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4,4′-[(5-Carboxy-1,3-phenylene)bis(oxy)]-dibenzoic acid

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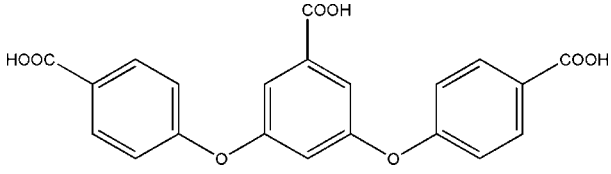
Key indicators: single-crystal X-ray study; T = 293 K; mean σ(C–C) = 0.002 Å; R factor = 0.046; wR factor = 0.121; data-to-parameter ratio = 15.5.

In the title compound, $C_{21}H_{14}O_8$, the central benzene ring makes dihedral angles of 77.8 (6) and 75.9 (5)° with the outer benzene rings. In the crystal, molecules are linked by O—H···O hydrogen bonds involving carboxyl groups, forming one-dimensional ladders. Two-dimensional layers are formed by interpenetration of these one-dimensional ladders.

Related literature

For general background, see: Moulton & Zaworotko,(2001); Kitagawa *et al.*,(2001); Lee *et al.*,(2009); Robin & Fromm, (2006). For the preparation of title compound, see: Neogi *et al.*(2009). For related structares, see: Lama *et al.*(2010); Pan *et al.* (2007).



Experimental

Crystal data

$C_{21}H_{14}O_8$
M_r = 394.32
Monoclinic, $C2/c$
a = 17.235 (3) Å
b = 13.419 (3) Å
c = 15.586 (3) Å
β = 96.24 (3)°
V = 3583.3 (12) Å^3
Z = 8
Mo $K\alpha$ radiation
μ = 0.11 mm^{-1}
T = 293 K
0.33 × 0.29 × 0.25 mm

Data collection

Bruker SMART CCD area-detector diffractometer
Absorption correction: multi-scan (*SADABS*; Bruker, 2003)
T_{min} = 0.316, T_{max} = 0.622
16922 measured reflections
4073 independent reflections
2862 reflections with $I > 2\sigma(I)$
R_{int} = 0.046

Refinement

$R[F^2 > 2\sigma(F^2)]$ = 0.046
$wR(F^2)$ = 0.121
S = 1.09
4073 reflections
262 parameters
H-atom parameters constrained
$\Delta\rho_{max}$ = 0.28 e Å^{-3}
$\Delta\rho_{min}$ = −0.22 e Å^{-3}

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

D—H···A	D—H	H···A	D···A	D—H···A
O3—H6···O4^{i}	0.82	1.87	2.6919 (16)	176.4
O6—H3···O8^{ii}	0.82	1.85	2.6615 (18)	169.9
O7—H8···O5^{iii}	0.82	1.82	2.6307 (18)	167.2

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

Data collection: *SMART* (Bruker, 2001); cell refinement: *SAINT-Plus* (Bruker, 2003); data reduction: *SAINT-Plus*; program(s) used to solve structure: *SHELXS97* (Sheldrick, 2008); program(s) used to refine structure: *SHELXL97* (Sheldrick, 2008); molecular graphics: *SHELXP97* (Sheldrick, 2008); software used to prepare material for publication: *SHELXTL*.

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Supplementary data and figures for this paper are available from the IUCr electronic archives (Reference: BQ2343).

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supplementary materials

Acta Cryst. (2012). E**68**, o1243 [doi:10.1107/S1600536812012275]

4,4′-[(5-Carboxy-1,3-phenylene)bis(oxy)]dibenzoic acid

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Comment

As a new kind of functional molecular materials, metal-organic frameworks have received extensive attention for their potential applications in gas storage, catalysis, optoelectronics, sensors, magnetism, luminescence, porous materials and so on. (Moulton & Zaworotko, 2001; Kitagawa *et al.*, 2001; Lee *et al.*, 2009). Organic molecules with O– and N-donors can be used as organic linkers in these coordination polymers (Robin & Fromm, 2006). In fact, there are many organic ligands which are linked by ether bond (Lama *et al.* 2010; Pan *et al.*, 2007). Here, we report the crystal structure of the title compound.

In the crystal structure, two benzene rings, *β*(composed of C_8—C_{13}) and *γ* (composed of C_{15}—C_{20}) are connected to the center ring (*α*, composed of C_1—C_6) by ether bond. The dihedral angle between *α* and *β* is 77.8 (6)°, and between *α* and *γ* is 75.9 (5)° (Fig. 1). Strong intermolecular O—H···O hydrogen bonds are formed between the carboxylic acid groups of neighboring molecules (Table 1), which link the molecules to one-dimensional supra-molecular ladder (Fig. 2). The interpenetration among the one-dimensional molecular ladders which are parallel produce two-dimensional layer (Fig. 3).

