

(1*RS*,2*RS*,3*SR*,5*RS*,7*RS*)-2,5-Dichloro-8-oxabicyclo[5.1.0]octan-3-ol

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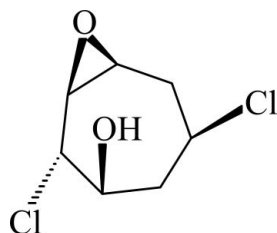
Received 25 March 2011; accepted 8 April 2011

Key indicators: single-crystal X-ray study; $T = 294$ K; mean $\sigma(\text{C}-\text{C}) = 0.008$ Å; R factor = 0.088; wR factor = 0.190; data-to-parameter ratio = 15.7.

In the title compound, $\text{C}_7\text{H}_{10}\text{Cl}_2\text{O}_2$, the seven-membered ring displays a chair conformation. In the crystal, the hydroxy H atom is equally disordered over two orientations, and links with an adjacent molecule *via* an $\text{O}-\text{H}\cdots\text{O}$ hydrogen bond in both cases. Weak intermolecular $\text{C}-\text{H}\cdots\text{O}$ hydrogen bonding is also a feature of the crystal structure.

Related literature

For background to *syn*-bis-epoxides, see: Balcı (1981); Akbulut *et al.* (1987); Menzek & Balcı (1993); Saraçoğlu *et al.* (1999). For background to unsaturated bicyclic endoperoxide, see: Menzek *et al.* (2005). For background to epoxide and bis-epoxide, see: Şengül *et al.* (2008).



Experimental

Crystal data

$\text{C}_7\text{H}_{10}\text{Cl}_2\text{O}_2$

$M_r = 197.05$

Orthorhombic, *Pbcn*

$a = 21.9202$ (5) Å

$b = 9.9343$ (3) Å

$c = 8.1005$ (2) Å

$V = 1763.98$ (8) Å³

$Z = 8$

Mo $K\alpha$ radiation

$\mu = 0.68$ mm⁻¹

$T = 294$ K

$0.32 \times 0.20 \times 0.15$ mm

Data collection

Rigaku R-AXIS RAPID-S

diffractometer

Absorption correction: multi-scan

(Blessing, 1995)

$T_{\min} = 0.845$, $T_{\max} = 0.900$

32813 measured reflections

1801 independent reflections

1267 reflections with $I > 2\sigma(I)$

$R_{\text{int}} = 0.110$

Refinement

$R[F^2 > 2\sigma(F^2)] = 0.088$

$wR(F^2) = 0.190$

$S = 1.18$

1801 reflections

115 parameters

2 restraints

H atoms treated by a mixture of independent and constrained refinement

$\Delta\rho_{\text{max}} = 0.28$ e Å⁻³

$\Delta\rho_{\text{min}} = -0.34$ e Å⁻³

Table 1

Hydrogen-bond geometry (Å, °).

$D-\text{H}\cdots A$	$D-\text{H}$	$\text{H}\cdots A$	$D\cdots A$	$D-\text{H}\cdots A$
$\text{O2}-\text{H2A}\cdots\text{O2}^{\text{i}}$	0.83 (11)	1.96 (11)	2.746 (7)	159 (12)
$\text{O2}-\text{H2B}\cdots\text{O2}^{\text{ii}}$	0.85	1.84	2.692 (8)	174
$\text{C2}-\text{H21}\cdots\text{O1}^{\text{iii}}$	0.97	2.43	3.398 (7)	172

Symmetry codes: (i) $-x + 1, -y + 2, -z$; (ii) $-x + 1, y, -z + \frac{1}{2}$; (iii) $-x + \frac{3}{2}, y - \frac{1}{2}, z$.

Data collection: *CrystalClear* (Rigaku/MSC, 2005); cell refinement: *CrystalClear*; data reduction: *CrystalClear*; program(s) used to solve structure: *SHELXS97* (Sheldrick, 2008); program(s) used to refine structure: *SHELXL97* (Sheldrick, 2008); molecular graphics: *ORTEP-3 for Windows* (Farrugia, 1997) and *Mercury* (Macrae *et al.*, 2006); software used to prepare material for publication: *WinGX* (Farrugia, 1999) and *PLATON* (Spek, 2009).

The authors are indebted to the Department of Chemistry, Atatürk University, Erzurum, Turkey, for the use of X-ray diffractometer purchased under grant No. 2003/219 of the University Research Fund.

Supplementary data and figures for this paper are available from the IUCr electronic archives (Reference: XU5182).

