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2-*tert*-Butyl-6-[(4-chloro-2-nitrophenyl)-diazenyl]-4-methylphenol

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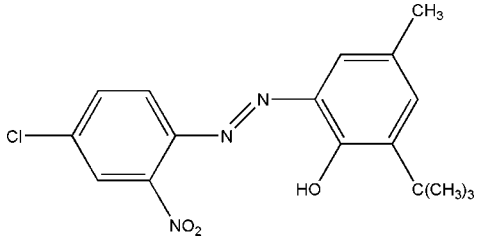
Key indicators: single-crystal X-ray study; T = 296 K; mean σ(C–C) = 0.003 Å; disorder in main residue; R factor = 0.044; wR factor = 0.141; data-to-parameter ratio = 16.3.

In the title compound, $C_{17}H_{18}ClN_3O_3$, the dihedral angle between the planes of the two benzene rings is 1.03 (7)°. The overall conformation of the molecule is influenced, in part, by electron delocalization and by an intramolecular bifurcated O—H···(O,N) hydrogen bonds. The O atoms of the nitro group, one of which serves as an H bond acceptor, are disordered over two sets of sites with refined occupancies of 0.56 (3) and 0.44 (3).

Related literature

For benzotriazoles as UV absorbers and their applications in industry, see: Ravichandran *et al.* (2002). *N*-oxides are a key type intermediates in the synthesis of benzotriazoles, see: Wen *et al.* (2006); Crawford (1999). For the use of green synthetic methods to obtain intermediates, see: Tanaka & Toda (2000).



Experimental

Crystal data

$C_{17}H_{18}ClN_3O_3$ M_r = 347.79
Monoclinic, $P2_1/c$
a = 14.578 (4) Å
b = 7.0616 (19) Å
c = 17.043 (5) Å
β = 101.233 (3)°
V = 1720.9 (8) Å^3
Z = 4
Mo $K\alpha$ radiation
μ = 0.24 mm^{-1}
T = 296 K
0.31 × 0.18 × 0.16 mm

Data collection

Bruker APEXII CCD diffractometer
Absorption correction: multi-scan (*SADABS*; Sheldrick, 1996)
T_{min} = 0.930, T_{max} = 0.963
14642 measured reflections
3927 independent reflections
2563 reflections with $I > 2\sigma(I)$
R_{int} = 0.033

Refinement

$R[F^2 > 2\sigma(F^2)]$ = 0.044
$wR(F^2)$ = 0.141
S = 1.03
3927 reflections
241 parameters
H-atom parameters constrained
$\Delta\rho_{max}$ = 0.22 e Å^{-3}
$\Delta\rho_{min}$ = −0.26 e Å^{-3}

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

D—H···A	D—H	H···A	D···A	D—H···A
O3—H3···O1′	0.82	2.28	2.933 (7)	136
O3—H3···O1	0.82	2.50	3.142 (12)	136
O3—H3···N2	0.82	1.84	2.553 (2)	145

Data collection: *APEX2* (Bruker, 2006); cell refinement: *SAINT* (Bruker, 2006); data reduction: *SAINT*; program(s) used to solve structure: *SHELXS97* (Sheldrick, 2008); program(s) used to refine structure: *SHELXL97* (Sheldrick, 2008); molecular graphics: *SHELXTL* (Sheldrick, 2008); software used to prepare material for publication: *SHELXTL*.

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Supplementary data and figures for this paper are available from the IUCr electronic archives (Reference: LH2824).

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supplementary materials

Acta Cryst. (2009). E**65**, o1520 [**doi:10.1107/S1600536809019631**]

2-*tert*-Butyl-6-[(4-chloro-2-nitrophenyl)diazenyl]-4-methylphenol

H.-L. Wen, X.-Q. Wu and B.-W. Lai

Comment

Benzotriazoles play an important role as a class of UV absorbers and have promising industrial applications (Ravichandran *et al.*, 2002). N-oxides are a key type intermediates in the synthesis of benzotriazoles (Wen *et al.*, 2006; Crawford, 1999) and the title compound is an important intermediate in the synthesis of 2-(2'-Hydroxy-3'-tert-butyl-5'-methylphenyl)-5-chloro benzotriazole (UV 326), a good ultraviolet absorber. Due to the growing awareness of environmental protection, the demand for clean and 'green' (i.e solvent free) chemical syntheses has been growing, so using these synthetic methods to form intermediates have received attention (Tanaka & Toda, 2000). Herein we report a 'green' synthetic method and the crystal structure of the title compound. In the title moleclue (Fig .1) the dihedral angle between the two benzene rings is 1.03 (7)°. The overall conformation of the molecule is influenced, in part, by electron delocalization and by intramolecular O—H···O and O—H···N hydrogen bonds.

