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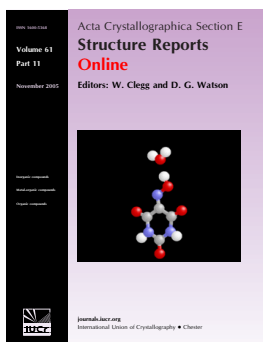
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# 1,3,5-Trichloro-2-methoxybenzene

**Sanjay Telu, Sean Parkin, Larry W. Robertson and Hans-Joachim Lehmler***Acta Cryst.* (2008). **E64**, o424

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## 1,3,5-Trichloro-2-methoxybenzene

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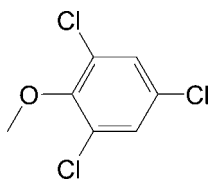
Received 14 December 2007; accepted 7 January 2008

Key indicators: single-crystal X-ray study;  $T = 90$  K; mean  $\sigma(\text{C}-\text{C}) = 0.003$  Å;  $R$  factor = 0.030;  $wR$  factor = 0.071; data-to-parameter ratio = 18.7.

The methoxy group of the title compound,  $\text{C}_7\text{H}_5\text{Cl}_3\text{O}$ , is rotated out of the plane of the aromatic ring system, with a dihedral angle of  $84.11(13)^\circ$ , due to the two bulky *ortho*-chloro substituents.

### Related literature

For similar structures of anisoles with two *ortho*-chloro substituents, see: Rissanen *et al.* (1987); Weller & Gerstner (1995); Wieczorek (1980). For other related literature, see: Brownlee *et al.* (1993); Curtis *et al.* (1972); Iimura *et al.* (1984); Kolehmainen & Knuutinen (1983); Oswald *et al.* (2005); Pereira *et al.* (2000); Rissanen *et al.* (1988); Vlachos *et al.* (2007); Zhang *et al.* (2006).



### Experimental

#### Crystal data

$\text{C}_7\text{H}_5\text{Cl}_3\text{O}$   
 $M_r = 211.46$   
Monoclinic,  $P2_1/n$   
 $a = 14.7538(5)$  Å  
 $b = 3.9846(2)$  Å  
 $c = 15.4810(7)$  Å  
 $\beta = 115.5031(19)^\circ$

$V = 821.42(6)$  Å<sup>3</sup>  
 $Z = 4$   
Mo  $K\alpha$  radiation  
 $\mu = 1.05$  mm<sup>-1</sup>  
 $T = 90.0(2)$  K  
 $0.25 \times 0.25 \times 0.12$  mm

#### Data collection

Nonius KappaCCD area-detector diffractometer  
Absorption correction: multi-scan (SCALEPACK; Otwinowski & Minor, 1997)  
 $T_{\min} = 0.780$ ,  $T_{\max} = 0.885$   
13489 measured reflections  
1888 independent reflections  
1558 reflections with  $I > 2\sigma(I)$   
 $R_{\text{int}} = 0.020$

#### Refinement

$R[F^2 > 2\sigma(F^2)] = 0.030$   
 $wR(F^2) = 0.070$   
 $S = 1.13$   
1888 reflections  
101 parameters  
H-atom parameters constrained  
 $\Delta\rho_{\text{max}} = 0.35$  e Å<sup>-3</sup>  
 $\Delta\rho_{\text{min}} = -0.31$  e Å<sup>-3</sup>

Data collection: COLLECT (Nonius, 1998); cell refinement: SCALEPACK (Otwinowski & Minor, 1997); data reduction: DENZO-SMN (Otwinowski & Minor, 1997); program(s) used to solve structure: SHELXS97 (Sheldrick, 2008); program(s) used to refine structure: SHELXL97 (Sheldrick, 2008); molecular graphics: XP in SHELXTL (Sheldrick, 1994); software used to prepare material for publication: SHELX97 and local procedures.

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Supplementary data and figures for this paper are available from the IUCr electronic archives (Reference: OM2200).