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supporting information

Acta Cryst. (2011). E67, o1145 [doi:10.1107/S1600536811013304]

(1*RS*,2*RS*,3*SR*,5*RS*,7*RS*)-2,5-Dichloro-8-oxabicyclo[5.1.0]octan-3-ol**Yasin Çetinkaya, Abdullah Menzek and Tuncer Hökelek****S1. Comment**

Unsaturated bicyclic endoperoxides are important compounds for versatile chemical transformations in organic chemistry. In these endoperoxides, diradicals formed by thermal cleavage of the weak O-O bonds react to give the syn-bis-epoxides (Balci, 1981; Akbulut *et al.*, 1987; Menzek & Balci, 1993; Saraçoğlu *et al.*, 1999).

Unsaturated bicyclic endoperoxide, (1), (Scheme 1) was synthesized by the literature method (Menzek *et al.*, 2005). Reaction of endoperoxide, (1), by heating at 453 (5) K gave a mixture of products (Scheme 1). The title compound, (2), was isolated from these mixtures. The other products were not identified. According to the NMR data of dichloride, (2), it was not easy to establish the exact configuration of the molecule. Therefore, the exact structure of dichloride, (2), was determined by X-ray single crystal analysis.

To rationalize the formation of dichloride, (2), we propose the following reaction mechanism as favourable mechanism (Scheme 1). Bis-epoxide, (3), is produced via diradicals formed by thermal cleavage of the weak O-O bonds. HCl formed by elimination from reaction products such as (3) can attack bis-epoxide, (3), to give intermediate (4). Cl⁻ can attack intermediate (4) to give dichloride, (2), as a nucleophile. An epoxide or bis-epoxide ring (Şengül *et al.*, 2008) may be opened by as a nucleophile.

In the title compound, the seven-membered ring A (C1-C7) is, of course, not planar. The planar moieties B (O1/C1/C7), C (C3-C5), D (C2/C3/C5/C6) and E (C1/C2/C6/C7) are oriented at dihedral angles of B/C = 68.75 (46)°, B/D = 13.84 (28)°, B/E = 74.48 (32)°, C/D = 54.91 (42)°, C/E = 6.00 (42)° and D/E = 60.65 (30)°.

In the crystal, intermolecular O—H...O and C—H...O hydrogen bonds link the molecules into a three-dimensional network (Table 1 and Fig. 2).

S2. Experimental

For the preparation of the title compound, a mixture of endoperoxide (0.5 g, 2.6 mmol) and benzene (5 ml) was placed into a test tube, sealed under vacuum and heated at 453 (5) K for 3 d. After cooling to room temperature, the solvent was evaporated. The residue was submitted to column chromatography (silica gel, 90 g) with AcOEt/hexane (1:6) as eluant. Dichloride (yield: 0.056 g, 9%, m. p. 366-368 K) and a mixture of unidentified products were obtained. Dichloride was crystallized from ethyl acetate/hexane (1:1) as colorless block crystals.

S3. Refinement

H1, H7, H21, H22 and H41, H42 atoms were positioned geometrically with C—H = 0.98 and 0.97 Å, respectively, and constrained to ride on their parent atoms, with $U_{\text{iso}}(\text{H}) = 1.2U_{\text{eq}}(\text{C})$. The remaining H-atoms were located in a difference Fourier map and refined isotropically. The H atom of the OH group was disordered over two orientations. During the refinement process, the disordered H2A and H2B atoms were refined with equal occupancies.

