

2,10-Dihydroxy-13-methyl-13-azatetracyclo[9.3.1.0^{2,10}.0^{3,8}]pentadeca-3(8),4,6-triene-9,15-dione

J. Suresh,^a U. C. Nithya,^a R. Suresh Kumar,^b S. Perumal^b and P. L. Nilantha Lakshman^{c*}

^aDepartment of Physics, The Madura College, Madurai 625 011, India, ^bSchool of Chemistry, Madurai Kamaraj University, Madurai 625 021, India, and ^cDepartment of Food Science and Technology, Faculty of Agriculture, University of Ruhuna, Mapalana, Kamburupitiya 81100, Sri Lanka
Correspondence e-mail: nilanthalakshman@yahoo.co.uk

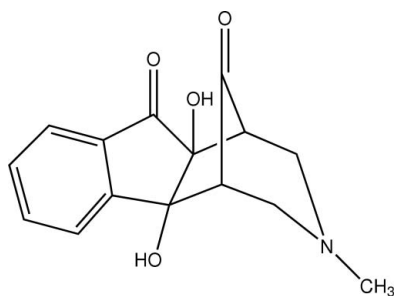
Received 22 April 2009; accepted 28 April 2009

Key indicators: single-crystal X-ray study; $T = 293$ K; mean $\sigma(\text{C}-\text{C}) = 0.002$ Å; R factor = 0.037; wR factor = 0.097; data-to-parameter ratio = 12.6.

In the title compound, $\text{C}_{15}\text{H}_{15}\text{NO}_4$, the *n*-methylpiperidone ring adopts a chair conformation and both five-membered rings adopt a twist conformation. An intramolecular $\text{O}-\text{H}\cdots\text{O}$ hydrogen bond is observed. Inversion-related molecules are linked into $R_2^2(10)$ dimers by pairs of $\text{O}-\text{H}\cdots\text{O}$ hydrogen bonds. The crystal structure is further stabilized by $\text{C}-\text{H}\cdots\text{O}$ hydrogen bonds.

Related literature

For the biological activity of piperidine compounds, see: Watson *et al.* (2000). For ring conformation details, see: Cremer & Pople (1975).



Experimental

Crystal data

$\text{C}_{15}\text{H}_{15}\text{NO}_4$

$M_r = 273.28$

Triclinic, $P\bar{1}$
 $a = 7.5616$ (7) Å
 $b = 8.9033$ (8) Å
 $c = 10.8091$ (11) Å
 $\alpha = 72.764$ (11)°
 $\beta = 80.486$ (12)°
 $\gamma = 72.369$ (11)°

$V = 660.09$ (11) Å³
 $Z = 2$
Mo $K\alpha$ radiation
 $\mu = 0.10$ mm⁻¹
 $T = 293$ K
 $0.18 \times 0.15 \times 0.11$ mm

Data collection

Nonius MACH-3 diffractometer
Absorption correction: ψ scan
(North *et al.*, 1968)
 $T_{\min} = 0.982$, $T_{\max} = 0.989$
2890 measured reflections
2316 independent reflections

2025 reflections with $I > 2\sigma(I)$
 $R_{\text{int}} = 0.010$
2 standard reflections
frequency: 60 min
intensity decay: none

Refinement

$R[F^2 > 2\sigma(F^2)] = 0.037$
 $wR(F^2) = 0.097$
 $S = 1.11$
2316 reflections

184 parameters
H-atom parameters constrained
 $\Delta\rho_{\text{max}} = 0.22$ e Å⁻³
 $\Delta\rho_{\text{min}} = -0.23$ e Å⁻³

Table 1

Hydrogen-bond geometry (Å, °).

