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## Structure Reports

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## 3-(4-Methoxyphenyl)isochroman-1-one

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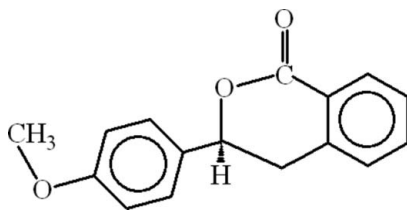
 Key indicators: single-crystal X-ray study;  $T = 296$  K; mean  $\sigma(C-C) = 0.002$  Å;

 $R$  factor = 0.042;  $wR$  factor = 0.117; data-to-parameter ratio = 18.9.

In the molecule of the title compound,  $C_{16}H_{14}O_3$ , the aromatic rings are oriented at a dihedral angle of  $72.02(6)^\circ$ . The heterocyclic ring adopts a twisted conformation. In the crystal structure, there are  $C-H \cdots \pi$  contacts between the heterocyclic and phenyl rings, and between the methyl group and methoxyphenyl ring.

## Related literature

For related structures, see: Schmalte *et al.* (1982); Schnebel *et al.* (2003). For bond-length data, see: Allen *et al.* (1987). For ring puckering parameters, see: Cremer & Pople (1975). For a description of the Cambridge Structural Database, see: Allen (2002).



## Experimental

## Crystal data

 $C_{16}H_{14}O_3$ 
 $M_r = 254.27$ 

 Monoclinic,  $P2_1/c$ 
 $a = 11.8933(7)$  Å

 $b = 14.6874(9)$  Å

 $c = 7.4521(4)$  Å

 $\beta = 101.040(2)^\circ$ 
 $V = 1277.66(13)$  Å<sup>3</sup>
 $Z = 4$ 

 Mo  $K\alpha$  radiation

 $\mu = 0.09$  mm<sup>-1</sup>
 $T = 296(2)$  K

 $0.24 \times 0.16 \times 0.12$  mm

## Data collection

Bruker KappaAPEXII CCD

diffractometer

Absorption correction: multi-scan

(SADABS; Bruker, 2005)

 $T_{\min} = 0.980$ ,  $T_{\max} = 0.996$ 

12426 measured reflections

3302 independent reflections

 2275 reflections with  $I > 2\sigma(I)$ 
 $R_{\text{int}} = 0.023$ 

## Refinement

 $R[F^2 > 2\sigma(F^2)] = 0.042$ 
 $wR(F^2) = 0.117$ 
 $S = 1.01$ 

3302 reflections

175 parameters

H atoms treated by a mixture of independent and constrained refinement

 $\Delta\rho_{\text{max}} = 0.14$  e Å<sup>-3</sup>
 $\Delta\rho_{\text{min}} = -0.22$  e Å<sup>-3</sup>

Table 1

Hydrogen-bond geometry (Å, °).

 $Cg1$  and  $Cg2$  are the centroids of the  $C2-C7$  and  $C10-C15$  rings, respectively.

$D-H \cdots A$	$D-H$	$H \cdots A$	$D \cdots A$	$D-H \cdots A$
$C8-H8B \cdots Cg2^i$	0.97	2.94	3.8398 (16)	154
$C16-H16B \cdots Cg3^{ii}$	0.96	2.89	3.7804 (17)	154

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

Data collection: *APEX2* (Bruker, 2007); cell refinement: *APEX2*; data reduction: *SAINT* (Bruker, 2007); 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 *PLATON* (Spek, 2003); software used to prepare material for publication: *WinGX* (Farrugia, 1999) and *PLATON*.

The authors acknowledge the Higher Education Commission, Islamabad, Pakistan, for funding the purchase of the diffractometer at GCU, Lahore.

Supplementary data and figures for this paper are available from the IUCr electronic archives (Reference: HK2543).

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**supplementary materials**

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### 3-(4-Methoxyphenyl)isochroman-1-one

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#### Comment

The title compound was prepared in order to evaluate its potential as antibacterial and antifungal agents. The CCDC search (Allen, 2002) showed that the crystal structures of *rac-exo*-tricarboxyl-(h6-3-phenylisochromanone)-chromium (Schnebel *et al.*, 2003) and 3,4-dihydro-8-hydroxy-3-(4-hydroxyphenyl)-isocoumarin (Schmalle *et al.*, 1982) have been reported, which have close resemblance as far as isochromane and attached phenyl ring is considered.