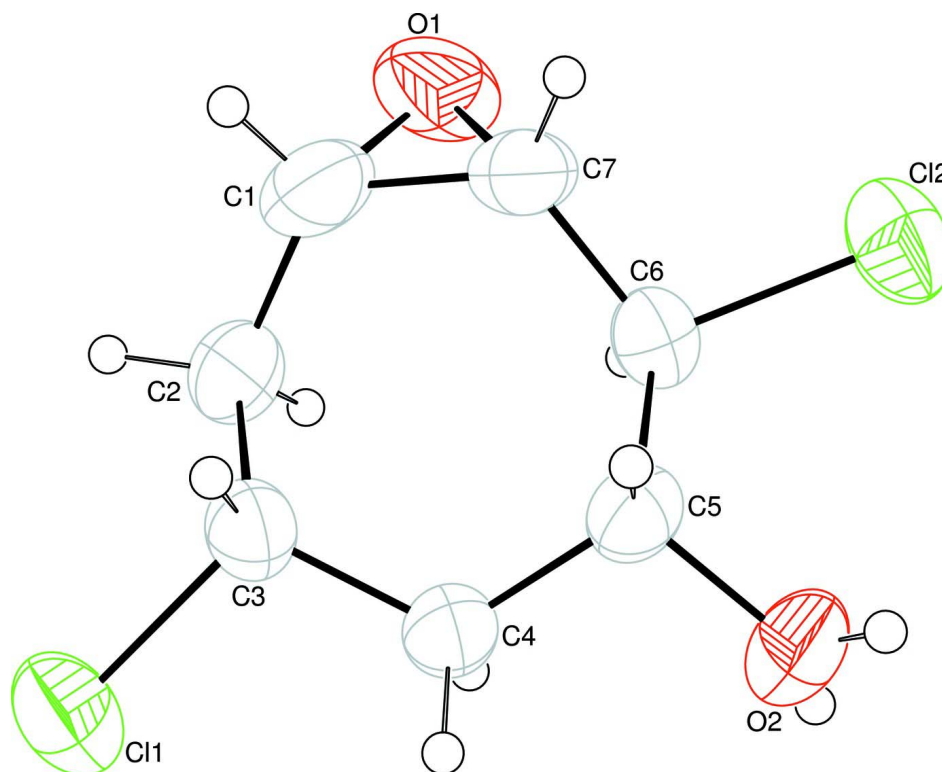


Figure 1

The molecular structure of the title molecule with the atom-numbering scheme. Displacement ellipsoids are drawn at the 40% probability level.

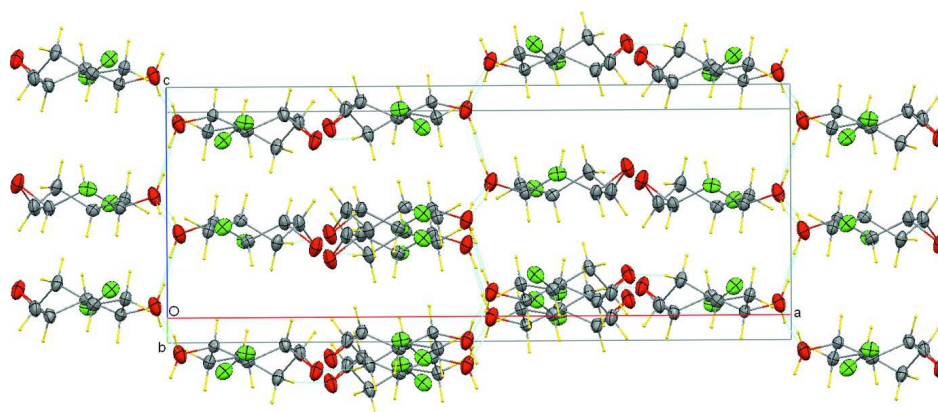


Figure 2

A view of the crystal packing of the title compound. The O-H...O and C-H...O hydrogen bonds are shown as dashed lines.

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Crystal data

C₇H₁₀Cl₂O₂

M_r = 197.05

Orthorhombic, *Pbcn*

Hall symbol: -*P* 2_n 2_{ab}

a = 21.9202 (5) Å

b = 9.9343 (3) Å

c = 8.1005 (2) Å

V = 1763.98 (8) Å³

Z = 8

F(000) = 816

$D_x = 1.484 \text{ Mg m}^{-3}$
 Mo $K\alpha$ radiation, $\lambda = 0.71073 \text{ \AA}$
 Cell parameters from 4978 reflections
 $\theta = 2.3\text{--}26.4^\circ$

$\mu = 0.68 \text{ mm}^{-1}$
 $T = 294 \text{ K}$
 Block, colorless
 $0.32 \times 0.20 \times 0.15 \text{ mm}$

Data collection

Rigaku R-AXIS RAPID-S
 diffractometer
 Radiation source: fine-focus sealed tube
 Graphite monochromator
 ω scans
 Absorption correction: multi-scan
 (Blessing, 1995)
 $T_{\min} = 0.845$, $T_{\max} = 0.900$

32813 measured reflections
 1801 independent reflections
 1267 reflections with $I > 2\sigma(I)$
 $R_{\text{int}} = 0.110$
 $\theta_{\max} = 26.4^\circ$, $\theta_{\min} = 2.3^\circ$
 $h = -27 \rightarrow 27$
 $k = -12 \rightarrow 12$
 $l = -9 \rightarrow 10$

