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Ethyl 2-{3-[(6-chloropyridin-3-yl)methyl]-2-(nitroimino)imidazolidin-1-yl}acetate

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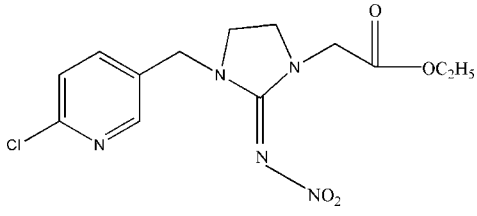
Key indicators: single-crystal X-ray study; T = 293 K; mean σ(C–C) = 0.003 Å; R factor = 0.038; wR factor = 0.090; data-to-parameter ratio = 14.3.

In the title compound, $C_{13}H_{16}ClN_5O_4$, the imidazole ring is in a slight envelope conformation. The dihedral angle between the pyridine ring and the four essentially planar atoms [maximum deviation 0.015 (2) Å] of the imidazole ring is 80.8 (1)°. In, the crystal, weak C—H···O and C—H···N hydrogen bonds are present. In addition, there are weak π–π stacking interactions between symmetry-related pyridine rings with a centroid–centroid distance of 3.807 (1) Å.

Related literature

For background to the insecticidal applications of imidaclo-prid [systematic name: (*E*)-1-(6-chloro-3-pyridylmethyl)-*N*-nitroimidazolidin-2-ylideneamine], see: Deshmukh *et al.* (2011, 2012); Zhao *et al.* (2010). For related structures, see: Kapoor *et al.* (2011, 2012); Kant *et al.* (2012).



Experimental

Crystal data

$C_{13}H_{16}ClN_5O_4$
M_r = 341.76
Monoclinic, $P2_1/c$
a = 7.8136 (2) Å
b = 19.3483 (4) Å
c = 10.1926 (2) Å
β = 100.346 (2)°
V = 1515.86 (6) Å^3
Z = 4
Mo $K\alpha$ radiation
μ = 0.28 mm^{-1}
T = 293 K
0.3 × 0.2 × 0.1 mm

Data collection

Oxford Diffraction Xcalibur Sapphire3 diffractometer
Absorption correction: multi-scan (*CrysAlis PRO RED*; Oxford Diffraction, 2010)
T_{min} = 0.868, T_{max} = 1.000
47458 measured reflections
2983 independent reflections
2387 reflections with $I > 2\sigma(I)$
R_{int} = 0.046

Refinement

$R[F^2 > 2\sigma(F^2)]$ = 0.038
$wR(F^2)$ = 0.090
S = 1.02
2983 reflections
209 parameters
H-atom parameters constrained
$\Delta\rho_{max}$ = 0.23 e Å^{-3}
$\Delta\rho_{min}$ = −0.29 e Å^{-3}

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

D—H···A	D—H	H···A	D···A	D—H···A
C9—H9B···O15^{i}	0.97	2.50	3.358 (2)	147
C17—H17A···N1^{ii}	0.97	2.57	3.509 (2)	163

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

Data collection: *CrysAlis PRO CCD* (Oxford Diffraction, 2010); cell refinement: *CrysAlis PRO CCD*; data reduction: *CrysAlis PRO RED* (Oxford Diffraction, 2010); program(s) used to solve structure: *SHELXS97* (Sheldrick, 2008); program(s) used to refine structure: *SHELXL97* (Sheldrick, 2008); molecular graphics: *ORTEP-3* (Farrugia, 1997) and *PLATON* (Spek, 2009); software used to prepare material for publication: *PLATON*.

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Supplementary data and figures for this paper are available from the IUCr electronic archives (Reference: LH5425).

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supporting information

Acta Cryst. (2012). E**68**, o987 [https://doi.org/10.1107/S160053681200918X]

Ethyl 2-{3-[(6-chloropyridin-3-yl)methyl]-2-(nitroimino)imidazolidin-1-yl}acetate

Kamini Kapoor, Madhukar B. Deshmukh, Chetan S. Shripanavar, Vivek K. Gupta and Rajni Kant

S1. Comment

The discovery of imidacloprid has been referred to as a milestone in the past three decades of insecticidal research. The nitroguanidine moiety of imidacloprid is also a common site for metabolism *via* cleavage to the guanidine and reduction to di-nitro-imidacloprid. The insecticidal activity of nitroguanidine was found to be 10,000 fold higher than that of natural insecticide nicotine (Deshmukh *et al.*, 2012). In mammalian systems the nitro group of imidacloprid has been postulated to be reduced to nitrosoguanidine and aminoguanidine and then cleaved to the guanidine and urea derivatives (Deshmukh *et al.*, 2011). Therefore, in a search for new neonicotinoid insecticides with improved profiles, neonicotinoid derivatives containing N-oxalyl groups were designed and synthesized (Zhao *et al.*, 2010).