In the molecule of the title compound (Fig. 1), the bond lengths (Allen *et al.*, 1987) and angles are within normal ranges. Rings A (C2–C7) and C (C10–C15) are, of course, planar, and they are oriented at a dihedral angle of 72.02 (6)°. Ring B (O2/C1/C2/C7–C9) is not planar, having total puckering amplitude,  $Q_T$ , of 0.483 (2) Å and twisted conformation [ $\varphi = 41.63$  (3)° and  $\theta = 116.64$  (3)°] (Cremer & Pople, 1975).

In the crystal structure, there are C—H $\cdots\pi$  contacts (Table 1) between the heterocyclic and phenyl rings and the methyl group and methoxyphenyl ring, in which they may be effective in the stabilization of the structure.

#### Experimental

As shown in Fig. 3, a mixture of homophthalic acid, (1), (1.98 g, 11.0 mmol) and 4-methoxybenzoyl chloride, (2), (7.85 g, 46 mmol) was heated under reflux at 473 K. After concentration, the residue was chromatographed on silica gel column using petroleum ether (333–353 K) to give 3-(4-methoxyphenyl)isocoumarin, (3). 2-[2'-Oxoethyl-2'-(4'-methoxyphenyl)]benzoic acid, (4), was obtained by refluxing a solution of (3) (4 g, 15.9 mmol) in ethanol (200 ml) and potassium hydroxide (5%, 200 ml) for 4 h. NaBH<sub>4</sub> (1.6 g) was added to a solution of (4) (4.81 g, 17.8 mmol) in sodium hydroxide (1%, 180 ml) and the resulting solution was stirred overnight at room temperature. After being acidified with HCl, the whole mixture was extracted with dichloromethane (2 × 15 ml). Usual work-up gave crude racemic hydroxy-acid, (5), which was dissolved in acetic anhydride (5 ml) and heated under reflux for 2 h to get the title compound (6). The crude compound was purified by column chromatography on silica gel with petroleum ether and recrystallized in ethanol.

#### Refinement

H9 atom was located in difference syntheses and refined as [C—H = 0.968 (15) Å;  $U_{\text{iso}}(\text{H}) = 0.0527 \text{ \AA}^2$ ]. The remaining H atoms were positioned geometrically, with C—H = 0.93, 0.97 and 0.96 Å for aromatic, methylene and methyl H, respectively, and constrained to ride on their parent atoms with  $U_{\text{iso}}(\text{H}) = xU_{\text{eq}}(\text{C})$ , where  $x = 1.5$  for methyl H and  $x = 1.2$  for all other H atoms.

## Figures

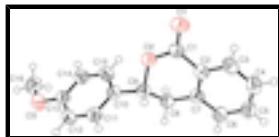


Fig. 1. The molecular structure of the title molecule, with the atom-numbering scheme. Displacement ellipsoids are drawn at the 50% probability level.

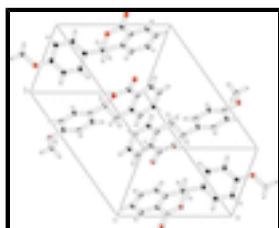


Fig. 2. A packing diagram of the title compound.

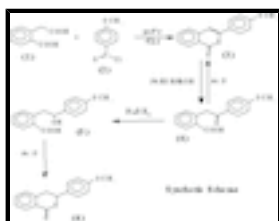


Fig. 3. Preparation of the title compound.

## 3-(4-Methoxyphenyl)isochroman-1-one

### Crystal data

$C_{16}H_{14}O_3$

$M_r = 254.27$

Monoclinic,  $P2_1/c$

Hall symbol: -P 2ybc

$a = 11.8933$  (7) Å

$b = 14.6874$  (9) Å

$c = 7.4521$  (4) Å

$\beta = 101.040$  (2)°

$V = 1277.66$  (13) Å<sup>3</sup>

$Z = 4$

$F(000) = 536$

$D_x = 1.332$  Mg m<sup>-3</sup>

Mo  $K\alpha$  radiation,  $\lambda = 0.71073$  Å

Cell parameters from 3302 reflections

$\theta = 2.2$ – $28.8$ °

$\mu = 0.09$  mm<sup>-1</sup>

$T = 296$  K

Prismatic, yellow

$0.24 \times 0.16 \times 0.12$  mm

### Data collection

Bruker KappaAPEXII CCD  
diffractometer

Radiation source: fine-focus sealed tube  
graphite

Detector resolution: 7.40 pixels mm<sup>-1</sup>  
 $\omega$  scans

Absorption correction: multi-scan  
(*SADABS*; Bruker, 2005)