$D-H\cdots A$	$D-H$	$H\cdots A$	$D\cdots A$	$D-H\cdots A$
$\text{O3}-\text{H3}\cdots\text{O2}$	0.82	2.11	2.6080 (15)	119
$\text{O2}-\text{H2}\cdots\text{O1}^{\text{i}}$	0.82	1.90	2.7168 (15)	175
$\text{C2}-\text{H2A}\cdots\text{O1}^{\text{i}}$	0.98	2.59	3.3778 (18)	137
$\text{C18}-\text{H18A}\cdots\text{O1}^{\text{ii}}$	0.96	2.57	3.432 (2)	149
$\text{C5}-\text{H5}\cdots\text{O3}^{\text{iii}}$	0.98	2.46	3.4381 (18)	173

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

Data collection: *CAD-4 EXPRESS* (Enraf-Nonius, 1994); cell refinement: *CAD-4 EXPRESS*; data reduction: *XCAD4* (Harms & Wocadlo, 1996); program(s) used to solve structure: *SHELXS97* (Sheldrick, 2008); program(s) used to refine structure: *SHELXL97* (Sheldrick, 2008); molecular graphics: *PLATON* (Spek, 2009); software used to prepare material for publication: *SHELXL97*.

JS and UCN thank the Management of the Madura College, Madurai, for their constant support.

Supplementary data and figures for this paper are available from the IUCr electronic archives (Reference: CI2789).

References

- Cremer, D. & Pople, J. A. (1975). *J. Am. Chem. Soc.* **97**, 1354–1358.
Enraf-Nonius (1994). *CAD-4 EXPRESS*. Enraf-Nonius, Delft, The Netherlands.
Harms, K. & Wocadlo, S. (1996). *XCAD4*. University of Marburg, Germany.
North, A. C. T., Phillips, D. C. & Mathews, F. S. (1968). *Acta Cryst.* **A24**, 351–359.
Sheldrick, G. M. (2008). *Acta Cryst.* **A64**, 112–122.
Spek, A. L. (2009). *Acta Cryst.* **D65**, 148–155.
Watson, P. S., Jiang, B. & Scott, B. (2000). *Org. Lett.* **2**, 3679–3681.

supplementary materials

Acta Cryst. (2009). E65, o1216 [doi:10.1107/S1600536809015785]

2,10-Dihydroxy-13-methyl-13-azatetracyclo[9.3.1.0^{2,10}.0^{3,8}]pentadeca-3(8),4,6-triene-9,15-dione

J. Suresh, U. C. Nithya, R. S. Kumar, S. Perumal and P. L. N. Lakshman

Comment

The piperidine ring is a distinct structural feature of a variety of alkaloid natural products and drug candidates. Watson *et al.* (2000) observed that during the past decade there were thousands of piperidine compounds mentioned in clinical and preclinical studies. Piperidinones, though relatively less prominent, have also been regarded as precursors of a host of biologically active compounds and natural alkaloids, prior to their conversion to piperidines. Ninhydrin is a chemical used to detect ammonia or primary and secondary amines.

In the molecule of the title compound, (Fig. 1), the six-membered ring A (N1/C4/C5/C1/C2/C3), and the five membered rings B (C1/C2/C7/C6/C5) and C (C6-C10) are not planar. Rings B and C both adopt twist conformations, as indicated by Cremer & Pople (1975) puckering parameters $Q = 0.455$ (2) Å and $\Phi = 160.4$ (2)° for ring B, and $Q = 0.149$ (2) Å and $\Phi = 21.6$ (6)° for ring C. Ring A adopts a chair conformation.

In the crystal structure, the molecules are linked to form dimers by intermolecular O—H···O hydrogen bonds (Table 1), generating a graph set motif of $R_2^2(10)$ (Fig.2). In addition, the structure is stabilized by C—H···O and van der Waals interactions.

Experimental

A mixture of 1-methyl-4-piperidinone (0.2 g, 0.002 mol), ninhydrin (0.315 g, 0.002 mol) and sarcosine (0.156 g, 0.002 mol) in methanol (30 ml) was refluxed in a water bath for 10 h. After completion of the reaction as monitored by TLC, the excess solvent was removed under vacuum and the residue was subjected to flash column chromatography using petroleum ether-ethyl acetate mixture (8:2 v/v) as eluent to obtain crystals of title compound in 8% yield along with a other product (yield 13%, m.p. 447–448 K).

Refinement

The H atoms were placed in calculated positions and allowed to ride on their carrier atoms, with C-H = 0.93–0.98 Å, O-H = 0.82 Å and $U_{\text{iso}}(\text{H}) = 1.2U_{\text{eq}}(\text{C})$ and $1.5U_{\text{eq}}(\text{C}_{\text{methyl}}, \text{O})$.

