Acta Crystallographica Section E
Structure Reports Online
ISSN 1600-5368

2,6-Bis(2,4-dichlorobenzylidene)cyclohexanone

Huan-Mei Guo, Li Liu and Fang-Fang Jian*

Microscale Science Institute, Department of Chemistry and Chemical Engineering, Weifang University, Weifang 261061, People's Republic of China
Correspondence e-mail: ffjian2008@163.com

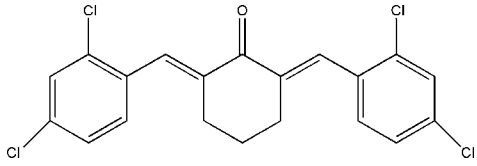
Received 10 July 2008; accepted 22 July 2008

Key indicators: single-crystal X-ray study; T = 293 K; mean σ(C–C) = 0.004 Å; R factor = 0.050; wR factor = 0.118; data-to-parameter ratio = 20.2.

The title compound, $C_{20}H_{14}Cl_4O$, was prepared from a mixture of 2,4-dichlorobenzophenone and cyclohexanone. The dihedral angles formed by the cyclohexane ring and two benzene rings are 39.18 (2) and 60.72 (2)°. There are some weak intramolecular C—H···O and C—H···Cl hydrogen-bond contacts in the crystal structure.

Related literature

For related literature, see: Butcher *et al.* (2006); Deli *et al.* (1984); Jia *et al.* (1989); Yu *et al.* (2000).



Experimental

Crystal data

$C_{20}H_{14}Cl_4O$
M_r = 412.11
Orthorhombic, *Pbca*
a = 14.469 (2) Å
b = 8.0602 (12) Å
c = 31.554 (4) Å
V = 3679.9 (9) Å^3
Z = 8
Mo $K\alpha$ radiation
μ = 0.65 mm^{-1}
T = 293 (2) K
0.20 × 0.15 × 0.10 mm

Data collection

Bruker SMART CCD area-detector diffractometer
Absorption correction: multi-scan (*SADABS*; Bruker, 1997)
T_{min} = 0.881, T_{max} = 0.938
21985 measured reflections
4570 independent reflections
2735 reflections with $I > 2\sigma(I)$
R_{int} = 0.053

Refinement

$R[F^2 > 2\sigma(F^2)]$ = 0.050
$wR(F^2)$ = 0.118
S = 1.03
4570 reflections
226 parameters
H-atom parameters constrained
$\Delta\rho_{max}$ = 0.24 e Å^{-3}
$\Delta\rho_{min}$ = −0.28 e Å^{-3}

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

D—H···A	D—H	H···A	D···A	D—H···A
C7—H7A···Cl2	0.93	2.74	3.055 (2)	101
C7—H7A···O1	0.93	2.33	2.732 (3)	105
C14—H14A···O1	0.93	2.39	2.759 (3)	103

Data collection: *SMART* (Bruker, 1997); cell refinement: *SAINT* (Bruker, 1997); 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*.

The authors thank the National Science Foundation of Weifang University (grant: No.2008Z04).

Supplementary data and figures for this paper are available from the IUCr electronic archives (Reference: AT2592).

References

Bruker (1997). *SADABS*, *SMART* and *SAINT*. Bruker AXS Inc., Madison, Wisconsin, USA.
Butcher, R. J., Yathirajan, H. S., Sarojini, B. K., Narayana, B. & Indira, J. (2006). *Acta Cryst.* E**62**, o1910–o1912.
Deli, J., Lorand, T., Szabo, D. & Foldesi, A. (1984). *Pharmazie*, **39**, 539–544.
Jia, Z., Quail, J. W., Arora, V. K. & Dimmock, J. R. (1989). *Acta Cryst.* C**45**, 285–289.
Sheldrick, G. M. (2008). *Acta Cryst.* A**64**, 112–122.
Yu, R. C., Yakimansky, A. V., Kothe, H., Voigt-Martin, I. G., Schollmeyer, D., Jansen, J., Zandbergen, H. & Tenkovtsev, A. V. (2000). *Acta Cryst.* A**56**, 436–450.