Experimental

The title compound was synthesized *via* the solid phase reaction of 4-chloro-2-nitroaniline and 2-*tert*-butyl-4-substituted phenol at room temperature. After intensive grinding a mixture of 4-chloro-2-nitrobenzenamine 1.72 g (10 mmol), 2-*tert*-butyl-4-methylphenol 1.72 g (10.5 mmol), $NaNO_2$ 0.69 g (10 mmol), and $KHSO_4$ 1.36 g (10 mmol) in a mortar for 15 min at 293 K, the product was washed with hot water. A few purple crystals suitable for X-ray diffraction analysis were obtained upon recrystallization in ethanol after several days (m. p. 445–446 K), which gave the product in 93% yield and higher than 99% purity (by HPLC).

Refinement

All H atoms were included in calculated positions with O—H = 0.82Å; *C*—*H*(methyl) = 0.96 Å, C—H(aromatic) = 0.93 Å, and U_{iso}(H) = 1.5U_{eq}(C_{methyl},O) and U_{iso}(H) = 1.2U_{eq}(C) for aromatic H atoms. The O atoms of the nitro group are disordered over two sites with refined occupancies of 0.56 (3) and 0.44 (3).

Figures

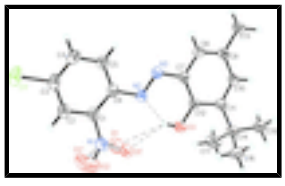


Fig. 1. : The molecular structure of the title compound, showing 30% probability displacement ellipsoids. The minor comonent of disorder is shown with open bonds and hydrogen bonds are shown with dashed lines.

2-*tert*-Butyl-6-[(4-chloro-2-nitrophenyl)diazenyl]-4-methylphenol

Crystal data

$C_{17}H_{18}ClN_3O_3$	F_{000} = 728
M_r = 347.79	D_x = 1.342 Mg m^{-3}
Monoclinic, $P2_1/c$	Mo $K\alpha$ radiation λ = 0.71073 Å
Hall symbol: -P 2ybc	Cell parameters from 4044 reflections
a = 14.578 (4) Å	θ = 2.6–27.2º
b = 7.0616 (19) Å	μ = 0.24 mm^{-1}
c = 17.043 (5) Å	T = 296 K
β = 101.233 (3)º	Block, purple
V = 1720.9 (8) Å^3	0.31 × 0.18 × 0.16 mm
Z = 4	

Data collection

Bruker APEXII CCD diffractometer	3927 independent reflections
Radiation source: fine-focus sealed tube	2563 reflections with $I > 2\sigma(I)$
Monochromator: graphite	R_{int} = 0.033
T = 296 K	θ_{max} = 27.5º
φ and ω scans	θ_{min} = 2.4º
Absorption correction: multi-scan (SADABS; Sheldrick, 1996)	h = −18→18
T_{min} = 0.930, T_{max} = 0.963	k = −9→8
14642 measured reflections	l = −22→22

Refinement

Refinement on F^2	Secondary atom site location: difference Fourier map
Least-squares matrix: full	Hydrogen site location: inferred from neighbouring sites
$R[F^2 > 2\sigma(F^2)]$ = 0.044	H-atom parameters constrained
$wR(F^2)$ = 0.141	$w = 1/[\sigma^2(F_o^2) + (0.0591P)^2 + 0.5672P]$ where $P = (F_o^2 + 2F_c^2)/3$
S = 1.03	$(\Delta/\sigma)_{max}$ = 0.002
3927 reflections	$\Delta\rho_{max}$ = 0.22 e Å^{-3}
241 parameters	$\Delta\rho_{min}$ = −0.26 e Å^{-3}
Primary atom site location: structure-invariant direct methods	Extinction correction: none