Figures

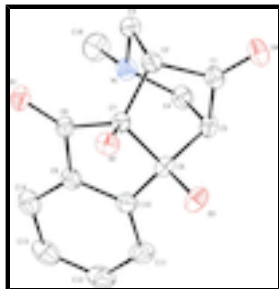


Fig. 1. The molecular structure of the title compound, showing 30% probability displacement ellipsoids and the atom-numbering scheme. H atoms have been omitted for clarity.

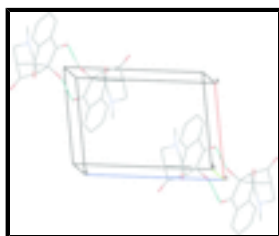


Fig. 2. Partial packing view down the *b* axis. Hydrogen bonds are shown as dashed lines.

2,10-Dihydroxy-13-methyl-13-azatetracyclo[9.3.1.0^{2,10}.0^{3,8}]pentadeca- 3(8),4,6-triene-9,15-dione

Crystal data

$C_{15}H_{15}NO_4$

$M_r = 273.28$

Triclinic, $P\bar{1}$

Hall symbol: -P 1

$a = 7.5616$ (7) Å

$b = 8.9033$ (8) Å

$c = 10.8091$ (11) Å

$\alpha = 72.764$ (11)°

$\beta = 80.486$ (12)°

$\gamma = 72.369$ (11)°

$V = 660.09$ (11) Å³

$Z = 2$

$F_{000} = 288$

$D_x = 1.375$ Mg m⁻³

Mo $K\alpha$ radiation

$\lambda = 0.71073$ Å

Cell parameters from 25 reflections

$\theta = 2-25^\circ$

$\mu = 0.10$ mm⁻¹

$T = 293$ K

Block, colourless

$0.18 \times 0.15 \times 0.11$ mm

Data collection

Nonius MACH-3
diffractometer

Radiation source: fine-focus sealed tube

Monochromator: graphite

$T = 293$ K

$\omega-2\theta$ scans

Absorption correction: ψ scan
(North *et al.*, 1968)

$T_{\min} = 0.982$, $T_{\max} = 0.989$

2890 measured reflections

$R_{\text{int}} = 0.010$

$\theta_{\text{max}} = 25.0^\circ$

$\theta_{\text{min}} = 2.5^\circ$

$h = -1 \rightarrow 8$

$k = -10 \rightarrow 10$

$l = -12 \rightarrow 12$

2 standard reflections

every 60 min

2316 independent reflections
 2025 reflections with $I > 2\sigma(I)$

intensity decay: none

Refinement

Refinement on F^2	Secondary atom site location: difference Fourier map
Least-squares matrix: full	Hydrogen site location: inferred from neighbouring sites
$R[F^2 > 2\sigma(F^2)] = 0.037$	H-atom parameters constrained
$wR(F^2) = 0.097$	$w = 1/[\sigma^2(F_o^2) + (0.0454P)^2 + 0.1839P]$
$S = 1.11$	where $P = (F_o^2 + 2F_c^2)/3$
2316 reflections	$(\Delta/\sigma)_{\max} = 0.001$
184 parameters	$\Delta\rho_{\max} = 0.22 \text{ e } \text{\AA}^{-3}$
Primary atom site location: structure-invariant direct methods	$\Delta\rho_{\min} = -0.23 \text{ e } \text{\AA}^{-3}$
	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 (\AA^2)