Refinement

Refinement on F^2
 Least-squares matrix: full
 $R[F^2 > 2\sigma(F^2)] = 0.088$
 $wR(F^2) = 0.190$
 $S = 1.18$
 1801 reflections
 115 parameters
 2 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
 $w = 1/[\sigma^2(F_o^2) + (0.020P)^2 + 5.1831P]$
 where $P = (F_o^2 + 2F_c^2)/3$
 $(\Delta/\sigma)_{\max} < 0.001$
 $\Delta\rho_{\max} = 0.28 \text{ e \AA}^{-3}$
 $\Delta\rho_{\min} = -0.34 \text{ e \AA}^{-3}$

Special details

Geometry. All esds (except the esd in the dihedral angle between two l.s. planes) are estimated using the full covariance matrix. The cell esds are taken into account individually in the estimation of esds in distances, angles and torsion angles; correlations between esds in cell parameters are only used when they are defined by crystal symmetry. An approximate (isotropic) treatment of cell esds is used for estimating esds 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 $F^2 > 2\sigma(F^2)$ 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)

	<i>x</i>	<i>y</i>	<i>z</i>	$U_{\text{iso}}^*/U_{\text{eq}}$	Occ. (<1)
Cl1	0.59049 (9)	0.41547 (16)	0.1157 (2)	0.0883 (6)	
Cl2	0.62620 (9)	1.07361 (16)	0.1076 (2)	0.0841 (6)	
O1	0.7389 (2)	0.8227 (5)	0.1601 (7)	0.1017 (17)	
O2	0.5199 (2)	0.8930 (5)	0.0928 (7)	0.0941 (16)	
H2A	0.511 (7)	0.945 (11)	0.017 (12)	0.090*	0.50
H2B	0.5050	0.8950	0.1900	0.090*	0.50
C1	0.7119 (3)	0.7145 (7)	0.0682 (10)	0.084 (2)	
H1	0.7361	0.6806	-0.0247	0.101*	
C2	0.6766 (2)	0.6091 (6)	0.1650 (9)	0.0732 (18)	
H21	0.7003	0.5271	0.1751	0.088*	
H22	0.6672	0.6421	0.2748	0.088*	

C3	0.6182 (3)	0.5825 (6)	0.0692 (8)	0.0604 (15)
H3	0.628 (3)	0.578 (7)	-0.042 (9)	0.10 (2)*
C4	0.5669 (2)	0.6810 (5)	0.1042 (8)	0.0610 (14)
H41	0.5303	0.6467	0.0516	0.073*
H42	0.5595	0.6806	0.2223	0.073*
C5	0.5748 (2)	0.8269 (6)	0.0507 (8)	0.0582 (14)
H5	0.580 (2)	0.832 (5)	-0.066 (7)	0.057 (16)*
C6	0.6294 (3)	0.8926 (5)	0.1265 (7)	0.0558 (13)
H6	0.636 (2)	0.873 (5)	0.231 (7)	0.066 (18)*
C7	0.6885 (2)	0.8499 (6)	0.0502 (8)	0.0703 (17)
H7	0.6993	0.8954	-0.0531	0.084*

Atomic displacement parameters (Å²)

	U^{11}	U^{22}	U^{33}	U^{12}	U^{13}	U^{23}
Cl1	0.1032 (13)	0.0592 (9)	0.1024 (14)	-0.0091 (8)	-0.0189 (11)	0.0044 (9)
Cl2	0.1150 (14)	0.0573 (9)	0.0799 (11)	-0.0025 (9)	0.0052 (10)	-0.0046 (8)
O1	0.067 (3)	0.095 (4)	0.143 (5)	-0.021 (3)	-0.030 (3)	0.028 (3)
O2	0.067 (3)	0.084 (3)	0.132 (5)	0.024 (2)	0.023 (3)	0.012 (3)
C1	0.049 (3)	0.097 (5)	0.107 (6)	0.011 (3)	0.009 (4)	0.011 (5)
C2	0.055 (3)	0.069 (4)	0.096 (5)	0.014 (3)	-0.016 (3)	0.010 (3)
C3	0.066 (4)	0.052 (3)	0.063 (4)	0.001 (3)	0.001 (3)	-0.005 (3)
C4	0.049 (3)	0.058 (3)	0.075 (4)	0.001 (2)	-0.009 (3)	-0.004 (3)
C5	0.050 (3)	0.064 (3)	0.061 (4)	0.008 (3)	-0.001 (3)	0.004 (3)
C6	0.066 (3)	0.049 (3)	0.052 (3)	0.001 (2)	-0.009 (3)	0.000 (3)
C7	0.052 (3)	0.075 (4)	0.084 (4)	-0.006 (3)	0.009 (3)	0.014 (3)