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 (Å^2)

	x	*y*	*z*	U_{iso}*/U_{eq}	Occ. (<1)
Cl1	1.37753 (4)	0.52156 (11)	0.44085 (5)	0.0871 (3)	
O1	0.9625 (4)	0.559 (3)	0.2905 (6)	0.109 (4)	0.56 (3)
O2	1.0758 (7)	0.659 (2)	0.2334 (6)	0.115 (4)	0.56 (3)
O3	0.81516 (9)	0.7932 (3)	0.36241 (8)	0.0666 (5)	
H3	0.8703	0.7643	0.3663	0.100*	
O1'	0.9670 (8)	0.689 (4)	0.2804 (4)	0.100 (5)	0.44 (3)
O2'	1.0687 (12)	0.545 (5)	0.2372 (10)	0.156 (8)	0.44 (3)
N1	1.04252 (15)	0.6167 (4)	0.29115 (12)	0.0765 (6)	
N2	0.98233 (10)	0.7293 (2)	0.43581 (9)	0.0475 (4)	
N3	0.96277 (10)	0.7767 (2)	0.50347 (9)	0.0468 (4)	
C1	1.10633 (13)	0.6202 (3)	0.36932 (12)	0.0528 (5)	
C2	1.19824 (14)	0.5718 (3)	0.36947 (14)	0.0597 (5)	
H2	1.2171	0.5395	0.3221	0.072*	
C3	1.26084 (14)	0.5723 (3)	0.44035 (15)	0.0599 (6)	
C4	1.23301 (14)	0.6162 (3)	0.51114 (14)	0.0606 (5)	
H4	1.2760	0.6122	0.5592	0.073*	
C5	1.14154 (13)	0.6660 (3)	0.51035 (12)	0.0537 (5)	
H5	1.1233	0.6963	0.5582	0.064*	
C6	1.07582 (12)	0.6716 (3)	0.43917 (11)	0.0468 (4)	
C7	0.87202 (12)	0.8321 (3)	0.50305 (10)	0.0425 (4)	
C8	0.85301 (13)	0.8758 (3)	0.57888 (11)	0.0465 (4)	
H8	0.9010	0.8707	0.6236	0.056*	
C9	0.76528 (13)	0.9255 (3)	0.58763 (10)	0.0473 (4)	
C10	0.69528 (13)	0.9379 (3)	0.51815 (11)	0.0485 (5)	
H10	0.6358	0.9748	0.5242	0.058*	
C11	0.70815 (12)	0.8993 (3)	0.44168 (10)	0.0485 (5)	
C12	0.79931 (12)	0.8412 (3)	0.43353 (10)	0.0465 (4)	
C13	0.74126 (16)	0.9610 (4)	0.66868 (11)	0.0646 (6)	
H13A	0.7066	0.8553	0.6831	0.097*	
H13B	0.7042	1.0738	0.6666	0.097*	
H13C	0.7978	0.9763	0.7078	0.097*	
C14	0.62867 (14)	0.9137 (4)	0.36802 (12)	0.0658 (6)	
C15	0.53856 (16)	0.9904 (5)	0.39088 (14)	0.0868 (9)	

H15A	0.5192	0.9069	0.4290	0.130*
H15B	0.4901	0.9979	0.3439	0.130*
H15C	0.5502	1.1142	0.4139	0.130*
C16	0.65620 (18)	1.0504 (5)	0.30642 (14)	0.0959 (10)
H16A	0.6645	1.1753	0.3290	0.144*
H16B	0.6077	1.0528	0.2594	0.144*
H16C	0.7136	1.0087	0.2924	0.144*
C17	0.60711 (17)	0.7157 (5)	0.33203 (15)	0.0931 (10)
H17A	0.6629	0.6623	0.3190	0.140*
H17B	0.5598	0.7250	0.2844	0.140*
H17C	0.5852	0.6360	0.3701	0.140*

Atomic displacement parameters (Å^2)