	x	y	z	$U_{\text{iso}}^*/U_{\text{eq}}$
O1	0.29809 (15)	0.41843 (13)	0.00498 (10)	0.0446 (3)
O2	-0.04087 (14)	0.29519 (13)	0.08496 (10)	0.0421 (3)
H2	-0.1139	0.3846	0.0581	0.063*
O3	-0.02584 (16)	0.06248 (14)	0.30007 (11)	0.0499 (3)
H3	-0.0724	0.0852	0.2313	0.075*
N1	0.29935 (17)	0.38335 (15)	0.28995 (12)	0.0388 (3)
O4	-0.19177 (16)	0.40313 (16)	0.45054 (11)	0.0538 (3)
C8	0.2679 (2)	0.30815 (17)	0.09849 (13)	0.0353 (3)
C4	0.2670 (2)	0.26334 (19)	0.41090 (14)	0.0403 (3)
H4A	0.2445	0.3121	0.4832	0.048*
H4B	0.3768	0.1712	0.4252	0.048*
C10	0.3067 (2)	0.04870 (17)	0.24458 (14)	0.0372 (3)
C7	0.07412 (19)	0.30768 (16)	0.17048 (13)	0.0322 (3)
C5	0.0984 (2)	0.20371 (18)	0.40485 (13)	0.0375 (3)
H5	0.0730	0.1229	0.4847	0.045*
C6	0.1144 (2)	0.14491 (16)	0.28236 (13)	0.0348 (3)

supplementary materials

C9	0.3957 (2)	0.14516 (17)	0.14346 (14)	0.0372 (3)
C1	-0.0622 (2)	0.35556 (19)	0.37706 (14)	0.0383 (3)
C3	0.1375 (2)	0.52059 (18)	0.25385 (15)	0.0406 (3)
H3A	0.1661	0.5959	0.1728	0.049*
H3B	0.0969	0.5788	0.3209	0.049*
C2	-0.0144 (2)	0.44610 (17)	0.23864 (13)	0.0362 (3)
H2A	-0.1220	0.5275	0.1970	0.043*
C14	0.5784 (2)	0.0851 (2)	0.09741 (16)	0.0486 (4)
H14	0.6375	0.1503	0.0301	0.058*
C11	0.3986 (2)	-0.11196 (19)	0.30063 (17)	0.0496 (4)
H11	0.3394	-0.1777	0.3674	0.059*
C12	0.5804 (3)	-0.1720 (2)	0.25488 (18)	0.0565 (5)
H12	0.6442	-0.2794	0.2917	0.068*
C13	0.6699 (2)	-0.0751 (2)	0.15496 (19)	0.0572 (5)
H13	0.7927	-0.1183	0.1263	0.069*
C18	0.4702 (3)	0.4293 (3)	0.2839 (2)	0.0624 (5)
H18A	0.4881	0.5036	0.2010	0.094*
H18B	0.5737	0.3334	0.2949	0.094*
H18C	0.4615	0.4811	0.3517	0.094*

Atomic displacement parameters (\AA^2)

	U^{11}	U^{22}	U^{33}	U^{12}	U^{13}	U^{23}
O1	0.0438 (6)	0.0456 (6)	0.0332 (5)	-0.0098 (5)	0.0005 (4)	0.0021 (5)
O2	0.0419 (6)	0.0433 (6)	0.0416 (6)	-0.0060 (5)	-0.0146 (5)	-0.0116 (5)
O3	0.0522 (7)	0.0488 (6)	0.0518 (7)	-0.0269 (5)	-0.0081 (5)	-0.0022 (5)
N1	0.0360 (7)	0.0438 (7)	0.0393 (7)	-0.0155 (5)	0.0001 (5)	-0.0116 (5)
O4	0.0413 (6)	0.0706 (8)	0.0475 (7)	-0.0118 (6)	0.0104 (5)	-0.0234 (6)
C8	0.0365 (8)	0.0379 (7)	0.0285 (7)	-0.0084 (6)	-0.0037 (6)	-0.0053 (6)
C4	0.0413 (8)	0.0455 (8)	0.0329 (7)	-0.0085 (6)	-0.0075 (6)	-0.0095 (6)
C10	0.0422 (8)	0.0343 (7)	0.0341 (7)	-0.0068 (6)	-0.0073 (6)	-0.0090 (6)
C7	0.0320 (7)	0.0335 (7)	0.0289 (7)	-0.0067 (6)	-0.0057 (5)	-0.0053 (6)
C5	0.0396 (8)	0.0413 (8)	0.0268 (7)	-0.0130 (6)	0.0003 (6)	-0.0013 (6)
C6	0.0369 (8)	0.0322 (7)	0.0338 (7)	-0.0120 (6)	-0.0039 (6)	-0.0030 (6)
C9	0.0378 (8)	0.0387 (8)	0.0326 (7)	-0.0050 (6)	-0.0052 (6)	-0.0101 (6)
C1	0.0350 (8)	0.0468 (8)	0.0360 (8)	-0.0141 (6)	0.0002 (6)	-0.0135 (6)
C3	0.0491 (9)	0.0361 (8)	0.0370 (8)	-0.0128 (7)	-0.0008 (6)	-0.0100 (6)
C2	0.0349 (7)	0.0354 (7)	0.0339 (7)	-0.0028 (6)	-0.0053 (6)	-0.0080 (6)
C14	0.0396 (8)	0.0557 (10)	0.0453 (9)	-0.0037 (7)	-0.0015 (7)	-0.0160 (8)
C11	0.0600 (10)	0.0342 (8)	0.0482 (9)	-0.0044 (7)	-0.0111 (8)	-0.0064 (7)
C12	0.0617 (11)	0.0404 (9)	0.0592 (11)	0.0085 (8)	-0.0191 (9)	-0.0161 (8)
C13	0.0442 (9)	0.0592 (11)	0.0625 (11)	0.0096 (8)	-0.0102 (8)	-0.0286 (9)
C18	0.0456 (10)	0.0723 (12)	0.0787 (13)	-0.0281 (9)	0.0013 (9)	-0.0244 (10)