supporting information

Acta Cryst. (2008). E**64**, o1626 [doi:10.1107/S1600536808023003]

2,6-Bis(2,4-dichlorobenzylidene)cyclohexanone

Huan-Mei Guo, Li Liu and Fang-Fang Jian

S1. Comment

As useful precursors to potentially bioactive pyrimidine derivatives, a,a-bis(substituted benzylidene) cycloalkanones have attracted considerable attention for many years (Deli *et al.*, 1984). In recent years, a series of non-linear optically active bis(benzylidene) ketones have been synthesized (Yu *et al.*, 2000). As part of our search for new non-linear optically active compounds, we synthesized the title compound (I), and describe its structure here.

In the structure of (I) (Fig. 1), all of the bond lengthes and bond angles fall in the normal range (Yu *et al.*, 2000; Jia *et al.*, 1989; Butcher *et al.*, 2006). There are some weak C—H···O and C—H···Cl intramolecular hydrogen bonds in the crystal structure.

S2. Experimental

A mixture of the 2,4-dichlorobenzophenone (0.2 mol), and cyclohexanone (0.1 mol) and 10% NaOH (10 ml) was stirred in ethanol (30 mL) for 5 h to afford the title compound [yield: 82%]. Single crystals suitable for X-ray measurements were obtained by recrystallization from ethanol at room temperature.

S3. Refinement

H atoms were fixed geometrically and allowed to ride on their attached atoms, with C—H = 0.93–0.97 Å, and with U_{iso}(H) = 1.2U_{eq}(C).

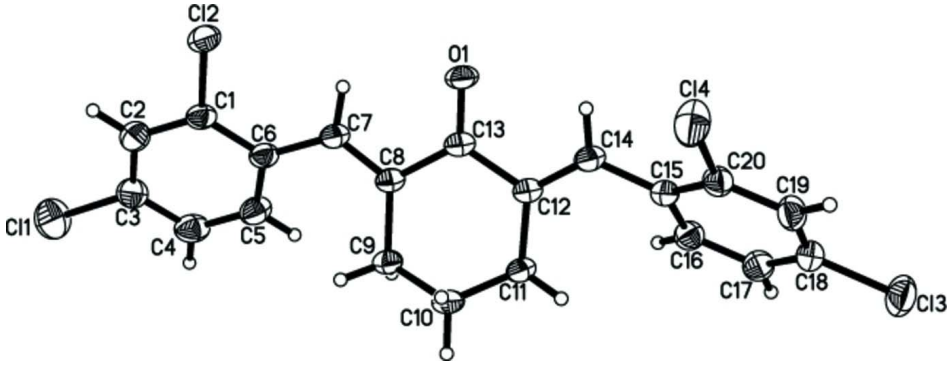


Figure 1

The structure of the title compound, showing 30% probability displacement ellipsoids and the atom-numbering scheme.

2,6-Bis(2,4-dichlorobenzylidene)cyclohexanone

Crystal data

$C_{20}H_{14}Cl_4O$
M_r = 412.11

Orthorhombic, *Pbca*
Hall symbol: -P 2ac 2ab

a = 14.469 (2) Å
b = 8.0602 (12) Å
c = 31.554 (4) Å
V = 3679.9 (9) Å^3
Z = 8
F(000) = 1680

D_x = 1.488 Mg m^{-3}
Mo $K\alpha$ radiation, λ = 0.71073 Å
μ = 0.65 mm^{-1}
T = 293 K
Bar, yellow
0.20 × 0.15 × 0.10 mm

