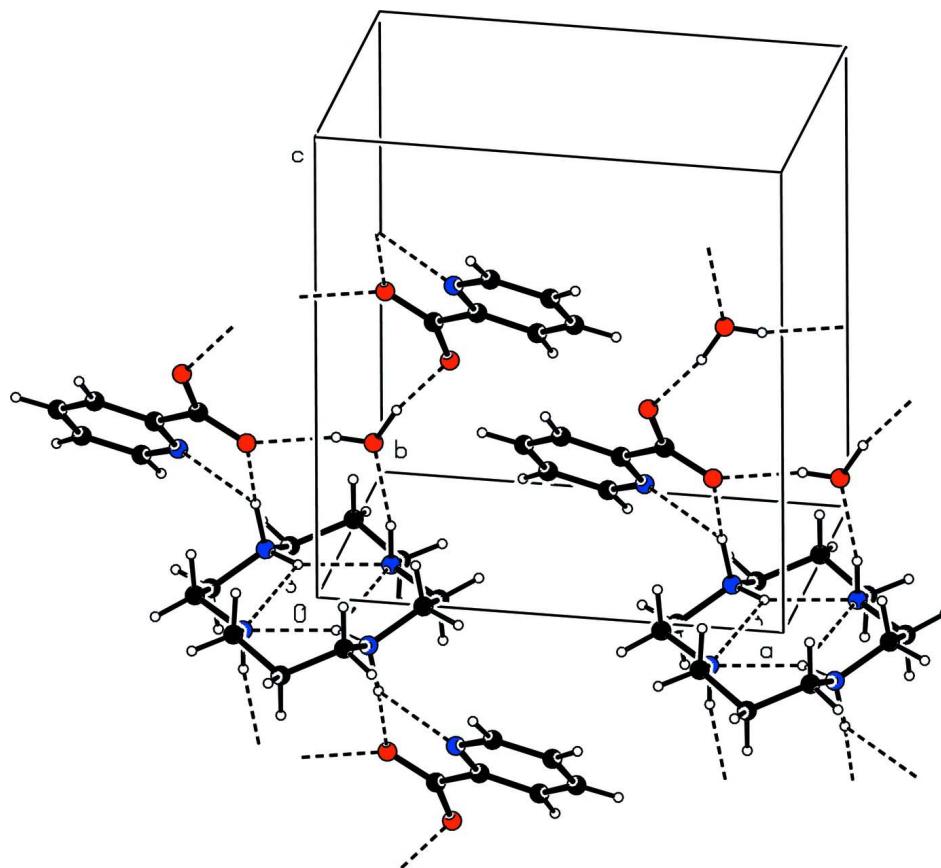
A solution of 1,4,8,11-tetraazacyclotetradecane (0.100 g, 0.50 mmol) and pyridine-2-carboxylic acid (0.123 g, 1.00 mmol) in H_2O (10 ml) was stirred for 3 h at 60 $^\circ\text{C}$. The solvent was removed under vacuum and the residue was washed with acetone/ether, to give a white powder (0.221 g). Crystals suitable for X-ray analysis were obtained by slow evaporation from an acetone solution.

S3. Refinement

H atoms were positioned geometrically and allowed to ride on their respective parent atoms [C—H = 0.95 (CH) or 0.99 (CH_2) Å, N—H = 0.92 Å, O—H = 0.84 Å and $U_{\text{iso}}(\text{H}) = 1.2U_{\text{eq}}(\text{C}, \text{N})$ or $1.5U_{\text{eq}}(\text{O})$].

**Figure 1**

The structure of (I), with displacement ellipsoids drawn at the 40% probability level for non-H atoms [Symmetry code: (a) $-x, -y, -z$].

**Figure 2**

View of the unit-cell contents of (I). Hydrogen bonding interactions are drawn with dashed lines.