The molecular structure of the title compound is shown in Fig. 1. The bond lengths and angles are comparable to those common to related structures (Kapoor *et al.*, 2011,2012; Kant *et al.*, 2012). The imidazole ring is in a slight envelope conformation with atom C9 forming the flap. The dihedral angle between the pyridine ring [N1/C2-C6] and the four essentially planar atoms [N8/N11/C10/C12 (maximum deviation 0.015 (2)Å for C12)] of the imidazole ring is 80.8 (1)°. In the crystal, molecules are connected by pairs of weak C—H···O hydrogen bonds into centrosymmetric dimers, which are in turn, linked into columns along [100] by weak C—H···N hydrogen bonds (Fig .2). In addition, there is a weak $\pi\cdots\pi$ interaction between the pyridine ring at (*x*, *y*, *z*) and the pyridine ring at (1 - *x*, 1 - *y*, - *z*) [centroid separation = 3.807 (1) Å, interplanar spacing = 3.368 Å and centroid shift = 1.77 Å].

S2. Experimental

Imidacloprid (10.20 g, 0.04 mol) in 30 ml acetone, ethyl chloroacetate (7.32 g, 0.06 mol) was refluxed for about 24 h in presence of 10 g m K_2CO_3. An aliquot of sample was taken to monitor the progress of reaction by TLC. After completion of reaction, the hot reaction mixture was filtered to remove excess K_2CO_3. Filtrate was then dried under reduced pressure giving a white solid, Yield 80%. The synthesized compound was dissolved in methanol, by the process of slow evaporation a fine crystalline compound separated out.

S3. Refinement

H atoms were positioned geometrically and were treated as riding on their parent C atoms, with C—H distances of 0.93–0.97 Å.

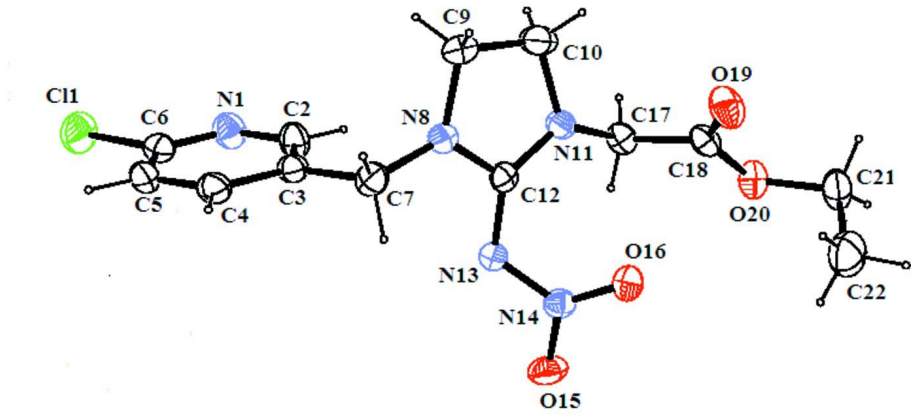

Figure 1

The molecular structure with ellipsoids are drawn at the 40% probability level. H atoms are shown as small spheres of arbitrary radii.