Data collection

Bruker SMART CCD area-detector diffractometer
Radiation source: fine-focus sealed tube
Graphite monochromator
φ and ω scans
Absorption correction: multi-scan (*SADABS*; Bruker, 1997)
T_{min} = 0.881, T_{max} = 0.938

21985 measured reflections
4570 independent reflections
2735 reflections with $I > 2\sigma(I)$
R_{int} = 0.053
θ_{max} = 28.4°, θ_{min} = 1.3°
h = −19→18
k = −10→10
l = −19→42

Refinement

Refinement on F^2
Least-squares matrix: full
$R[F^2 > 2\sigma(F^2)]$ = 0.050
$wR(F^2)$ = 0.118
S = 1.03
4570 reflections
226 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
$w = 1/[\sigma^2(F_o^2) + (0.0381P)^2 + 1.6278P]$ where $P = (F_o^2 + 2F_c^2)/3$
$(\Delta/\sigma)_{max}$ = 0.001
$\Delta\rho_{max}$ = 0.24 e Å^{-3}
$\Delta\rho_{min}$ = −0.28 e Å^{-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 (Å^2)

	x	y	z	U_{iso}*/U_{eq}
Cl1	−0.37152 (7)	0.07869 (15)	−0.49524 (3)	0.1044 (4)
Cl2	−0.51233 (4)	0.04759 (9)	−0.33808 (2)	0.05594 (19)
Cl3	0.06595 (6)	0.60906 (12)	−0.04572 (3)	0.0852 (3)
Cl4	−0.21635 (6)	0.20911 (12)	−0.08713 (2)	0.0825 (3)
O1	−0.36351 (10)	0.3253 (2)	−0.22630 (5)	0.0557 (5)
C1	−0.42738 (15)	0.1234 (3)	−0.37207 (8)	0.0468 (6)
C2	−0.43315 (18)	0.0787 (4)	−0.41435 (8)	0.0578 (7)
H2A	−0.4804	0.0098	−0.4238	0.069*
C3	−0.36783 (19)	0.1381 (4)	−0.44228 (9)	0.0635 (8)
C4	−0.29917 (19)	0.2442 (4)	−0.42879 (9)	0.0663 (8)

H4A	−0.2572	0.2881	−0.4481	0.080*
C5	−0.29343 (17)	0.2844 (4)	−0.38653 (9)	0.0592 (7)
H5A	−0.2465	0.3550	−0.3776	0.071*
C6	−0.35581 (16)	0.2229 (3)	−0.35643 (8)	0.0466 (6)
C7	−0.34846 (15)	0.2620 (3)	−0.31106 (8)	0.0461 (6)
H7A	−0.4041	0.2752	−0.2967	0.055*
C8	−0.27162 (14)	0.2811 (3)	−0.28761 (8)	0.0419 (6)
C9	−0.17413 (14)	0.2600 (3)	−0.30424 (8)	0.0499 (6)
H9A	−0.1571	0.3581	−0.3202	0.060*
H9B	−0.1727	0.1661	−0.3235	0.060*
C10	−0.10328 (15)	0.2324 (3)	−0.26939 (8)	0.0502 (6)
H10A	−0.1142	0.1261	−0.2559	0.060*
H10B	−0.0417	0.2310	−0.2815	0.060*
C11	−0.11002 (15)	0.3693 (3)	−0.23677 (8)	0.0457 (6)
H11A	−0.0612	0.3563	−0.2160	0.055*
H11B	−0.1020	0.4760	−0.2505	0.055*
C12	−0.20266 (14)	0.3642 (3)	−0.21491 (8)	0.0412 (5)
C13	−0.28552 (15)	0.3230 (3)	−0.24158 (7)	0.0423 (6)
C14	−0.21556 (16)	0.3858 (3)	−0.17333 (8)	0.0466 (6)
H14A	−0.2751	0.3680	−0.1632	0.056*
C15	−0.14545 (15)	0.4349 (3)	−0.14198 (8)	0.0443 (6)
C16	−0.08247 (16)	0.5614 (3)	−0.15088 (8)	0.0512 (6)
H16A	−0.0840	0.6112	−0.1775	0.061*
C17	−0.01834 (18)	0.6150 (3)	−0.12186 (8)	0.0554 (7)
H17A	0.0227	0.6995	−0.1288	0.067*
C18	−0.01524 (17)	0.5426 (3)	−0.08242 (8)	0.0539 (7)
C19	−0.07653 (18)	0.4177 (3)	−0.07173 (8)	0.0555 (7)
H19A	−0.0746	0.3691	−0.0450	0.067*
C20	−0.14056 (17)	0.3665 (3)	−0.10137 (8)	0.0490 (6)