Experimental

The title compound was synthesized by a modified literature method (Neogi *et al.*2009). Methyl 3,5-dihydroxylbenzoate (1.68 g,10 mmol) was dissolved in DMF (50 ml). To this solution was added K_2CO_3 (7 g,51 mmol) and 4-fluorobenzonitrile (2.4 g,20 mmol). The mixture was heated under reflux for 2 days. The resulting solution was poured in 250 ml ice-cold water and kept over-night. The yellow compound was filtered and washed several times with water. The yellow compound (3.73 g, 10 mmol) was allowed to reflux with 6 N NaOH solution (50 ml) for 12 h, cooled to room temperature and acidified with HCl (6 N). Colorless crystalline product was obtained and isolated by filtration, washed with water and dried in vacuum. $Zn(NO_3)_2$ (0.075 g,0.25 mmol), 4,4′-(5-carboxy-1,3-phenylene)bis(oxy)dibenzoic acid (0.098 g,0.25 mmol), were mixed in water (5 ml). The mixture were placed in a 25 ml Teflon-lined stainless steel autoclave and heated autogenously under pressure for 2 d at 393 K. After cooling to room temperature, the block-shaped colourless crystals were obtained.

Refinement

All hydrogen atoms bonded to O and C were fixed in ideal positions, with C—H = 0.93 (aromatic) and O—H = 0.82 Å, and treated as riding on their parent atoms with $\boldsymbol{U}_{iso}$(H)=0.08 Å^2.

Computing details

Data collection: *SMART* (Bruker, 2001); cell refinement: *SAINT-Plus* (Bruker, 2003); data reduction: *SAINT-Plus* (Bruker, 2003); program(s) used to solve structure: *SHELXS97* (Sheldrick, 2008); program(s) used to refine structure: *SHELXL97* (Sheldrick, 2008); molecular graphics: SHELXP97 (Sheldrick, 2008); software used to prepare material for publication: *SHELXTL* (Sheldrick, 2008).

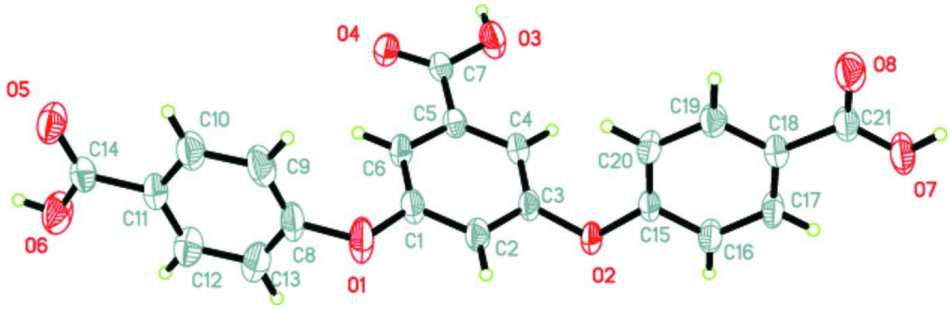

Figure 1

The molecular structure of the title compound with the atom-numbering scheme, with 50% probability displacement ellipsoids.

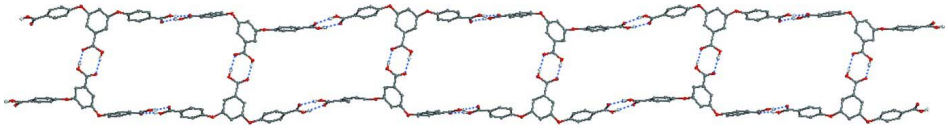

Figure 2

The packing of title compound, showing one ladder of molecules connected by O—H···O hydrogen bonds (dashed lines). H atoms not involved in hydrogen bonding have been omitted for clarity.