4,11-Diaza-1,8-diazoniacyclotetradecane bis(pyridine-2-carboxylate) dihydrate

Crystal data

$C_{10}H_{26}N_4^{2+} \cdot 2C_6H_4NO_2^- \cdot 2(H_2O)$

$M_r = 482.58$

Monoclinic, $P2_1/c$

Hall symbol: $-P\ 2_1/c$

$a = 10.2746\ (8)\ \text{\AA}$

$b = 12.0551\ (9)\ \text{\AA}$

$c = 10.3244\ (8)\ \text{\AA}$

$\beta = 93.104\ (2)^\circ$

$V = 1276.92\ (17)\ \text{\AA}^3$

$Z = 2$

$F(000) = 520$

$D_x = 1.255\ \text{Mg m}^{-3}$

Mo $K\alpha$ radiation, $\lambda = 0.71073\ \text{\AA}$

Cell parameters from 1456 reflections

$\theta = 2.6\text{--}24.3^\circ$

$\mu = 0.09\ \text{mm}^{-1}$

$T = 200\ \text{K}$

Block, colourless

$0.22 \times 0.17 \times 0.11\ \text{mm}$

Data collection

Bruker SMART 1000 CCD

diffractometer

Radiation source: fine-focus sealed tube

Graphite monochromator

ω scans

Absorption correction: multi-scan

(*SADABS*; Sheldrick, 1996)

$T_{\min} = 0.868$, $T_{\max} = 1.000$

9364 measured reflections

3152 independent reflections

1540 reflections with $I > 2\sigma(I)$

$R_{\text{int}} = 0.070$

$\theta_{\max} = 28.3^\circ$, $\theta_{\min} = 2.0^\circ$

$h = -11 \rightarrow 13$

$k = -16 \rightarrow 15$

$l = -13 \rightarrow 13$

Refinement

Refinement on F^2
 Least-squares matrix: full
 $R[F^2 > 2\sigma(F^2)] = 0.055$
 $wR(F^2) = 0.123$
 $S = 1.02$
 3152 reflections
 154 parameters
 0 restraints
 Primary atom site location: structure-invariant
 direct methods

Secondary atom site location: difference Fourier
 map
 Hydrogen site location: inferred from
 neighbouring sites
 H-atom parameters constrained
 $w = 1/[\sigma^2(F_o^2) + (0.0315P)^2]$
 where $P = (F_o^2 + 2F_c^2)/3$
 $(\Delta/\sigma)_{\max} < 0.001$
 $\Delta\rho_{\max} = 0.22 \text{ e } \text{\AA}^{-3}$
 $\Delta\rho_{\min} = -0.25 \text{ e } \text{\AA}^{-3}$

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 (\AA^2)

	x	y	z	$U_{\text{iso}}^*/U_{\text{eq}}$
O1	0.84295 (15)	0.05500 (12)	0.30790 (13)	0.0377 (4)
O2	0.71238 (16)	0.00216 (13)	0.46269 (15)	0.0433 (5)
N1	0.67291 (18)	0.21986 (15)	0.23751 (17)	0.0333 (5)
C1	0.5839 (2)	0.29517 (19)	0.1972 (2)	0.0383 (6)
H1	0.6109	0.3538	0.1434	0.046*
C2	0.4558 (2)	0.2926 (2)	0.2290 (2)	0.0434 (6)
H2	0.3960	0.3481	0.1984	0.052*
C3	0.4166 (2)	0.2074 (2)	0.3066 (2)	0.0449 (7)
H3	0.3282	0.2017	0.3283	0.054*
C4	0.5066 (2)	0.1306 (2)	0.3524 (2)	0.0384 (6)
H4	0.4818	0.0725	0.4082	0.046*
C5	0.6339 (2)	0.13915 (18)	0.31597 (19)	0.0278 (5)
C6	0.7385 (2)	0.05841 (18)	0.3658 (2)	0.0312 (5)
N2	0.14631 (18)	0.09306 (16)	0.05713 (17)	0.0374 (5)
H21	0.1475	0.0811	0.1452	0.045*
N3	0.12754 (18)	-0.12813 (15)	-0.05896 (17)	0.0360 (5)
H31	0.1456	-0.1194	-0.1447	0.043*
H32	0.0565	-0.0845	-0.0438	0.043*
C7	0.0412 (3)	0.2719 (2)	0.0986 (3)	0.0510 (7)
H7A	0.0561	0.3529	0.0946	0.061*
H7B	0.0435	0.2503	0.1912	0.061*
C8	0.1501 (2)	0.2132 (2)	0.0340 (2)	0.0446 (7)
H8A	0.1424	0.2276	-0.0605	0.054*
H8B	0.2349	0.2430	0.0681	0.054*
C9	0.2570 (2)	0.0350 (2)	0.0057 (2)	0.0459 (7)