Atomic displacement parameters (Å²)

	U^{11}	U^{22}	U^{33}	U^{12}	U^{13}	U^{23}
Cl1	0.1071 (7)	0.1580 (10)	0.0482 (5)	−0.0068 (7)	0.0000 (5)	−0.0108 (5)
Cl2	0.0381 (3)	0.0687 (4)	0.0610 (4)	−0.0027 (3)	0.0010 (3)	0.0002 (3)
Cl3	0.0747 (5)	0.1149 (7)	0.0660 (5)	−0.0291 (5)	−0.0131 (4)	−0.0144 (5)
Cl4	0.0959 (6)	0.0945 (6)	0.0572 (5)	−0.0482 (5)	−0.0024 (4)	0.0122 (4)
O1	0.0313 (9)	0.0847 (14)	0.0509 (10)	−0.0009 (8)	0.0073 (8)	0.0041 (10)
C1	0.0355 (13)	0.0554 (16)	0.0497 (15)	0.0080 (11)	−0.0013 (11)	0.0045 (12)
C2	0.0506 (16)	0.0682 (19)	0.0546 (17)	0.0054 (13)	−0.0080 (13)	−0.0023 (14)
C3	0.0559 (17)	0.090 (2)	0.0445 (15)	0.0099 (16)	−0.0043 (14)	−0.0017 (15)
C4	0.0524 (16)	0.093 (2)	0.0531 (17)	0.0033 (16)	0.0063 (14)	0.0141 (16)
C5	0.0459 (15)	0.077 (2)	0.0549 (17)	−0.0076 (13)	−0.0001 (13)	0.0080 (15)
C6	0.0355 (12)	0.0555 (15)	0.0488 (15)	0.0069 (11)	−0.0019 (11)	0.0027 (12)
C7	0.0335 (12)	0.0547 (15)	0.0501 (15)	0.0002 (10)	0.0026 (11)	0.0034 (12)
C8	0.0331 (12)	0.0459 (14)	0.0468 (14)	−0.0012 (10)	0.0041 (10)	0.0056 (11)
C9	0.0331 (12)	0.0648 (17)	0.0517 (15)	−0.0026 (11)	0.0061 (11)	−0.0020 (13)
C10	0.0318 (12)	0.0576 (16)	0.0612 (16)	0.0049 (11)	0.0045 (12)	−0.0007 (13)

C11	0.0316 (12)	0.0537 (15)	0.0518 (15)	−0.0006 (11)	−0.0006 (11)	0.0023 (12)
C12	0.0327 (12)	0.0436 (14)	0.0473 (14)	0.0007 (10)	0.0024 (11)	0.0048 (11)
C13	0.0331 (12)	0.0458 (14)	0.0481 (14)	0.0018 (10)	0.0034 (11)	0.0096 (11)
C14	0.0340 (12)	0.0552 (15)	0.0507 (15)	−0.0003 (11)	0.0016 (11)	0.0050 (12)
C15	0.0370 (12)	0.0527 (15)	0.0431 (14)	0.0021 (11)	0.0027 (11)	−0.0019 (12)
C16	0.0490 (14)	0.0575 (16)	0.0471 (15)	−0.0007 (12)	0.0020 (12)	0.0047 (13)
C17	0.0513 (15)	0.0566 (17)	0.0583 (17)	−0.0101 (13)	0.0042 (13)	−0.0005 (14)
C18	0.0484 (14)	0.0633 (17)	0.0499 (16)	−0.0018 (13)	−0.0033 (12)	−0.0111 (14)
C19	0.0629 (17)	0.0642 (18)	0.0393 (14)	−0.0059 (14)	−0.0008 (12)	−0.0001 (13)
C20	0.0487 (14)	0.0513 (16)	0.0469 (15)	−0.0062 (12)	0.0052 (12)	−0.0013 (12)