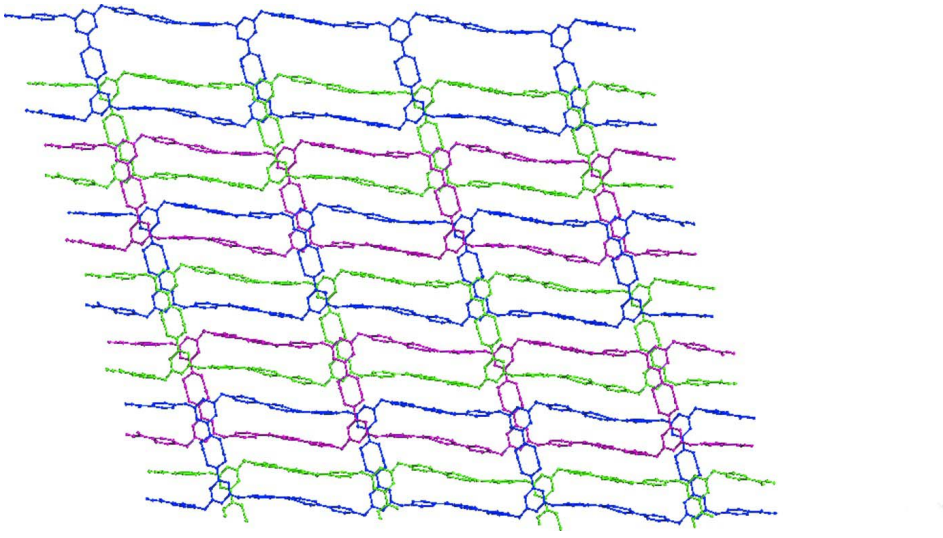

Figure 3

The interpenetration among the one-dimensional ladders, showing two-dimensional layer. H atoms not involved in hydrogen bonding have been omitted for clarity.

4,4′-[(5-Carboxy-1,3-phenylene)bis(oxy)]dibenzoic acid

Crystal data

$C_{21}H_{14}O_8$
$M_r = 394.32$
Monoclinic, $C2/c$
Hall symbol: -C 2yc
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$b = 13.419$ (3) Å
$c = 15.586$ (3) Å
$\beta = 96.24$ (3)°
$V = 3583.3$ (12) Å^3
$Z = 8$
$F(000) = 1632$
$D_x = 1.462$ Mg m^{-3}
Mo $K\alpha$ radiation, $\lambda = 0.71073$ Å
Cell parameters from 4073 reflections
$\theta = 3.0$–$27.5°$
$\mu = 0.11$ mm^{-1}
$T = 293$ K
Block, colourless
0.33 × 0.29 × 0.25 mm

Data collection

Bruker SMART CCD area-detector diffractometer
Radiation source: fine-focus sealed tube
Graphite monochromator
phi and ω scans
Absorption correction: multi-scan (*SADABS*; Bruker, 2003)
$T_{min} = 0.316$, $T_{max} = 0.622$
16922 measured reflections
4073 independent reflections
2862 reflections with $I > 2\sigma(I)$
$R_{int} = 0.046$
$\theta_{max} = 27.5°$, $\theta_{min} = 3.0°$
$h = -22 \rightarrow 22$
$k = -17 \rightarrow 17$
$l = -19 \rightarrow 20$

Refinement

Refinement on F^2
Least-squares matrix: full
$R[F^2 > 2\sigma(F^2)] = 0.046$
$wR(F^2) = 0.121$
$S = 1.09$
4073 reflections
262 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
$w = 1/[\sigma^2(F_o^2) + (0.0601P)^2 + 0.3436P]$ where $P = (F_o^2 + 2F_c^2)/3$
$(\Delta/\sigma)_{max} < 0.001$
$\Delta\rho_{max} = 0.28$ e Å^{-3}
$\Delta\rho_{min} = -0.22$ 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}
O1	0.16748 (7)	0.89288 (12)	−0.11510 (7)	0.0630 (4)
O2	−0.07879 (6)	0.73896 (9)	−0.14922 (7)	0.0404 (3)
O3	−0.06065 (6)	0.88133 (11)	0.15033 (7)	0.0503 (4)
H6	−0.0624	0.8964	0.2011	0.080*
O4	0.06498 (6)	0.92236 (9)	0.18120 (7)	0.0386 (3)