$T_{\min} = 0.980$ ,  $T_{\max} = 0.996$

12426 measured reflections

3302 independent reflections

2275 reflections with  $I > 2\sigma(I)$

$R_{\text{int}} = 0.023$

$\theta_{\max} = 28.8$ °,  $\theta_{\min} = 2.2$ °

$h = -16 \rightarrow 16$

$k = -19 \rightarrow 19$

$l = -9 \rightarrow 10$

Refinement

Refinement on $F^2$	Primary atom site location: structure-invariant direct methods
Least-squares matrix: full	Secondary atom site location: difference Fourier map
$R[F^2 > 2\sigma(F^2)] = 0.042$	Hydrogen site location: inferred from neighbouring sites
$wR(F^2) = 0.117$	H atoms treated by a mixture of independent and constrained refinement
$S = 1.01$	$w = 1/[\sigma^2(F_o^2) + (0.0481P)^2 + 0.2427P]$
3302 reflections	where $P = (F_o^2 + 2F_c^2)/3$
175 parameters	$(\Delta/\sigma)_{\max} < 0.001$
0 restraints	$\Delta\rho_{\max} = 0.14 \text{ e } \text{\AA}^{-3}$
	$\Delta\rho_{\min} = -0.22 \text{ e } \text{\AA}^{-3}$

Special details

**Geometry.** Bond distances, angles *etc.* have been calculated using the rounded fractional coordinates. All su's are estimated from the variances of the (full) variance-covariance matrix. The cell e.s.d.'s are taken into account in the estimation of distances, angles and torsion angles

**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 ( $\text{\AA}^2$ )

	$x$	$y$	$z$	$U_{\text{iso}}^*/U_{\text{eq}}$
O1	0.56862 (10)	0.12610 (9)	-0.37797 (13)	0.0668 (4)
O2	0.68506 (8)	0.13694 (7)	-0.11396 (12)	0.0495 (3)
O3	1.15702 (8)	0.09384 (8)	0.43236 (15)	0.0642 (4)
C1	0.57829 (12)	0.13397 (9)	-0.21507 (18)	0.0460 (4)
C2	0.48087 (11)	0.13874 (8)	-0.11934 (17)	0.0421 (4)
C3	0.37156 (12)	0.15327 (9)	-0.2200 (2)	0.0512 (5)
C4	0.27961 (12)	0.15613 (10)	-0.1325 (2)	0.0575 (5)
C5	0.29633 (12)	0.14322 (10)	0.0534 (2)	0.0576 (5)
C6	0.40443 (12)	0.12764 (10)	0.1541 (2)	0.0519 (5)
C7	0.49834 (11)	0.12588 (8)	0.06918 (17)	0.0422 (4)
C8	0.61822 (11)	0.11135 (10)	0.17022 (17)	0.0465 (4)
C9	0.70140 (11)	0.16318 (9)	0.07877 (17)	0.0440 (4)
C10	0.82407 (11)	0.14481 (9)	0.16272 (17)	0.0438 (4)
C11	0.88770 (12)	0.20841 (10)	0.27598 (19)	0.0525 (5)
C12	0.99780 (12)	0.18919 (11)	0.3646 (2)	0.0573 (5)
C13	1.04728 (11)	0.10621 (10)	0.34024 (18)	0.0484 (4)
C14	0.98541 (12)	0.04249 (10)	0.2268 (2)	0.0542 (5)
C15	0.87482 (12)	0.06259 (10)	0.1397 (2)	0.0529 (5)

## supplementary materials

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C16	1.20764 (14)	0.00719 (12)	0.4207 (2)	0.0664 (6)
H3	0.36043	0.16105	-0.34599	0.0615*
H4	0.20631	0.16679	-0.19903	0.0690*
H5	0.23396	0.14501	0.11186	0.0691*
H6	0.41438	0.11826	0.27955	0.0623*
H8A	0.62502	0.13197	0.29549	0.0558*
H8B	0.63641	0.04693	0.17237	0.0558*
H9	0.6859 (12)	0.2278 (10)	0.0814 (19)	0.0527*
H11	0.85577	0.26487	0.29252	0.0630*
H12	1.03908	0.23245	0.44148	0.0687*
H14	1.01772	-0.01364	0.20892	0.0651*
H15	0.83351	0.01922	0.06319	0.0635*
H16A	1.28412	0.00733	0.49143	0.0995*
H16B	1.16306	-0.03839	0.46749	0.0995*
H16C	1.21001	-0.00605	0.29534	0.0995*