Geometric parameters (Å, º)

Cl1—C3	1.739 (3)	C9—H9B	0.9700
Cl2—C1	1.742 (2)	C10—C11	1.512 (3)
Cl3—C18	1.735 (3)	C10—H10A	0.9700
Cl4—C20	1.736 (3)	C10—H10B	0.9700
O1—C13	1.227 (2)	C11—C12	1.508 (3)
C1—C2	1.384 (3)	C11—H11A	0.9700
C1—C6	1.400 (3)	C11—H11B	0.9700
C2—C3	1.378 (4)	C12—C14	1.337 (3)
C2—H2A	0.9300	C12—C13	1.502 (3)
C3—C4	1.378 (4)	C14—C15	1.471 (3)
C4—C5	1.375 (4)	C14—H14A	0.9300
C4—H4A	0.9300	C15—C16	1.396 (3)
C5—C6	1.401 (3)	C15—C20	1.397 (3)
C5—H5A	0.9300	C16—C17	1.373 (3)
C6—C7	1.470 (3)	C16—H16A	0.9300
C7—C8	1.345 (3)	C17—C18	1.375 (4)
C7—H7A	0.9300	C17—H17A	0.9300
C8—C13	1.504 (3)	C18—C19	1.383 (4)
C8—C9	1.515 (3)	C19—C20	1.380 (3)
C9—C10	1.520 (3)	C19—H19A	0.9300
C9—H9A	0.9700		
C2—C1—C6	122.2 (2)	H10A—C10—H10B	108.2
C2—C1—Cl2	117.4 (2)	C12—C11—C10	110.4 (2)
C6—C1—Cl2	120.4 (2)	C12—C11—H11A	109.6
C3—C2—C1	119.0 (3)	C10—C11—H11A	109.6
C3—C2—H2A	120.5	C12—C11—H11B	109.6
C1—C2—H2A	120.5	C10—C11—H11B	109.6
C2—C3—C4	120.8 (3)	H11A—C11—H11B	108.1
C2—C3—Cl1	119.9 (2)	C14—C12—C13	117.9 (2)
C4—C3—Cl1	119.3 (2)	C14—C12—C11	124.7 (2)
C5—C4—C3	119.3 (3)	C13—C12—C11	117.3 (2)
C5—C4—H4A	120.3	O1—C13—C12	120.7 (2)
C3—C4—H4A	120.3	O1—C13—C8	120.4 (2)
C4—C5—C6	122.4 (3)	C12—C13—C8	118.93 (19)