Geometric parameters (\AA , $^\circ$)

O1—C8	1.2285 (17)	C5—C1	1.509 (2)
O2—C7	1.4171 (16)	C5—C6	1.537 (2)
O2—H2	0.82	C5—H5	0.98

O3—C6	1.4219 (17)	C9—C14	1.387 (2)
O3—H3	0.82	C1—C2	1.517 (2)
N1—C3	1.453 (2)	C3—C2	1.541 (2)
N1—C18	1.455 (2)	C3—H3A	0.97
N1—C4	1.4613 (19)	C3—H3B	0.97
O4—C1	1.2088 (18)	C2—H2A	0.98
C8—C9	1.471 (2)	C14—C13	1.385 (3)
C8—C7	1.5419 (19)	C14—H14	0.93
C4—C5	1.539 (2)	C11—C12	1.381 (3)
C4—H4A	0.97	C11—H11	0.93
C4—H4B	0.97	C12—C13	1.388 (3)
C10—C11	1.386 (2)	C12—H12	0.93
C10—C9	1.391 (2)	C13—H13	0.93
C10—C6	1.507 (2)	C18—H18A	0.96
C7—C2	1.5443 (19)	C18—H18B	0.96
C7—C6	1.5720 (18)	C18—H18C	0.96
C7—O2—H2	109.5	C14—C9—C8	128.62 (14)
C6—O3—H3	109.5	C10—C9—C8	109.95 (13)
C3—N1—C18	114.32 (13)	O4—C1—C5	128.54 (14)
C3—N1—C4	113.97 (12)	O4—C1—C2	126.94 (14)
C18—N1—C4	112.86 (13)	C5—C1—C2	104.23 (11)
O1—C8—C9	126.53 (13)	N1—C3—C2	105.67 (11)
O1—C8—C7	124.00 (12)	N1—C3—H3A	110.6
C9—C8—C7	108.44 (12)	C2—C3—H3A	110.6
N1—C4—C5	110.50 (11)	N1—C3—H3B	110.6
N1—C4—H4A	109.6	C2—C3—H3B	110.6
C5—C4—H4A	109.6	H3A—C3—H3B	108.7
N1—C4—H4B	109.6	C1—C2—C3	103.05 (11)
C5—C4—H4B	109.6	C1—C2—C7	103.33 (11)
H4A—C4—H4B	108.1	C3—C2—C7	109.41 (11)
C11—C10—C9	120.35 (15)	C1—C2—H2A	113.4
C11—C10—C6	128.39 (14)	C3—C2—H2A	113.4
C9—C10—C6	111.20 (12)	C7—C2—H2A	113.4
O2—C7—C8	108.50 (11)	C13—C14—C9	117.82 (16)
O2—C7—C2	114.39 (11)	C13—C14—H14	121.1
C8—C7—C2	115.40 (12)	C9—C14—H14	121.1
O2—C7—C6	108.69 (11)	C12—C11—C10	118.30 (16)
C8—C7—C6	103.21 (11)	C12—C11—H11	120.8
C2—C7—C6	105.84 (11)	C10—C11—H11	120.8
C1—C5—C6	100.91 (11)	C11—C12—C13	121.32 (15)
C1—C5—C4	105.49 (12)	C11—C12—H12	119.3
C6—C5—C4	112.08 (11)	C13—C12—H12	119.3
C1—C5—H5	112.5	C14—C13—C12	120.77 (16)
C6—C5—H5	112.5	C14—C13—H13	119.6
C4—C5—H5	112.5	C12—C13—H13	119.6
O3—C6—C10	114.30 (12)	N1—C18—H18A	109.5
O3—C6—C5	106.98 (11)	N1—C18—H18B	109.5
C10—C6—C5	114.94 (12)	H18A—C18—H18B	109.5
O3—C6—C7	111.58 (11)	N1—C18—H18C	109.5