### Atomic displacement parameters ( $\text{\AA}^2$ )

	$U^{11}$	$U^{22}$	$U^{33}$	$U^{12}$	$U^{13}$	$U^{23}$
O1	0.0692 (7)	0.0953 (8)	0.0350 (5)	0.0026 (6)	0.0079 (5)	-0.0001 (5)
O2	0.0472 (5)	0.0654 (6)	0.0364 (5)	-0.0014 (4)	0.0096 (4)	0.0001 (4)
O3	0.0460 (6)	0.0818 (8)	0.0601 (6)	0.0054 (5)	-0.0018 (5)	-0.0082 (5)
C1	0.0519 (8)	0.0472 (7)	0.0378 (6)	0.0003 (6)	0.0062 (6)	0.0032 (5)
C2	0.0466 (7)	0.0375 (6)	0.0410 (7)	-0.0019 (5)	0.0057 (5)	0.0020 (5)
C3	0.0539 (8)	0.0477 (8)	0.0480 (8)	-0.0011 (6)	-0.0003 (6)	0.0040 (6)
C4	0.0443 (8)	0.0557 (8)	0.0688 (10)	0.0005 (6)	0.0013 (7)	0.0014 (7)
C5	0.0464 (8)	0.0594 (9)	0.0694 (10)	-0.0051 (6)	0.0170 (7)	-0.0047 (7)
C6	0.0513 (8)	0.0585 (8)	0.0477 (8)	-0.0056 (6)	0.0138 (6)	-0.0006 (6)
C7	0.0446 (7)	0.0408 (6)	0.0409 (7)	-0.0030 (5)	0.0072 (5)	0.0008 (5)
C8	0.0465 (7)	0.0574 (8)	0.0351 (6)	0.0001 (6)	0.0064 (5)	0.0041 (5)
C9	0.0465 (7)	0.0474 (7)	0.0375 (6)	-0.0012 (6)	0.0069 (5)	-0.0026 (5)
C10	0.0435 (7)	0.0493 (7)	0.0396 (6)	-0.0033 (5)	0.0106 (5)	-0.0023 (5)
C11	0.0501 (8)	0.0503 (8)	0.0565 (8)	-0.0003 (6)	0.0089 (6)	-0.0116 (6)
C12	0.0514 (8)	0.0615 (9)	0.0561 (9)	-0.0065 (7)	0.0033 (7)	-0.0168 (7)
C13	0.0416 (7)	0.0636 (9)	0.0401 (7)	-0.0007 (6)	0.0079 (5)	-0.0026 (6)
C14	0.0522 (8)	0.0547 (8)	0.0547 (8)	0.0056 (6)	0.0074 (7)	-0.0085 (7)
C15	0.0514 (8)	0.0522 (8)	0.0525 (8)	-0.0039 (6)	0.0033 (6)	-0.0126 (6)
C16	0.0561 (9)	0.0836 (12)	0.0574 (9)	0.0146 (8)	0.0060 (7)	0.0092 (8)

### Geometric parameters ( $\text{\AA}$ , $^\circ$ )

O1—C1	1.2028 (16)	C12—C13	1.380 (2)
O2—C1	1.3475 (17)	C13—C14	1.376 (2)
O2—C9	1.4636 (15)	C14—C15	1.383 (2)
O3—C13	1.3655 (17)	C3—H3	0.9300
O3—C16	1.418 (2)	C4—H4	0.9300
C1—C2	1.4741 (19)	C5—H5	0.9300
C2—C3	1.387 (2)	C6—H6	0.9300
C2—C7	1.3931 (18)	C8—H8A	0.9700