H9A	0.3388	0.0598	0.0519	0.055*
H9B	0.2625	0.0525	-0.0875	0.055*
C10	0.2406 (2)	-0.0876 (2)	0.0229 (2)	0.0485 (7)
H10A	0.3206	-0.1264	-0.0015	0.058*
H10B	0.2270	-0.1042	0.1151	0.058*
C11	0.0922 (3)	-0.24601 (19)	-0.0372 (3)	0.0487 (7)
H11A	0.0928	-0.2609	0.0571	0.058*
H11B	0.1577	-0.2949	-0.0748	0.058*
O3	0.11622 (16)	0.04470 (16)	0.33220 (14)	0.0609 (6)
H3A	0.0351	0.0418	0.3385	0.091*
H3B	0.1593	0.0314	0.4019	0.091*

Atomic displacement parameters (Å²)

	U^{11}	U^{22}	U^{33}	U^{12}	U^{13}	U^{23}
O1	0.0313 (10)	0.0450 (11)	0.0374 (9)	0.0095 (8)	0.0064 (8)	0.0074 (8)
O2	0.0425 (11)	0.0484 (11)	0.0389 (9)	-0.0014 (8)	0.0010 (8)	0.0163 (8)
N1	0.0335 (12)	0.0338 (11)	0.0327 (10)	0.0028 (9)	0.0026 (9)	0.0033 (9)
C1	0.0439 (16)	0.0346 (14)	0.0361 (13)	0.0048 (12)	-0.0003 (12)	0.0055 (11)
C2	0.0398 (16)	0.0423 (16)	0.0476 (15)	0.0118 (13)	-0.0042 (13)	-0.0027 (13)
C3	0.0305 (15)	0.0573 (18)	0.0473 (15)	0.0062 (13)	0.0050 (12)	0.0018 (14)
C4	0.0377 (15)	0.0464 (15)	0.0314 (13)	-0.0027 (13)	0.0038 (11)	0.0052 (12)
C5	0.0289 (13)	0.0303 (13)	0.0239 (11)	-0.0024 (11)	-0.0012 (10)	-0.0007 (10)
C6	0.0365 (15)	0.0284 (13)	0.0281 (12)	-0.0041 (11)	-0.0045 (11)	-0.0015 (11)
N2	0.0427 (13)	0.0406 (12)	0.0296 (10)	0.0016 (10)	0.0071 (9)	-0.0003 (9)
N3	0.0392 (12)	0.0361 (12)	0.0333 (10)	0.0082 (10)	0.0086 (9)	0.0035 (9)
C7	0.074 (2)	0.0301 (14)	0.0499 (16)	-0.0049 (14)	0.0144 (16)	0.0006 (12)
C8	0.0459 (17)	0.0431 (16)	0.0448 (15)	-0.0104 (13)	0.0036 (13)	0.0033 (13)
C9	0.0392 (16)	0.0610 (18)	0.0377 (14)	-0.0040 (14)	0.0035 (12)	-0.0116 (13)
C10	0.0424 (16)	0.0643 (19)	0.0383 (14)	0.0160 (14)	-0.0020 (13)	-0.0044 (13)
C11	0.0565 (18)	0.0364 (16)	0.0554 (17)	0.0108 (13)	0.0231 (15)	0.0159 (13)
O3	0.0392 (11)	0.1048 (16)	0.0385 (10)	-0.0009 (11)	0.0012 (9)	0.0222 (10)

Geometric parameters (Å, °)

O1—C6	1.257 (2)	N3—H31	0.9200
O2—C6	1.250 (2)	N3—H32	0.9200
N1—C1	1.339 (3)	C7—C8	1.509 (3)
N1—C5	1.341 (2)	C7—C11 ⁱ	1.511 (4)
C1—C2	1.374 (3)	C7—H7A	0.9900
C1—H1	0.9500	C7—H7B	0.9900
C2—C3	1.377 (3)	C8—H8A	0.9900
C2—H2	0.9500	C8—H8B	0.9900
C3—C4	1.374 (3)	C9—C10	1.499 (3)
C3—H3	0.9500	C9—H9A	0.9900
C4—C5	1.385 (3)	C9—H9B	0.9900
C4—H4	0.9500	C10—H10A	0.9900
C5—C6	1.519 (3)	C10—H10B	0.9900