supplementary materials

C10—C6—C7	104.97 (11)	H18A—C18—H18C	109.5
C5—C6—C7	103.62 (11)	H18B—C18—H18C	109.5
C14—C9—C10	121.43 (14)		
C3—N1—C4—C5	54.45 (16)	C6—C10—C9—C14	176.26 (14)
C18—N1—C4—C5	-172.98 (13)	C11—C10—C9—C8	178.38 (14)
O1—C8—C7—O2	66.57 (17)	C6—C10—C9—C8	-4.31 (17)
C9—C8—C7—O2	-102.49 (12)	O1—C8—C9—C14	4.9 (3)
O1—C8—C7—C2	-63.28 (18)	C7—C8—C9—C14	173.60 (15)
C9—C8—C7—C2	127.66 (12)	O1—C8—C9—C10	-174.49 (14)
O1—C8—C7—C6	-178.24 (13)	C7—C8—C9—C10	-5.78 (16)
C9—C8—C7—C6	12.70 (14)	C6—C5—C1—O4	138.05 (16)
N1—C4—C5—C1	-56.97 (15)	C4—C5—C1—O4	-105.16 (17)
N1—C4—C5—C6	51.96 (16)	C6—C5—C1—C2	-47.81 (13)
C11—C10—C6—O3	-48.1 (2)	C4—C5—C1—C2	68.98 (13)
C9—C10—C6—O3	134.84 (13)	C18—N1—C3—C2	168.31 (13)
C11—C10—C6—C5	76.15 (19)	C4—N1—C3—C2	-59.82 (15)
C9—C10—C6—C5	-100.88 (14)	O4—C1—C2—C3	98.33 (17)
C11—C10—C6—C7	-170.70 (15)	C5—C1—C2—C3	-75.95 (13)
C9—C10—C6—C7	12.27 (15)	O4—C1—C2—C7	-147.76 (15)
C1—C5—C6—O3	-80.03 (13)	C5—C1—C2—C7	37.97 (14)
C4—C5—C6—O3	168.14 (11)	N1—C3—C2—C1	69.03 (14)
C1—C5—C6—C10	151.91 (12)	N1—C3—C2—C7	-40.39 (14)
C4—C5—C6—C10	40.09 (16)	O2—C7—C2—C1	106.41 (13)
C1—C5—C6—C7	37.97 (13)	C8—C7—C2—C1	-126.65 (12)
C4—C5—C6—C7	-73.85 (14)	C6—C7—C2—C1	-13.22 (14)
O2—C7—C6—O3	-23.90 (15)	O2—C7—C2—C3	-144.35 (12)
C8—C7—C6—O3	-138.96 (12)	C8—C7—C2—C3	-17.42 (15)
C2—C7—C6—O3	99.41 (13)	C6—C7—C2—C3	96.02 (13)
O2—C7—C6—C10	100.42 (12)	C10—C9—C14—C13	0.3 (2)
C8—C7—C6—C10	-14.64 (14)	C8—C9—C14—C13	-178.97 (15)
C2—C7—C6—C10	-136.28 (11)	C9—C10—C11—C12	1.0 (2)
O2—C7—C6—C5	-138.66 (11)	C6—C10—C11—C12	-175.81 (15)
C8—C7—C6—C5	106.28 (12)	C10—C11—C12—C13	-0.3 (3)
C2—C7—C6—C5	-15.36 (14)	C9—C14—C13—C12	0.4 (3)
C11—C10—C9—C14	-1.1 (2)	C11—C12—C13—C14	-0.4 (3)