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## **supplementary materials**

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## 1,3,5-Trichloro-2-methoxybenzene

S. Telu, S. Parkin, L. W. Robertson and H.-J. Lehmler

### Comment

Chlorinated anisoles are persistent environmental pollutants known for their toxicity and tendency to bioaccumulate (Brownlee *et al.*, 1993). Especially anisoles with chlorine substitution in the 2- and 6- position are a well known cause for off-flavors in water, fish and chicken eggs (Brownlee *et al.*, 1993; Curtis *et al.*, 1972; Zhang *et al.*, 2006). The title compound is a major contributor to "cork taint", a well known off-flavor in the wine industry (Pereira *et al.*, 2000; Vlachos *et al.*, 2007). We herein report the molecular structure of the title compound to aid in structure activity relationship (SAR) studies of the toxicity of chlorinated anisoles.

It has been reported that *ortho*-disubstitution causes drastic changes in the spatial arrangement of the methoxy group of chlorinated anisols, both in solution and in the solid state (Kolehmainen & Knuutinen, 1983; Rissanen *et al.*, 1987). In the title compound, the methoxy group was rotated out of the plane of the phenyl ring with a dihedral angle of 84.11 (13)°. The angle was calculated between the plane of the benzene ring (C1 through C6) and the methoxy group (atoms C1,O1,C7). Comparable anisoles have similar dihedral angles ranging from 76.8 to 92.1° (Imura *et al.*, 1984; Rissanen *et al.*, 1987; Rissanen *et al.*, 1988; Weller & Gerstner, 1995; Wieczorek, 1980). In contrast, the methoxy group of non-*ortho* substituted anisoles lies within the plane of the phenyl ring.

The O—CH<sub>3</sub> bond length of the title compound (O1—C7: 1.446 (2) Å) was larger compared to non-*ortho* substituted anisoles, which have a mean bond length of 1.422 Å. Furthermore, the C<sub>Ar</sub>—O—CH<sub>3</sub> bond angle of the title compound (C1—O1—C7: 114.60 (15)°) was smaller compared to non-*ortho* substituted anisoles, which is on average 117.73°. Chlorine disubstitution *ortho* to an aromatic methoxy group had a similar effect on the O—CH<sub>3</sub> bond length and the C<sub>Ar</sub>—O—CH<sub>3</sub> bond angles in several structurally related compounds (Imura *et al.*, 1984; Rissanen *et al.*, 1987; Rissanen *et al.*, 1988; Weller & Gerstner, 1995; Wieczorek, 1980).

Ortho disubstitution forced the methoxy group out of the plane of the aromatic ring, so that the CH<sub>3</sub> group (C7) of the title compound was located 1.192 (4) Å above and the O1 atom -0.079 (3) Å below the calculated least-squares plane of the benzene ring. In other, structurally related compounds with *ortho* dichloro substitution, the O atoms are positioned 0.030 to 0.133 Å below and the methoxy C atoms 1.119 to 1.252 Å above the calculated last-squares plane (Imura *et al.*, 1984; Rissanen *et al.*, 1987; Rissanen *et al.*, 1988; Weller & Gerstner, 1995; Wieczorek, 1980). Overall, this difference in the spatial arrangement of the methoxy group is thought to explain the off-flavor of anisoles with *ortho* disubstitution.

### Experimental

Crystals of the title compound suitable for crystal structure analysis were obtained from TCI America (Portland, Oregon, USA.).

## Refinement

H atoms were found in difference Fourier maps and subsequently placed in idealized positions with constrained C—H distances of 0.98 Å ( $C_{Me}H$ ) and 0.95 Å ( $C_{Ar}H$ ) with  $U_{iso}(H)$  values set to either  $1.5U_{eq}(C_{Me}H)$  or  $1.2U_{eq}(C_{Ar}H)$  of the attached C atom respectively.

## Figures

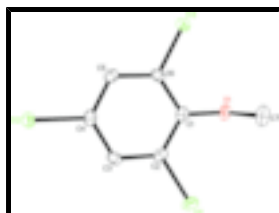


Fig. 1. View of the title compound showing the atom-labelling scheme. Displacement ellipsoids are drawn at the 50% probability level.