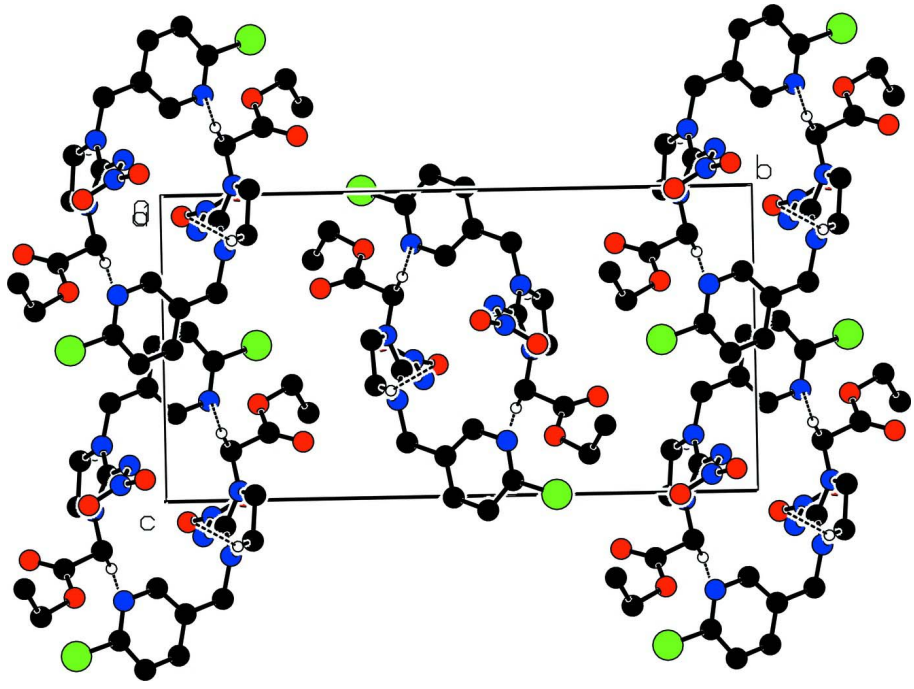

Figure 2

Part of the crystal structure with weak hydrogen bonds shown as dashed lines.

Ethyl 2-{3-[(6-chloropyridin-3-yl)methyl]-2-(nitroimino)imidazolidin-1-yl}acetate

Crystal data

$C_{13}H_{16}ClN_5O_4$
$M_r = 341.76$

Monoclinic, $P2_1/c$
Hall symbol: -P 2ybc

a = 7.8136 (2) Å
b = 19.3483 (4) Å
c = 10.1926 (2) Å
β = 100.346 (2)°
V = 1515.86 (6) Å^3
Z = 4
F(000) = 712
D_x = 1.498 Mg m^{-3}

Mo $K\alpha$ radiation, λ = 0.71073 Å
Cell parameters from 21411 reflections
θ = 3.6–29.1°
μ = 0.28 mm^{-1}
T = 293 K
Plate, white
0.3 × 0.2 × 0.1 mm

Data collection

Oxford Diffraction Xcalibur Sapphire3 diffractometer
Radiation source: fine-focus sealed tube
Graphite monochromator
Detector resolution: 16.1049 pixels mm^{-1}
ω scans
Absorption correction: multi-scan (*CrysAlis PRO* RED; Oxford Diffraction, 2010)
T_{min} = 0.868, T_{max} = 1.000

47458 measured reflections
2983 independent reflections
2387 reflections with $I > 2\sigma(I)$
R_{int} = 0.046
θ_{max} = 26.0°, θ_{min} = 3.7°
h = −9→9
k = −23→23
l = −12→12

Refinement

Refinement on F^2
Least-squares matrix: full
$R[F^2 > 2\sigma(F^2)]$ = 0.038
$wR(F^2)$ = 0.090
S = 1.02
2983 reflections
209 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
$w = 1/[\sigma^2(F_o^2) + (0.0372P)^2 + 0.6841P]$ where $P = (F_o^2 + 2F_c^2)/3$
$(\Delta/\sigma)_{max}$ = 0.001
$\Delta\rho_{max}$ = 0.23 e Å^{-3}
$\Delta\rho_{min}$ = −0.29 e Å^{-3}

Special details

Experimental. *CrysAlis PRO*, Oxford Diffraction Ltd., Version 1.171.34.40 (release 27–08-2010 CrysAlis171. NET) (compiled Aug 27 2010,11:50:40) Empirical absorption correction using spherical harmonics, implemented in SCALE3 ABSPACK scaling algorithm.
IR (cm-1): 3008, 2991, 2908, 1743, 1558, 1548. 1H NMR ?: 1.29(t, J: 7.5 Hz, CH3),3.59(t, J: 7.5 Hz, CH2), 3.84(t, J: 7.5 Hz, CH2), 4.06 (s, CH2), 4.24 (q, J:7.5 Hz, OCH2), 4.50 (s, CH2), 7.37 (d, J: 8.2 Hz, Py1H), 7.74 (dd, J1: 7.5,J2: 2.5 Hz, Py1H), 8.32 (s, Py1H) p.p.m.. LCMS/MS (ESI, m/*z*): 342.0891 (*M*+H)+,295.0881, 261.1297, 170.0910.
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 (Å^2)

	x	y	z	U_{iso}*/U_{eq}
Cl1	0.49235 (8)	0.34128 (3)	−0.00544 (6)	0.06124 (18)
N1	0.4056 (2)	0.42363 (8)	0.17333 (16)	0.0437 (4)
C2	0.3295 (2)	0.48018 (10)	0.21371 (18)	0.0402 (4)