Hydrogen-bond geometry (Å, °)

<i>D</i> —H... <i>A</i>	<i>D</i> —H	H... <i>A</i>	<i>D</i> ... <i>A</i>	<i>D</i> —H... <i>A</i>
O3—H3...O2	0.82	2.11	2.6080 (15)	119
O2—H2...O1 ⁱ	0.82	1.90	2.7168 (15)	175
C2—H2A...O1 ⁱ	0.98	2.59	3.3778 (18)	137
C18—H18A...O1 ⁱⁱ	0.96	2.57	3.432 (2)	149
C5—H5...O3 ⁱⁱⁱ	0.98	2.46	3.4381 (18)	173

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

Fig. 1

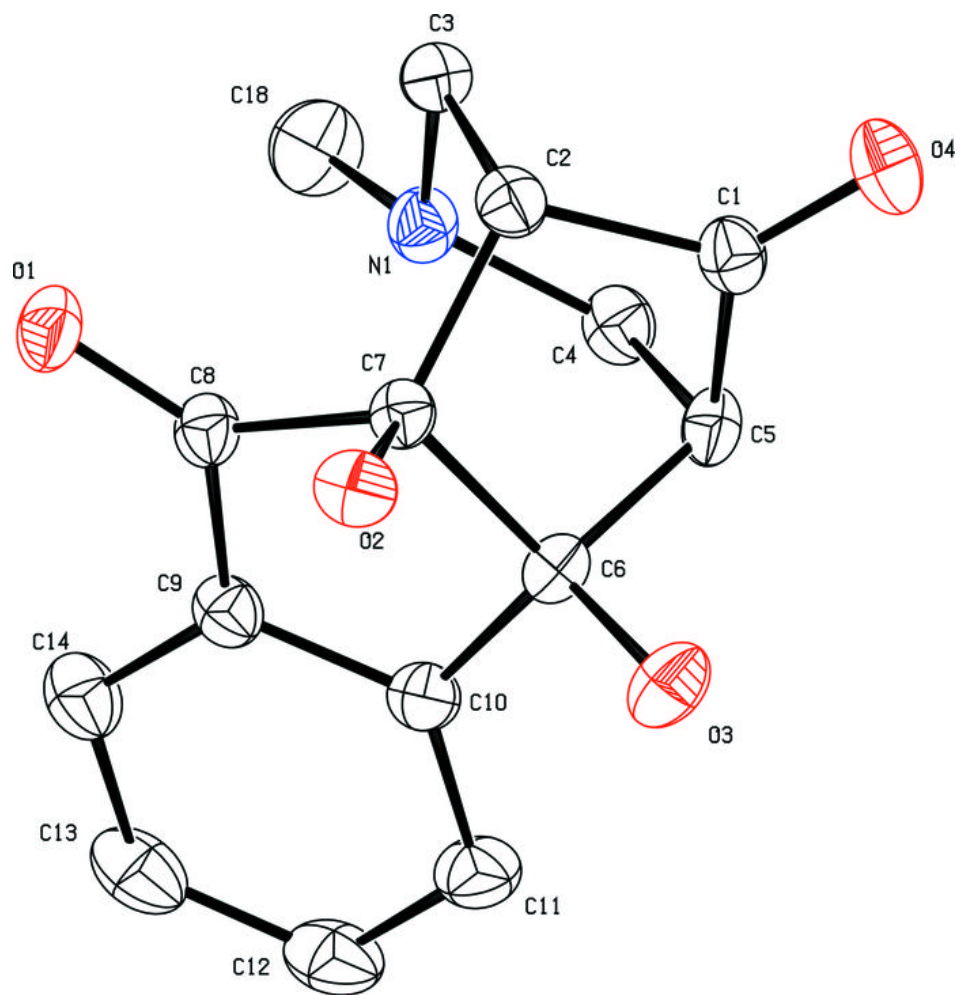


Fig. 2