C3—C4	1.377 (2)	C8—H8B	0.9700
C4—C5	1.375 (2)	C9—H9	0.968 (15)
C5—C6	1.378 (2)	C11—H11	0.9300
C6—C7	1.385 (2)	C12—H12	0.9300
C7—C8	1.4947 (19)	C14—H14	0.9300
C8—C9	1.5102 (19)	C15—H15	0.9300
C9—C10	1.4977 (19)	C16—H16A	0.9600
C10—C11	1.3831 (19)	C16—H16B	0.9600
C10—C15	1.375 (2)	C16—H16C	0.9600
C11—C12	1.378 (2)		
O1…C16 <sup>i</sup>	3.367 (2)	C16…H14	2.5100
O3…C4 <sup>ii</sup>	3.4157 (18)	H3…O1	2.5800
O3…C3 <sup>ii</sup>	3.3841 (18)	H3…O3 <sup>vi</sup>	2.8300
O1…H3	2.5800	H3…C5 <sup>iv</sup>	3.0300
O1…H6 <sup>iii</sup>	2.8500	H4…O3 <sup>vi</sup>	2.9000
O1…H9 <sup>iv</sup>	2.608 (15)	H5…O3 <sup>x</sup>	2.8200
O1…H8A <sup>iii</sup>	2.6500	H5…C13 <sup>x</sup>	3.1000
O1…H16A <sup>i</sup>	2.8600	H6…O1 <sup>xi</sup>	2.8500
O2…H11 <sup>iv</sup>	2.6900	H6…H8A	2.4900
O2…H15	2.6400	H8A…O1 <sup>xi</sup>	2.6500
O2…H16C <sup>i</sup>	2.7800	H8A…H6	2.4900
O3…H3 <sup>ii</sup>	2.8300	H8B…C15	2.9000
O3…H4 <sup>ii</sup>	2.9000	H8B…C2 <sup>viii</sup>	3.0500
O3…H5 <sup>v</sup>	2.8200	H8B…C3 <sup>viii</sup>	2.9700
C3…C6 <sup>iv</sup>	3.395 (2)	H9…C2	2.915 (14)
C3…O3 <sup>vi</sup>	3.3841 (18)	H9…H11	2.3700
C3…C5 <sup>iv</sup>	3.465 (2)	H9…O1 <sup>vii</sup>	2.608 (15)
C4…O3 <sup>vi</sup>	3.4157 (18)	H9…C1 <sup>vii</sup>	2.967 (15)
C5…C3 <sup>vii</sup>	3.465 (2)	H11…H9	2.3700
C6…C3 <sup>vii</sup>	3.395 (2)	H11…O2 <sup>vii</sup>	2.6900
C16…O1 <sup>i</sup>	3.367 (2)	H14…C16	2.5100
C1…H9 <sup>iv</sup>	2.967 (15)	H14…H16B	2.3600
C2…H8B <sup>viii</sup>	3.0500	H14…H16C	2.2600
C2…H9	2.915 (14)	H15…O2	2.6400
C3…H16A <sup>vi</sup>	3.0700	H15…C4 <sup>viii</sup>	3.0000
C3…H8B <sup>viii</sup>	2.9700	H15…C5 <sup>viii</sup>	2.8800
C4…H15 <sup>viii</sup>	3.0000	H16A…C3 <sup>ii</sup>	3.0700
C5…H3 <sup>vii</sup>	3.0300	H16A…O1 <sup>i</sup>	2.8600
C5…H15 <sup>viii</sup>	2.8800	H16B…C14	2.7600
C13…H5 <sup>v</sup>	3.1000	H16B…H14	2.3600
C14…H16B	2.7600	H16B…C15 <sup>ix</sup>	3.0700
C14…H16C	2.7200	H16C…C14	2.7200