H2	0.3394	0.4877	0.3049	0.048*
C3	0.2376 (2)	0.52773 (9)	0.12776 (16)	0.0316 (4)
C4	0.2223 (2)	0.51489 (10)	−0.00787 (17)	0.0376 (4)
H4	0.1600	0.5453	−0.0692	0.045*
C5	0.2988 (2)	0.45741 (10)	−0.05220 (18)	0.0406 (4)
H5	0.2903	0.4481	−0.1426	0.049*
C6	0.3884 (2)	0.41447 (10)	0.04397 (19)	0.0387 (4)
C7	0.1554 (2)	0.59159 (10)	0.17545 (16)	0.0362 (4)
H7A	0.1805	0.6309	0.1230	0.043*
H7B	0.0302	0.5855	0.1602	0.043*
N8	0.21587 (18)	0.60656 (8)	0.31527 (14)	0.0337 (3)
C9	0.3777 (2)	0.64367 (10)	0.36251 (18)	0.0377 (4)
H9A	0.3764	0.6890	0.3218	0.045*
H9B	0.4774	0.6180	0.3440	0.045*
C10	0.3805 (2)	0.64912 (11)	0.5113 (2)	0.0459 (5)
H10A	0.4723	0.6208	0.5610	0.055*
H10B	0.3969	0.6966	0.5414	0.055*
N11	0.20831 (18)	0.62338 (7)	0.52662 (14)	0.0314 (3)
C12	0.1228 (2)	0.59805 (8)	0.41189 (16)	0.0278 (3)
N13	−0.02640 (17)	0.56113 (7)	0.38209 (14)	0.0324 (3)
N14	−0.15702 (18)	0.57746 (7)	0.44638 (14)	0.0330 (3)
O15	−0.28030 (16)	0.53623 (7)	0.43068 (15)	0.0490 (4)
O16	−0.15984 (16)	0.63170 (7)	0.51017 (13)	0.0454 (3)
C17	0.1705 (2)	0.61047 (9)	0.65820 (16)	0.0331 (4)
H17A	0.2751	0.5946	0.7165	0.040*
H17B	0.0836	0.5743	0.6531	0.040*
C18	0.1042 (2)	0.67502 (9)	0.71615 (17)	0.0347 (4)
O19	0.13493 (19)	0.73282 (7)	0.68446 (14)	0.0505 (4)
O20	0.01369 (17)	0.65861 (6)	0.81063 (12)	0.0414 (3)
C21	−0.0639 (3)	0.71622 (11)	0.8720 (2)	0.0502 (5)
H21A	0.0215	0.7527	0.8939	0.060*
H21B	−0.0980	0.7007	0.9541	0.060*
C22	−0.2189 (3)	0.74367 (13)	0.7804 (3)	0.0658 (7)
H22A	−0.1828	0.7651	0.7049	0.099*
H22B	−0.2765	0.7772	0.8266	0.099*
H22C	−0.2975	0.7064	0.7505	0.099*

Atomic displacement parameters (Å²)

	U^{11}	U^{22}	U^{33}	U^{12}	U^{13}	U^{23}
Cl1	0.0666 (4)	0.0457 (3)	0.0760 (4)	0.0005 (3)	0.0250 (3)	−0.0183 (3)
N1	0.0488 (10)	0.0409 (9)	0.0396 (9)	0.0051 (7)	0.0028 (7)	−0.0019 (7)
C2	0.0482 (11)	0.0440 (11)	0.0273 (9)	0.0050 (9)	0.0036 (8)	−0.0007 (8)
C3	0.0283 (9)	0.0374 (10)	0.0288 (9)	−0.0033 (7)	0.0048 (7)	0.0013 (7)
C4	0.0338 (10)	0.0483 (11)	0.0296 (9)	−0.0034 (8)	0.0025 (7)	0.0026 (8)
C5	0.0402 (10)	0.0523 (12)	0.0296 (9)	−0.0105 (9)	0.0071 (8)	−0.0089 (8)
C6	0.0335 (9)	0.0387 (10)	0.0447 (11)	−0.0064 (8)	0.0089 (8)	−0.0093 (8)
C7	0.0343 (9)	0.0445 (10)	0.0296 (9)	0.0042 (8)	0.0056 (7)	0.0040 (8)