	U^{11}	U^{22}	U^{33}	U^{12}	U^{13}	U^{23}
Cl1	0.0412 (3)	0.1022 (5)	0.1217 (6)	0.0083 (3)	0.0251 (3)	0.0142 (4)
O1	0.052 (2)	0.186 (11)	0.084 (3)	−0.001 (4)	−0.001 (2)	−0.047 (5)
O2	0.105 (4)	0.185 (9)	0.056 (3)	−0.001 (5)	0.016 (3)	0.012 (4)
O3	0.0420 (7)	0.1165 (13)	0.0417 (7)	0.0091 (8)	0.0094 (6)	−0.0118 (8)
O1'	0.070 (4)	0.173 (13)	0.055 (3)	0.043 (5)	0.007 (2)	−0.013 (4)
O2'	0.115 (7)	0.255 (18)	0.091 (7)	0.062 (10)	0.007 (5)	−0.080 (9)
N1	0.0612 (13)	0.1073 (17)	0.0606 (12)	0.0158 (13)	0.0106 (10)	−0.0208 (12)
N2	0.0388 (8)	0.0553 (10)	0.0491 (9)	−0.0003 (7)	0.0102 (7)	−0.0025 (7)
N3	0.0412 (8)	0.0512 (9)	0.0478 (9)	−0.0017 (7)	0.0083 (7)	−0.0007 (7)
C1	0.0465 (11)	0.0550 (12)	0.0571 (12)	0.0003 (9)	0.0106 (9)	−0.0026 (9)
C2	0.0503 (12)	0.0634 (14)	0.0702 (14)	0.0041 (10)	0.0232 (11)	0.0006 (11)
C3	0.0385 (10)	0.0570 (13)	0.0869 (16)	0.0007 (9)	0.0186 (11)	0.0079 (11)
C4	0.0429 (11)	0.0655 (13)	0.0699 (14)	−0.0021 (9)	0.0021 (10)	0.0061 (11)
C5	0.0455 (11)	0.0588 (12)	0.0561 (12)	−0.0017 (9)	0.0084 (9)	0.0023 (9)
C6	0.0394 (9)	0.0460 (10)	0.0554 (11)	−0.0029 (8)	0.0104 (8)	−0.0006 (8)
C7	0.0374 (9)	0.0472 (10)	0.0425 (9)	−0.0019 (7)	0.0072 (7)	−0.0006 (8)
C8	0.0469 (10)	0.0531 (11)	0.0378 (9)	−0.0022 (8)	0.0038 (8)	0.0008 (8)
C9	0.0515 (11)	0.0542 (11)	0.0380 (9)	−0.0012 (9)	0.0131 (8)	0.0004 (8)
C10	0.0418 (10)	0.0617 (12)	0.0441 (10)	0.0039 (8)	0.0139 (8)	0.0015 (9)
C11	0.0402 (10)	0.0668 (13)	0.0384 (9)	−0.0008 (9)	0.0074 (7)	0.0002 (9)
C12	0.0418 (10)	0.0606 (12)	0.0381 (9)	0.0003 (8)	0.0102 (8)	−0.0032 (8)
C13	0.0651 (13)	0.0909 (17)	0.0408 (10)	0.0048 (12)	0.0178 (10)	−0.0003 (11)
C14	0.0399 (11)	0.115 (2)	0.0415 (10)	0.0116 (11)	0.0065 (8)	−0.0021 (11)
C15	0.0470 (12)	0.156 (3)	0.0552 (13)	0.0292 (15)	0.0053 (10)	0.0027 (15)
C16	0.0679 (16)	0.170 (3)	0.0488 (13)	0.0225 (18)	0.0092 (11)	0.0315 (16)
C17	0.0536 (14)	0.154 (3)	0.0663 (15)	−0.0061 (16)	−0.0019 (11)	−0.0380 (17)

Geometric parameters (Å, °)

Cl1—C3	1.737 (2)	C8—H8	0.9300
O1—N1	1.233 (8)	C9—C10	1.407 (3)
O2—N1	1.217 (8)	C9—C13	1.511 (2)
O3—C12	1.322 (2)	C10—C11	1.379 (2)
O3—H3	0.8200	C10—H10	0.9300