N2—C9	1.459 (3)	C11—C7 ⁱ	1.511 (4)
N2—C8	1.468 (3)	C11—H11A	0.9900
N2—H21	0.9200	C11—H11B	0.9900
N3—C10	1.482 (3)	O3—H3A	0.8400
N3—C11	1.487 (3)	O3—H3B	0.8400
C1—N1—C5	117.3 (2)	C11 ⁱ —C7—H7A	108.9
N1—C1—C2	123.9 (2)	C8—C7—H7B	108.9
N1—C1—H1	118.1	C11 ⁱ —C7—H7B	108.9
C2—C1—H1	118.1	H7A—C7—H7B	107.7
C1—C2—C3	118.0 (2)	N2—C8—C7	111.34 (19)
C1—C2—H2	121.0	N2—C8—H8A	109.4
C3—C2—H2	121.0	C7—C8—H8A	109.4
C4—C3—C2	119.4 (2)	N2—C8—H8B	109.4
C4—C3—H3	120.3	C7—C8—H8B	109.4
C2—C3—H3	120.3	H8A—C8—H8B	108.0
C3—C4—C5	119.0 (2)	N2—C9—C10	109.6 (2)
C3—C4—H4	120.5	N2—C9—H9A	109.8
C5—C4—H4	120.5	C10—C9—H9A	109.8
N1—C5—C4	122.4 (2)	N2—C9—H9B	109.8
N1—C5—C6	116.15 (19)	C10—C9—H9B	109.8
C4—C5—C6	121.4 (2)	H9A—C9—H9B	108.2
O2—C6—O1	126.1 (2)	N3—C10—C9	110.3 (2)
O2—C6—C5	116.1 (2)	N3—C10—H10A	109.6
O1—C6—C5	117.80 (19)	C9—C10—H10A	109.6
C9—N2—C8	112.69 (19)	N3—C10—H10B	109.6
C9—N2—H21	108.4	C9—C10—H10B	109.6
C8—N2—H21	108.5	H10A—C10—H10B	108.1
C10—N3—C11	114.79 (19)	N3—C11—C7 ⁱ	110.9 (2)
C10—N3—H31	108.6	N3—C11—H11A	109.5
C11—N3—H31	108.6	C7 ⁱ —C11—H11A	109.5
C10—N3—H32	108.6	N3—C11—H11B	109.5
C11—N3—H32	108.6	C7 ⁱ —C11—H11B	109.5
H31—N3—H32	107.5	H11A—C11—H11B	108.0
C8—C7—C11 ⁱ	113.3 (2)	H3A—O3—H3B	113.9
C8—C7—H7A	108.9		
C5—N1—C1—C2	-1.6 (3)	C4—C5—C6—O2	-16.4 (3)
N1—C1—C2—C3	-0.2 (4)	N1—C5—C6—O1	-17.0 (3)
C1—C2—C3—C4	2.0 (4)	C4—C5—C6—O1	164.4 (2)
C2—C3—C4—C5	-1.9 (3)	C9—N2—C8—C7	-174.9 (2)
C1—N1—C5—C4	1.7 (3)	C11 ⁱ —C7—C8—N2	-68.7 (3)
C1—N1—C5—C6	-176.94 (19)	C8—N2—C9—C10	-175.26 (19)
C3—C4—C5—N1	0.0 (3)	C11—N3—C10—C9	-172.71 (19)
C3—C4—C5—C6	178.6 (2)	N2—C9—C10—N3	66.5 (2)
N1—C5—C6—O2	162.19 (18)	C10—N3—C11—C7 ⁱ	165.97 (19)

Symmetry code: (i) $-x, -y, -z$.

Hydrogen-bond geometry (Å, °)

<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>
N2—H21 \cdots O3	0.92	2.02	2.932 (2)	170
N3—H32 \cdots N2	0.92	2.53	2.926 (3)	106
N3—H32 \cdots N2 ⁱ	0.92	2.08	2.846 (3)	139
N3—H31 \cdots O1 ⁱⁱ	0.92	1.86	2.749 (2)	161
N3—H31 \cdots N1 ⁱⁱ	0.92	2.46	3.039 (2)	121
O3—H3A \cdots O1 ⁱⁱⁱ	0.84	1.99	2.808 (2)	165
O3—H3B \cdots O2 ^{iv}	0.84	1.91	2.739 (2)	168

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