N8	0.0303 (7)	0.0393 (8)	0.0320 (8)	−0.0024 (6)	0.0065 (6)	−0.0023 (6)
C9	0.0274 (9)	0.0414 (10)	0.0443 (10)	−0.0021 (8)	0.0063 (8)	0.0026 (8)
C10	0.0345 (10)	0.0563 (13)	0.0459 (11)	−0.0121 (9)	0.0043 (8)	−0.0037 (9)
N11	0.0316 (7)	0.0311 (8)	0.0309 (7)	−0.0033 (6)	0.0042 (6)	−0.0024 (6)
C12	0.0297 (9)	0.0224 (8)	0.0310 (9)	0.0045 (6)	0.0048 (7)	0.0003 (6)
N13	0.0293 (7)	0.0340 (8)	0.0346 (8)	−0.0030 (6)	0.0074 (6)	−0.0059 (6)
N14	0.0296 (8)	0.0326 (8)	0.0355 (8)	0.0027 (6)	0.0026 (6)	0.0025 (6)
O15	0.0304 (7)	0.0429 (8)	0.0743 (10)	−0.0071 (6)	0.0114 (6)	−0.0031 (7)
O16	0.0414 (7)	0.0428 (8)	0.0529 (8)	0.0053 (6)	0.0104 (6)	−0.0131 (6)
C17	0.0380 (10)	0.0303 (9)	0.0293 (9)	0.0014 (7)	0.0014 (7)	0.0010 (7)
C18	0.0392 (10)	0.0339 (10)	0.0282 (9)	−0.0018 (8)	−0.0013 (7)	−0.0026 (7)
O19	0.0711 (10)	0.0294 (7)	0.0525 (8)	−0.0076 (7)	0.0151 (7)	−0.0041 (6)
O20	0.0550 (8)	0.0366 (7)	0.0345 (7)	0.0050 (6)	0.0127 (6)	−0.0010 (5)
C21	0.0605 (13)	0.0481 (12)	0.0431 (11)	0.0052 (10)	0.0125 (10)	−0.0134 (9)
C22	0.0676 (15)	0.0568 (15)	0.0716 (16)	0.0174 (12)	0.0091 (12)	−0.0097 (12)

Geometric parameters (Å, º)

Cl1—C6	1.7510 (19)	C10—H10A	0.9700
N1—C6	1.313 (2)	C10—H10B	0.9700
N1—C2	1.345 (2)	N11—C12	1.332 (2)
C2—C3	1.380 (2)	N11—C17	1.446 (2)
C2—H2	0.9300	C12—N13	1.354 (2)
C3—C4	1.388 (2)	N13—N14	1.3458 (19)
C3—C7	1.512 (2)	N14—O16	1.2367 (19)
C4—C5	1.377 (3)	N14—O15	1.2388 (18)
C4—H4	0.9300	C17—C18	1.512 (2)
C5—C6	1.376 (3)	C17—H17A	0.9700
C5—H5	0.9300	C17—H17B	0.9700
C7—N8	1.448 (2)	C18—O19	1.200 (2)
C7—H7A	0.9700	C18—O20	1.332 (2)
C7—H7B	0.9700	O20—C21	1.462 (2)
N8—C12	1.335 (2)	C21—C22	1.488 (3)
N8—C9	1.459 (2)	C21—H21A	0.9700
C9—C10	1.517 (3)	C21—H21B	0.9700
C9—H9A	0.9700	C22—H22A	0.9600
C9—H9B	0.9700	C22—H22B	0.9600
C10—N11	1.469 (2)	C22—H22C	0.9600
C6—N1—C2	116.47 (16)	C9—C10—H10B	111.1
N1—C2—C3	123.83 (16)	H10A—C10—H10B	109.0
N1—C2—H2	118.1	C12—N11—C17	126.65 (14)
C3—C2—H2	118.1	C12—N11—C10	110.86 (14)
C2—C3—C4	117.05 (16)	C17—N11—C10	120.06 (14)
C2—C3—C7	122.91 (15)	N11—C12—N8	110.38 (14)
C4—C3—C7	120.04 (15)	N11—C12—N13	131.84 (15)
C5—C4—C3	120.43 (17)	N8—C12—N13	117.42 (14)
C5—C4—H4	119.8	N14—N13—C12	117.67 (13)