## 1,3,5-Trichloro-2-methoxybenzene

### Crystal data

$C_7H_5Cl_3O$

$M_r = 211.46$

Monoclinic,  $P2_1/n$

Hall symbol:  $-P\ 2_1/n$

$a = 14.7538(5)$  Å

$b = 3.9846(2)$  Å

$c = 15.4810(7)$  Å

$\beta = 115.5031(19)^\circ$

$V = 821.42(6)$  Å<sup>3</sup>

$Z = 4$

$F_{000} = 424$

$D_x = 1.710$  Mg m<sup>-3</sup>

Mo  $K\alpha$  radiation

$\lambda = 0.71073$  Å

Cell parameters from 2156 reflections

$\theta = 1.0\text{--}27.5^\circ$

$\mu = 1.05$  mm<sup>-1</sup>

$T = 90.0(2)$  K

Block, colourless

$0.25 \times 0.25 \times 0.12$  mm

### Data collection

Nonius KappaCCD area-detector diffractometer

Radiation source: fine-focus sealed tube

Monochromator: graphite

Detector resolution: 18 pixels mm<sup>-1</sup>

$T = 90.0(2)$  K

$\omega$  scans at fixed  $\chi = 55^\circ$

Absorption correction: multi-scan

(SCALEPACK; Otwinowski & Minor, 1997)

$T_{min} = 0.780$ ,  $T_{max} = 0.885$

13489 measured reflections

1888 independent reflections

1558 reflections with  $I > 2\sigma(I)$

$R_{int} = 0.020$

$\theta_{max} = 27.5^\circ$

$\theta_{min} = 1.6^\circ$

$h = -18 \rightarrow 19$

$k = -5 \rightarrow 5$

$l = -20 \rightarrow 19$

Refinement

Refinement on $F^2$	Secondary atom site location: difference Fourier map
Least-squares matrix: full	Hydrogen site location: inferred from neighbouring sites
$R[F^2 > 2\sigma(F^2)] = 0.030$	H-atom parameters constrained
$wR(F^2) = 0.070$	$w = 1/[\sigma^2(F_o^2) + (0.0244P)^2 + 0.6499P]$
$S = 1.13$	where $P = (F_o^2 + 2F_c^2)/3$
1888 reflections	$(\Delta/\sigma)_{\max} = 0.001$
101 parameters	$\Delta\rho_{\max} = 0.35 \text{ e } \text{\AA}^{-3}$
Primary atom site location: structure-invariant direct methods	$\Delta\rho_{\min} = -0.31 \text{ e } \text{\AA}^{-3}$
	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  $F^2 > 2\sigma(F^2)$  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 ( $\text{\AA}^2$ )

	$x$	$y$	$z$	$U_{\text{iso}}^*/U_{\text{eq}}$
Cl1	0.65436 (3)	0.31728 (13)	0.83292 (3)	0.02202 (13)
Cl2	0.36963 (3)	0.90882 (13)	0.91077 (3)	0.01965 (12)
Cl3	0.32350 (3)	0.84337 (13)	0.54775 (3)	0.02053 (13)
O1	0.51708 (9)	0.4751 (4)	0.63059 (9)	0.0189 (3)
C1	0.48575 (13)	0.5887 (5)	0.69659 (13)	0.0161 (4)
C2	0.54027 (13)	0.5242 (5)	0.79404 (13)	0.0161 (4)
C3	0.50643 (13)	0.6209 (5)	0.86111 (13)	0.0162 (4)
H3	0.5447	0.5753	0.9273	0.019*
C4	0.41531 (14)	0.7858 (5)	0.82874 (13)	0.0164 (4)
C5	0.35818 (13)	0.8564 (5)	0.73309 (13)	0.0158 (4)
H5	0.2958	0.9705	0.7126	0.019*
C6	0.39426 (13)	0.7561 (5)	0.66748 (12)	0.0147 (4)
C7	0.58710 (16)	0.6940 (6)	0.61549 (15)	0.0253 (5)
H7A	0.5565	0.9156	0.5951	0.038*
H7B	0.6039	0.5988	0.5659	0.038*
H7C	0.6483	0.7159	0.6753	0.038*