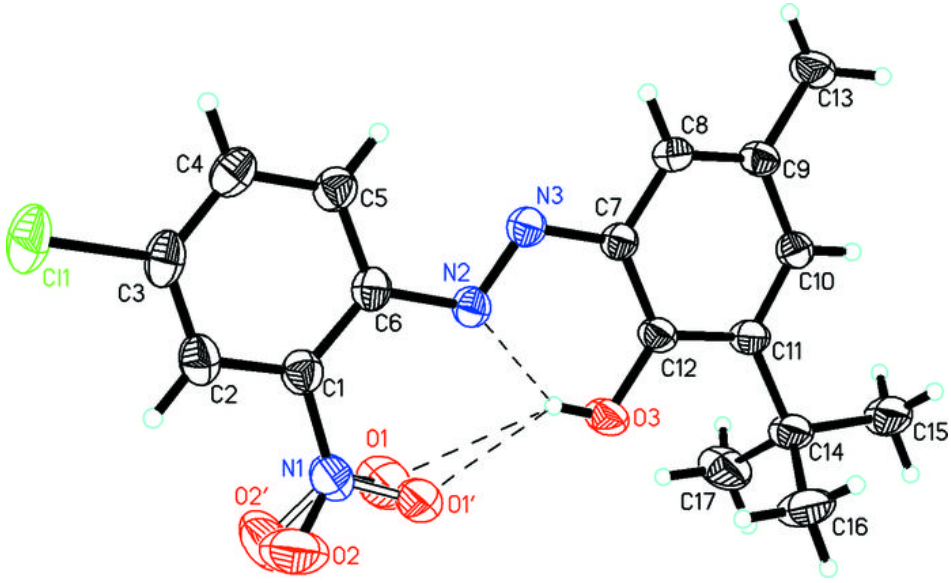
O1'—N1	1.195 (7)	C11—C12	1.423 (2)
O2'—N1	1.179 (10)	C11—C14	1.537 (3)
N1—C1	1.469 (3)	C13—H13A	0.9600
N2—N3	1.285 (2)	C13—H13B	0.9600
N2—C6	1.413 (2)	C13—H13C	0.9600
N3—C7	1.378 (2)	C14—C17	1.534 (4)
C1—C2	1.382 (3)	C14—C16	1.536 (4)
C1—C6	1.397 (3)	C14—C15	1.540 (3)
C2—C3	1.365 (3)	C15—H15A	0.9600
C2—H2	0.9300	C15—H15B	0.9600
C3—C4	1.381 (3)	C15—H15C	0.9600
C4—C5	1.376 (3)	C16—H16A	0.9600
C4—H4	0.9300	C16—H16B	0.9600
C5—C6	1.392 (3)	C16—H16C	0.9600
C5—H5	0.9300	C17—H17A	0.9600
C7—C8	1.408 (2)	C17—H17B	0.9600
C7—C12	1.429 (3)	C17—H17C	0.9600
C8—C9	1.362 (3)		
C12—O3—H3	109.5	C9—C10—H10	117.5
O2'—N1—O1'	119.6 (10)	C10—C11—C12	116.78 (16)
O2—N1—O1	126.8 (8)	C10—C11—C14	122.62 (17)
O2'—N1—C1	118.1 (7)	C12—C11—C14	120.59 (16)
O1'—N1—C1	122.2 (5)	O3—C12—C11	119.79 (16)
O2—N1—C1	116.6 (5)	O3—C12—C7	120.99 (16)
O1—N1—C1	116.4 (6)	C11—C12—C7	119.20 (15)
N3—N2—C6	114.81 (15)	C9—C13—H13A	109.5
N2—N3—C7	116.74 (15)	C9—C13—H13B	109.5
C2—C1—C6	122.12 (19)	H13A—C13—H13B	109.5
C2—C1—N1	116.10 (18)	C9—C13—H13C	109.5
C6—C1—N1	121.77 (17)	H13A—C13—H13C	109.5
C3—C2—C1	118.8 (2)	H13B—C13—H13C	109.5
C3—C2—H2	120.6	C17—C14—C16	111.1 (2)
C1—C2—H2	120.6	C17—C14—C11	109.2 (2)
C2—C3—C4	120.94 (19)	C16—C14—C11	110.12 (19)
C2—C3—Cl1	119.29 (17)	C17—C14—C15	107.7 (2)
C4—C3—Cl1	119.75 (18)	C16—C14—C15	107.5 (2)
C5—C4—C3	119.8 (2)	C11—C14—C15	111.17 (17)
C5—C4—H4	120.1	C14—C15—H15A	109.5
C3—C4—H4	120.1	C14—C15—H15B	109.5
C4—C5—C6	121.11 (19)	H15A—C15—H15B	109.5
C4—C5—H5	119.4	C14—C15—H15C	109.5
C6—C5—H5	119.4	H15A—C15—H15C	109.5
C5—C6—C1	117.11 (17)	H15B—C15—H15C	109.5
C5—C6—N2	122.56 (17)	C14—C16—H16A	109.5
C1—C6—N2	120.32 (17)	C14—C16—H16B	109.5
N3—C7—C8	114.70 (16)	H16A—C16—H16B	109.5
N3—C7—C12	124.97 (16)	C14—C16—H16C	109.5
C8—C7—C12	120.30 (16)	H16A—C16—H16C	109.5
C9—C8—C7	120.93 (17)	H16B—C16—H16C	109.5