## supplementary materials

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C15···H16B <sup>ix</sup>	3.0700	H16C···H14	2.2600
C15···H8B	2.9000	H16C···O2 <sup>i</sup>	2.7800
C1—O2—C9	119.41 (10)	C4—C3—H3	120.00
C13—O3—C16	117.70 (12)	C3—C4—H4	120.00
O1—C1—O2	117.72 (13)	C5—C4—H4	120.00
O1—C1—C2	124.12 (13)	C4—C5—H5	120.00
O2—C1—C2	118.15 (11)	C6—C5—H5	120.00
C1—C2—C3	119.19 (12)	C5—C6—H6	120.00
C1—C2—C7	120.16 (12)	C7—C6—H6	120.00
C3—C2—C7	120.63 (12)	C7—C8—H8A	110.00
C2—C3—C4	119.71 (13)	C7—C8—H8B	110.00
C3—C4—C5	119.89 (14)	C9—C8—H8A	110.00
C4—C5—C6	120.77 (14)	C9—C8—H8B	110.00
C5—C6—C7	120.27 (13)	H8A—C8—H8B	108.00
C2—C7—C6	118.72 (12)	O2—C9—H9	106.7 (8)
C2—C7—C8	118.10 (12)	C8—C9—H9	110.0 (9)
C6—C7—C8	123.18 (12)	C10—C9—H9	110.0 (9)
C7—C8—C9	110.17 (11)	C10—C11—H11	120.00
O2—C9—C8	109.56 (10)	C12—C11—H11	120.00
O2—C9—C10	107.35 (10)	C11—C12—H12	120.00
C8—C9—C10	112.99 (11)	C13—C12—H12	120.00
C9—C10—C11	120.57 (12)	C13—C14—H14	120.00
C9—C10—C15	121.45 (12)	C15—C14—H14	120.00
C11—C10—C15	117.85 (13)	C10—C15—H15	119.00
C10—C11—C12	120.85 (14)	C14—C15—H15	119.00
C11—C12—C13	120.51 (14)	O3—C16—H16A	109.00
O3—C13—C12	116.22 (13)	O3—C16—H16B	109.00
O3—C13—C14	124.44 (13)	O3—C16—H16C	109.00
C12—C13—C14	119.34 (13)	H16A—C16—H16B	109.00
C13—C14—C15	119.47 (13)	H16A—C16—H16C	109.00
C10—C15—C14	121.98 (13)	H16B—C16—H16C	109.00
C2—C3—H3	120.00		
C9—O2—C1—O1	168.75 (13)	C5—C6—C7—C8	178.66 (13)
C9—O2—C1—C2	-12.42 (17)	C2—C7—C8—C9	32.06 (16)
C1—O2—C9—C8	47.27 (15)	C6—C7—C8—C9	-147.57 (13)
C1—O2—C9—C10	170.31 (11)	C7—C8—C9—O2	-55.05 (14)
C16—O3—C13—C12	175.67 (13)	C7—C8—C9—C10	-174.68 (11)
C16—O3—C13—C14	-4.8 (2)	O2—C9—C10—C11	134.82 (12)
O1—C1—C2—C3	-13.4 (2)	O2—C9—C10—C15	-49.38 (16)
O1—C1—C2—C7	164.54 (14)	C8—C9—C10—C11	-104.28 (15)
O2—C1—C2—C3	167.82 (12)	C8—C9—C10—C15	71.51 (16)
O2—C1—C2—C7	-14.21 (18)	C9—C10—C11—C12	175.05 (13)
C1—C2—C3—C4	178.77 (12)	C15—C10—C11—C12	-0.9 (2)
C7—C2—C3—C4	0.81 (19)	C9—C10—C15—C14	-175.39 (13)
C1—C2—C7—C6	-177.78 (12)	C11—C10—C15—C14	0.5 (2)
C1—C2—C7—C8	2.58 (17)	C10—C11—C12—C13	0.8 (2)
C3—C2—C7—C6	0.17 (18)	C11—C12—C13—O3	179.38 (13)
C3—C2—C7—C8	-179.48 (12)	C11—C12—C13—C14	-0.2 (2)

C2—C3—C4—C5	-1.0 (2)	O3—C13—C14—C15	-179.72 (13)
C3—C4—C5—C6	0.2 (2)	C12—C13—C14—C15	-0.2 (2)
C4—C5—C6—C7	0.8 (2)	C13—C14—C15—C10	0.0 (2)
C5—C6—C7—C2	-1.0 (2)		

Symmetry codes: (i)  $-x+2, -y, -z$ ; (ii)  $x+1, y, z+1$ ; (iii)  $x, y, z-1$ ; (iv)  $x, -y+1/2, z-1/2$ ; (v)  $x+1, y, z$ ; (vi)  $x-1, y, z-1$ ; (vii)  $x, -y+1/2, z+1/2$ ; (viii)  $-x+1, -y, -z$ ; (ix)  $-x+2, -y, -z+1$ ; (x)  $x-1, y, z$ ; (xi)  $x, y, z+1$ .