C4—C5—H5A	118.8	C12—C14—C15	126.8 (2)
C6—C5—H5A	118.8	C12—C14—H14A	116.6
C1—C6—C5	116.1 (2)	C15—C14—H14A	116.6
C1—C6—C7	121.3 (2)	C16—C15—C20	116.1 (2)
C5—C6—C7	122.5 (2)	C16—C15—C14	120.7 (2)
C8—C7—C6	128.4 (2)	C20—C15—C14	123.1 (2)
C8—C7—H7A	115.8	C17—C16—C15	122.5 (2)
C6—C7—H7A	115.8	C17—C16—H16A	118.8
C7—C8—C13	116.5 (2)	C15—C16—H16A	118.8
C7—C8—C9	124.5 (2)	C16—C17—C18	119.5 (2)
C13—C8—C9	118.96 (19)	C16—C17—H17A	120.3
C8—C9—C10	113.2 (2)	C18—C17—H17A	120.3
C8—C9—H9A	108.9	C17—C18—C19	120.6 (2)
C10—C9—H9A	108.9	C17—C18—Cl3	119.7 (2)
C8—C9—H9B	108.9	C19—C18—Cl3	119.8 (2)
C10—C9—H9B	108.9	C20—C19—C18	118.9 (2)
H9A—C9—H9B	107.8	C20—C19—H19A	120.6
C11—C10—C9	110.0 (2)	C18—C19—H19A	120.6
C11—C10—H10A	109.7	C19—C20—C15	122.5 (2)
C9—C10—H10A	109.7	C19—C20—Cl4	117.9 (2)
C11—C10—H10B	109.7	C15—C20—Cl4	119.59 (19)
C9—C10—H10B	109.7		

C6—C1—C2—C3	−1.9 (4)	C11—C12—C13—O1	−176.1 (2)
Cl2—C1—C2—C3	179.2 (2)	C14—C12—C13—C8	−173.9 (2)
C1—C2—C3—C4	−1.9 (4)	C11—C12—C13—C8	3.0 (3)
C1—C2—C3—Cl1	178.0 (2)	C7—C8—C13—O1	5.7 (3)
C2—C3—C4—C5	3.2 (4)	C9—C8—C13—O1	−173.0 (2)
Cl1—C3—C4—C5	−176.7 (2)	C7—C8—C13—C12	−173.4 (2)
C3—C4—C5—C6	−0.8 (4)	C9—C8—C13—C12	7.9 (3)
C2—C1—C6—C5	4.1 (4)	C13—C12—C14—C15	−176.6 (2)
Cl2—C1—C6—C5	−176.99 (19)	C11—C12—C14—C15	6.7 (4)
C2—C1—C6—C7	−176.9 (2)	C12—C14—C15—C16	43.5 (4)
Cl2—C1—C6—C7	1.9 (3)	C12—C14—C15—C20	−140.1 (3)
C4—C5—C6—C1	−2.8 (4)	C20—C15—C16—C17	0.8 (4)
C4—C5—C6—C7	178.3 (3)	C14—C15—C16—C17	177.5 (2)
C1—C6—C7—C8	145.0 (3)	C15—C16—C17—C18	0.0 (4)
C5—C6—C7—C8	−36.1 (4)	C16—C17—C18—C19	−0.5 (4)
C6—C7—C8—C13	179.4 (2)	C16—C17—C18—Cl3	179.7 (2)
C6—C7—C8—C9	−2.0 (4)	C17—C18—C19—C20	0.3 (4)
C7—C8—C9—C10	−161.0 (2)	Cl3—C18—C19—C20	−180.0 (2)
C13—C8—C9—C10	17.6 (3)	C18—C19—C20—C15	0.6 (4)
C8—C9—C10—C11	−53.5 (3)	C18—C19—C20—Cl4	179.8 (2)
C9—C10—C11—C12	64.1 (3)	C16—C15—C20—C19	−1.1 (4)
C10—C11—C12—C14	138.1 (3)	C14—C15—C20—C19	−177.7 (2)
C10—C11—C12—C13	−38.5 (3)	C16—C15—C20—Cl4	179.69 (18)
C14—C12—C13—O1	7.0 (4)	C14—C15—C20—Cl4	3.1 (3)

Hydrogen-bond geometry (Å, º)

D—H···*A*	*D*—H	H···*A*	*D*···*A*	*D*—H···*A*
C7—H7*A*···Cl2	0.93	2.74	3.055 (2)	101
C7—H7*A*···O1	0.93	2.33	2.732 (3)	105
C14—H14*A*···O1	0.93	2.39	2.759 (3)	103