## supplementary materials

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

	$U^{11}$	$U^{22}$	$U^{33}$	$U^{12}$	$U^{13}$	$U^{23}$
C11	0.0146 (2)	0.0252 (3)	0.0231 (2)	0.00407 (19)	0.00509 (19)	-0.0021 (2)
C12	0.0178 (2)	0.0266 (3)	0.0163 (2)	0.00018 (18)	0.00906 (18)	-0.00204 (18)
C13	0.0190 (2)	0.0279 (3)	0.0139 (2)	0.00294 (19)	0.00632 (18)	0.00251 (18)
O1	0.0171 (7)	0.0230 (8)	0.0205 (7)	-0.0022 (6)	0.0119 (6)	-0.0046 (6)
C1	0.0160 (9)	0.0152 (10)	0.0188 (9)	-0.0037 (7)	0.0092 (7)	-0.0036 (7)
C2	0.0123 (8)	0.0142 (10)	0.0208 (9)	-0.0003 (7)	0.0061 (7)	-0.0008 (7)
C3	0.0152 (9)	0.0177 (10)	0.0129 (8)	-0.0030 (7)	0.0035 (7)	-0.0009 (7)
C4	0.0177 (9)	0.0160 (10)	0.0184 (9)	-0.0037 (7)	0.0106 (8)	-0.0031 (7)
C5	0.0128 (8)	0.0155 (10)	0.0192 (9)	-0.0002 (7)	0.0069 (7)	0.0005 (7)
C6	0.0138 (9)	0.0168 (10)	0.0120 (8)	-0.0023 (7)	0.0040 (7)	0.0002 (7)
C7	0.0273 (11)	0.0255 (11)	0.0310 (11)	-0.0054 (9)	0.0201 (9)	-0.0052 (9)

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

C1—O1	1.367 (2)	C13—C6	1.7264 (18)
C1—C2	1.394 (3)	C4—C5	1.381 (3)
C1—C6	1.395 (3)	C5—C6	1.393 (3)
C2—C3	1.386 (3)	C5—H5	0.9500
C2—C11	1.7329 (19)	O1—C7	1.446 (2)
C3—C4	1.382 (3)	C7—H7A	0.9800
C3—H3	0.9500	C7—H7B	0.9800
C12—C4	1.7447 (18)	C7—H7C	0.9800
O1—C1—C2	121.63 (16)	C4—C5—H5	120.8
O1—C1—C6	120.55 (16)	C6—C5—H5	120.8
C2—C1—C6	117.70 (16)	C5—C6—C1	121.46 (16)
C3—C2—C1	122.21 (17)	C5—C6—C13	118.74 (14)
C3—C2—C11	118.77 (14)	C1—C6—C13	119.79 (14)
C1—C2—C11	119.02 (14)	C1—O1—C7	114.60 (15)
C4—C3—C2	117.92 (17)	O1—C7—H7A	109.5
C4—C3—H3	121.0	O1—C7—H7B	109.5
C2—C3—H3	121.0	H7A—C7—H7B	109.5
C5—C4—C3	122.34 (17)	O1—C7—H7C	109.5
C5—C4—C12	118.33 (14)	H7A—C7—H7C	109.5
C3—C4—C12	119.33 (14)	H7B—C7—H7C	109.5
C4—C5—C6	118.36 (17)		
O1—C1—C2—C3	-176.17 (17)	C12—C4—C5—C6	-179.55 (14)
C6—C1—C2—C3	-0.1 (3)	C4—C5—C6—C1	-0.1 (3)
O1—C1—C2—C11	4.0 (3)	C4—C5—C6—C13	-179.94 (14)
C6—C1—C2—C11	-179.91 (14)	O1—C1—C6—C5	176.18 (17)
C1—C2—C3—C4	0.1 (3)	C2—C1—C6—C5	0.1 (3)
C11—C2—C3—C4	179.96 (15)	O1—C1—C6—C13	-4.0 (3)
C2—C3—C4—C5	-0.2 (3)	C2—C1—C6—C13	179.92 (14)
C2—C3—C4—C12	179.51 (15)	C2—C1—O1—C7	-86.2 (2)
C3—C4—C5—C6	0.1 (3)	C6—C1—O1—C7	97.9 (2)

Fig. 1