Geometric parameters (Å, °)

Cl1—C3	1.806 (6)	C3—C2	1.520 (8)
Cl2—C6	1.806 (6)	C3—C4	1.518 (7)
O1—C1	1.434 (8)	C3—H3	0.93 (7)
O1—C7	1.443 (7)	C4—C5	1.522 (8)
O2—C5	1.413 (7)	C4—H41	0.9700
O2—H2B	0.853 (5)	C4—H42	0.9700
O2—H2A	0.82 (2)	C5—H5	0.95 (5)
C1—C7	1.447 (9)	C6—C5	1.497 (8)
C1—H1	0.9800	C6—C7	1.497 (8)
C2—C1	1.521 (8)	C6—H6	0.88 (6)
C2—H21	0.9700	C7—H7	0.9800
C2—H22	0.9700		
C5—O2—H2B	124.0 (5)	C3—C4—H41	107.7
C5—O2—H2A	109 (10)	C3—C4—H42	107.7
H2B—O2—H2A	125 (10)	C5—C4—H41	107.7
C1—O1—C7	60.4 (4)	C5—C4—H42	107.7
O1—C1—C2	117.3 (6)	H42—C4—H41	107.1
O1—C1—C7	60.1 (4)	O2—C5—C4	106.1 (5)

O1—C1—H1	115.7	O2—C5—C6	112.3 (5)
C2—C1—H1	115.7	O2—C5—H5	109 (3)
C7—C1—C2	120.7 (5)	C4—C5—H5	110 (3)
C7—C1—H1	115.7	C6—C5—C4	112.9 (5)
C1—C2—H21	110.4	C6—C5—H5	107 (3)
C1—C2—H22	110.4	C12—C6—H6	108 (4)
C3—C2—C1	106.6 (5)	C5—C6—C12	111.6 (4)
C3—C2—H21	110.4	C5—C6—C7	113.6 (5)
C3—C2—H22	110.4	C5—C6—H6	116 (4)
H21—C2—H22	108.6	C7—C6—C12	106.3 (4)
C11—C3—H3	104 (4)	C7—C6—H6	101 (4)
C2—C3—C11	109.6 (4)	O1—C7—C1	59.5 (4)
C2—C3—H3	108 (4)	O1—C7—C6	117.4 (6)
C4—C3—C11	107.7 (4)	O1—C7—H7	115.4
C4—C3—C2	114.6 (5)	C1—C7—C6	122.0 (5)
C4—C3—H3	113 (4)	C1—C7—H7	115.4
C3—C4—C5	118.4 (5)	C6—C7—H7	115.4
C7—O1—C1—C2	-111.5 (6)	C3—C4—C5—O2	177.8 (5)
C1—O1—C7—C6	112.7 (6)	C3—C4—C5—C6	-58.7 (7)
O1—C1—C7—C6	-105.2 (7)	C12—C6—C5—O2	-44.3 (6)
C2—C1—C7—O1	105.9 (8)	C12—C6—C5—C4	-164.2 (4)
C2—C1—C7—C6	0.7 (10)	C7—C6—C5—O2	-164.5 (5)
C3—C2—C1—O1	135.8 (5)	C7—C6—C5—C4	75.6 (7)
C3—C2—C1—C7	66.0 (8)	C12—C6—C7—C1	168.9 (6)
C11—C3—C2—C1	154.6 (5)	C12—C6—C7—O1	99.4 (5)
C4—C3—C2—C1	-84.1 (7)	C5—C6—C7—O1	-137.5 (5)
C11—C3—C4—C5	-170.8 (4)	C5—C6—C7—C1	-68.0 (8)
C2—C3—C4—C5	66.8 (7)		

Hydrogen-bond geometry (\AA , $^\circ$)

$D-H\cdots A$	$D-H$	$H\cdots A$	$D\cdots A$	$D-H\cdots A$
O2—H2A \cdots O2 ⁱ	0.83 (11)	1.96 (11)	2.746 (7)	159 (12)
O2—H2B \cdots O2 ⁱⁱ	0.85	1.84	2.692 (8)	174
C2—H21 \cdots O1 ⁱⁱⁱ	0.97	2.43	3.398 (7)	172

Symmetry codes: (i) $-x+1, -y+2, -z$; (ii) $-x+1, y, -z+1/2$; (iii) $-x+3/2, y-1/2, z$.