C3—C4—H4	119.8	O16—N14—O15	121.84 (14)
C6—C5—C4	116.69 (16)	O16—N14—N13	122.81 (14)
C6—C5—H5	121.7	O15—N14—N13	115.21 (14)
C4—C5—H5	121.7	N11—C17—C18	111.17 (14)
N1—C6—C5	125.52 (17)	N11—C17—H17A	109.4
N1—C6—Cl1	115.38 (15)	C18—C17—H17A	109.4
C5—C6—Cl1	119.10 (14)	N11—C17—H17B	109.4
N8—C7—C3	113.43 (14)	C18—C17—H17B	109.4
N8—C7—H7A	108.9	H17A—C17—H17B	108.0
C3—C7—H7A	108.9	O19—C18—O20	125.08 (17)
N8—C7—H7B	108.9	O19—C18—C17	124.43 (17)
C3—C7—H7B	108.9	O20—C18—C17	110.43 (15)
H7A—C7—H7B	107.7	C18—O20—C21	116.27 (15)
C12—N8—C7	125.22 (14)	O20—C21—C22	110.91 (16)
C12—N8—C9	111.84 (14)	O20—C21—H21A	109.5
C7—N8—C9	122.24 (14)	C22—C21—H21A	109.5
N8—C9—C10	102.70 (14)	O20—C21—H21B	109.5
N8—C9—H9A	111.2	C22—C21—H21B	109.5
C10—C9—H9A	111.2	H21A—C21—H21B	108.0
N8—C9—H9B	111.2	C21—C22—H22A	109.5
C10—C9—H9B	111.2	C21—C22—H22B	109.5
H9A—C9—H9B	109.1	H22A—C22—H22B	109.5
N11—C10—C9	103.54 (14)	C21—C22—H22C	109.5
N11—C10—H10A	111.1	H22A—C22—H22C	109.5
C9—C10—H10A	111.1	H22B—C22—H22C	109.5
N11—C10—H10B	111.1		

C6—N1—C2—C3	−0.3 (3)	C17—N11—C12—N8	−165.04 (15)
N1—C2—C3—C4	0.9 (3)	C10—N11—C12—N8	−2.9 (2)
N1—C2—C3—C7	−179.01 (17)	C17—N11—C12—N13	7.7 (3)
C2—C3—C4—C5	−0.9 (3)	C10—N11—C12—N13	169.83 (18)
C7—C3—C4—C5	179.07 (16)	C7—N8—C12—N11	−173.32 (15)
C3—C4—C5—C6	0.3 (3)	C9—N8—C12—N11	−2.8 (2)
C2—N1—C6—C5	−0.4 (3)	C7—N8—C12—N13	12.8 (2)
C2—N1—C6—Cl1	178.84 (13)	C9—N8—C12—N13	−176.68 (14)
C4—C5—C6—N1	0.5 (3)	N11—C12—N13—N14	38.2 (3)
C4—C5—C6—Cl1	−178.79 (13)	N8—C12—N13—N14	−149.51 (15)
C2—C3—C7—N8	13.8 (2)	C12—N13—N14—O16	14.3 (2)
C4—C3—C7—N8	−166.19 (15)	C12—N13—N14—O15	−169.90 (15)
C3—C7—N8—C12	−107.54 (18)	C12—N11—C17—C18	−111.85 (18)
C3—C7—N8—C9	82.8 (2)	C10—N11—C17—C18	87.50 (19)
C12—N8—C9—C10	6.9 (2)	N11—C17—C18—O19	−24.7 (2)
C7—N8—C9—C10	177.74 (16)	N11—C17—C18—O20	157.88 (14)
N8—C9—C10—N11	−7.84 (19)	O19—C18—O20—C21	4.8 (3)
C9—C10—N11—C12	7.0 (2)	C17—C18—O20—C21	−177.77 (14)
C9—C10—N11—C17	170.43 (15)	C18—O20—C21—C22	74.9 (2)

Hydrogen-bond geometry (Å, º)

D—H···*A*	*D*—H	H···*A*	*D*···*A*	*D*—H···*A*
C9—H9*B*···O15^{i}	0.97	2.50	3.358 (2)	147
C17—H17*A*···N1^{ii}	0.97	2.57	3.509 (2)	163

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