C9—C8—H8	119.5	C14—C17—H17A	109.5
C7—C8—H8	119.5	C14—C17—H17B	109.5
C8—C9—C10	117.79 (16)	H17A—C17—H17B	109.5
C8—C9—C13	122.19 (17)	C14—C17—H17C	109.5
C10—C9—C13	120.00 (17)	H17A—C17—H17C	109.5
C11—C10—C9	124.95 (17)	H17B—C17—H17C	109.5
C11—C10—H10	117.5		
C6—N2—N3—C7	−179.36 (15)	N2—N3—C7—C8	177.70 (16)
O2'—N1—C1—C2	12 (2)	N2—N3—C7—C12	−0.3 (3)
O1'—N1—C1—C2	−165.3 (16)	N3—C7—C8—C9	−177.44 (18)
O2—N1—C1—C2	−32.9 (10)	C12—C7—C8—C9	0.7 (3)
O1—N1—C1—C2	142.5 (11)	C7—C8—C9—C10	−2.1 (3)
O2'—N1—C1—C6	−168 (2)	C7—C8—C9—C13	176.00 (19)
O1'—N1—C1—C6	14.0 (16)	C8—C9—C10—C11	1.5 (3)
O2—N1—C1—C6	146.4 (10)	C13—C9—C10—C11	−176.6 (2)
O1—N1—C1—C6	−38.2 (11)	C9—C10—C11—C12	0.6 (3)
C6—C1—C2—C3	0.7 (3)	C9—C10—C11—C14	179.5 (2)
N1—C1—C2—C3	−179.9 (2)	C10—C11—C12—O3	176.69 (19)
C1—C2—C3—C4	1.4 (3)	C14—C11—C12—O3	−2.2 (3)
C1—C2—C3—Cl1	−177.30 (16)	C10—C11—C12—C7	−2.0 (3)
C2—C3—C4—C5	−2.0 (3)	C14—C11—C12—C7	179.05 (19)
Cl1—C3—C4—C5	176.67 (17)	N3—C7—C12—O3	0.7 (3)
C3—C4—C5—C6	0.5 (3)	C8—C7—C12—O3	−177.22 (18)
C4—C5—C6—C1	1.5 (3)	N3—C7—C12—C11	179.40 (18)
C4—C5—C6—N2	−177.41 (19)	C8—C7—C12—C11	1.5 (3)
C2—C1—C6—C5	−2.1 (3)	C10—C11—C14—C17	−114.1 (2)
N1—C1—C6—C5	178.6 (2)	C12—C11—C14—C17	64.7 (3)
C2—C1—C6—N2	176.80 (19)	C10—C11—C14—C16	123.6 (2)
N1—C1—C6—N2	−2.5 (3)	C12—C11—C14—C16	−57.5 (3)
N3—N2—C6—C5	0.2 (3)	C10—C11—C14—C15	4.6 (3)
N3—N2—C6—C1	−178.68 (17)	C12—C11—C14—C15	−176.6 (2)

Hydrogen-bond geometry (Å, °)

D—H···*A*	*D*—H	H···*A*	*D*···*A*	*D*—H···*A*
O3—H3···O1'	0.82	2.28	2.933 (7)	136
O3—H3···O1	0.82	2.50	3.142 (12)	136
O3—H3···N2	0.82	1.84	2.553 (2)	145

Fig. 1


C13
C8
C9
C4
C5
N3
C7
Cl1
C3
N2
C10
C6
C11
C12
C2
C1
O3
O1
C15
N1
C14
O2′
O1′
C17
C16
O2