*Hydrogen-bond geometry* ( $\text{\AA}, ^\circ$ )

$D-H\cdots A$	$D-H$	$H\cdots A$	$D\cdots A$	$D-H\cdots A$
C8—H8B $\cdots$ Cg2 <sup>viii</sup>	0.97	2.94	3.8398 (16)	154
C16—H16B $\cdots$ Cg3 <sup>ix</sup>	0.96	2.89	3.7804 (17)	154

Symmetry codes: (viii)  $-x+1, -y, -z$ ; (ix)  $-x+2, -y, -z+1$ .

Fig. 1

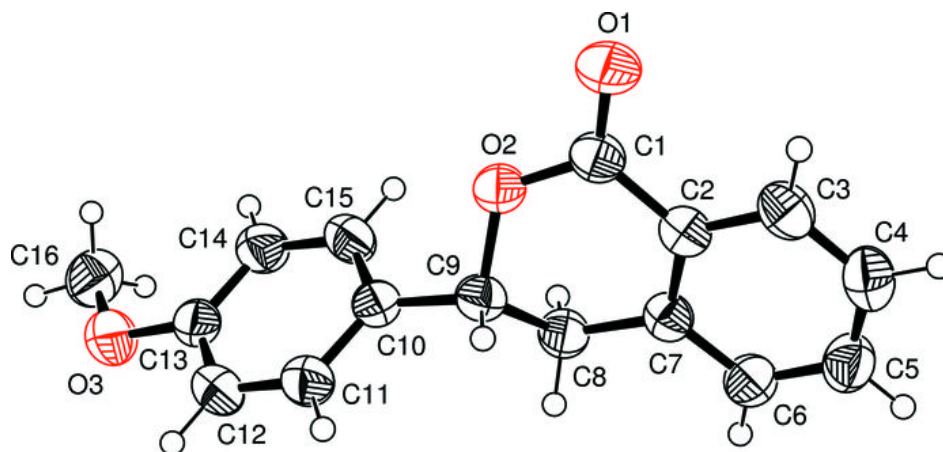


Fig. 2

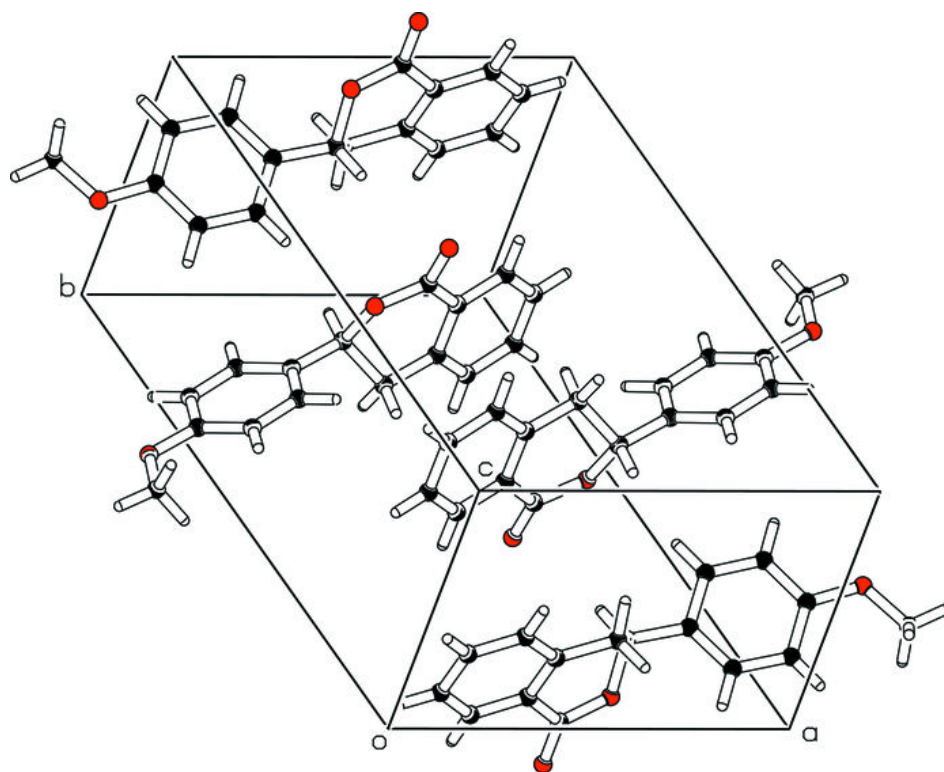


Fig. 3