O5	0.44526 (7)	1.12170 (10)	0.10342 (8)	0.0533 (3)
O6	0.48890 (7)	0.96517 (11)	0.10924 (10)	0.0622 (4)
H3	0.5260	0.9916	0.1379	0.080*
O7	−0.42509 (6)	0.81151 (10)	−0.31119 (8)	0.0517 (3)
H8	−0.4653	0.8389	−0.3320	0.080*
O8	−0.38899 (7)	0.96958 (10)	−0.28672 (8)	0.0510 (3)
C1	0.10101 (8)	0.87058 (14)	−0.07678 (10)	0.0383 (4)
C2	0.04339 (9)	0.82152 (13)	−0.12950 (10)	0.0389 (4)
H11A	0.0509	0.8057	−0.1861	0.080*
C3	−0.02518 (8)	0.79647 (12)	−0.09712 (10)	0.0335 (4)
C4	−0.03815 (8)	0.82052 (12)	−0.01343 (10)	0.0338 (4)
H9A	−0.0852	0.8047	0.0074	0.080*
C5	0.02053 (8)	0.86866 (12)	0.03845 (9)	0.0305 (3)
C6	0.09117 (8)	0.89385 (12)	0.00791 (9)	0.0340 (4)
H13A	0.1305	0.9254	0.0435	0.080*
C7	0.01100 (8)	0.89365 (12)	0.13006 (10)	0.0318 (3)
C8	0.23310 (9)	0.92824 (16)	−0.06384 (10)	0.0457 (5)
C9	0.24061 (9)	1.02836 (17)	−0.04567 (12)	0.0520 (5)
H3A	0.2011	1.0727	−0.0652	0.080*
C10	0.30817 (10)	1.06221 (15)	0.00240 (12)	0.0475 (4)
H2A	0.3139	1.1296	0.0154	0.080*
C11	0.36696 (8)	0.99590 (14)	0.03100 (10)	0.0379 (4)
C12	0.35866 (10)	0.89573 (15)	0.01005 (12)	0.0491 (5)
H6A	0.3984	0.8512	0.0282	0.080*
C13	0.29151 (10)	0.86173 (16)	−0.03783 (11)	0.0510 (5)
H5A	0.2860	0.7946	−0.0522	0.080*
C14	0.43795 (9)	1.03153 (14)	0.08419 (11)	0.0393 (4)
C15	−0.15218 (8)	0.77687 (12)	−0.17470 (9)	0.0310 (3)
C16	−0.20659 (9)	0.70800 (13)	−0.21092 (11)	0.0396 (4)
H17A	−0.1938	0.6408	−0.2131	0.080*
C17	−0.27973 (9)	0.74089 (13)	−0.24351 (11)	0.0407 (4)
H16A	−0.3166	0.6952	−0.2673	0.080*
C18	−0.29914 (8)	0.84138 (13)	−0.24126 (10)	0.0340 (4)
C19	−0.24389 (9)	0.90878 (13)	−0.20428 (10)	0.0380 (4)
H20A	−0.2565	0.9761	−0.2023	0.080*
C20	−0.17048 (9)	0.87698 (13)	−0.17049 (10)	0.0367 (4)
H19A	−0.1340	0.9223	−0.1453	0.080*
C21	−0.37538 (9)	0.87785 (13)	−0.28147 (10)	0.0376 (4)

Atomic displacement parameters (Å²)

	U^{11}	U^{22}	U^{33}	U^{12}	U^{13}	U^{23}
O1	0.0313 (6)	0.1242 (13)	0.0339 (6)	−0.0307 (7)	0.0050 (5)	−0.0151 (7)
O2	0.0226 (5)	0.0450 (7)	0.0501 (7)	0.0008 (5)	−0.0117 (5)	−0.0106 (5)
O3	0.0275 (6)	0.0879 (10)	0.0359 (6)	−0.0055 (6)	0.0051 (5)	−0.0026 (6)
O4	0.0332 (6)	0.0474 (7)	0.0347 (6)	−0.0087 (5)	0.0009 (5)	−0.0024 (5)
O5	0.0394 (7)	0.0487 (9)	0.0676 (9)	−0.0075 (6)	−0.0131 (6)	−0.0023 (6)
O6	0.0382 (7)	0.0610 (10)	0.0805 (9)	0.0024 (7)	−0.0247 (6)	−0.0065 (7)
O7	0.0291 (6)	0.0479 (8)	0.0730 (9)	0.0008 (5)	−0.0179 (6)	−0.0011 (6)
O8	0.0401 (7)	0.0423 (8)	0.0657 (8)	0.0062 (6)	−0.0167 (6)	−0.0016 (6)

C1	0.0226 (7)	0.0587 (12)	0.0329 (8)	−0.0054 (7)	0.0002 (6)	0.0002 (8)
C2	0.0266 (8)	0.0567 (12)	0.0317 (8)	−0.0007 (7)	−0.0045 (6)	−0.0038 (7)
C3	0.0207 (7)	0.0385 (9)	0.0386 (8)	0.0009 (6)	−0.0088 (6)	−0.0009 (7)
C4	0.0215 (7)	0.0412 (10)	0.0378 (8)	−0.0012 (6)	−0.0016 (6)	0.0027 (7)
C5	0.0242 (7)	0.0348 (9)	0.0316 (7)	0.0010 (6)	−0.0014 (6)	0.0048 (6)
C6	0.0242 (7)	0.0442 (10)	0.0320 (8)	−0.0050 (7)	−0.0036 (6)	0.0010 (7)
C7	0.0254 (7)	0.0353 (9)	0.0340 (8)	0.0012 (6)	0.0000 (6)	0.0043 (6)
C8	0.0257 (8)	0.0812 (15)	0.0303 (8)	−0.0177 (9)	0.0038 (7)	−0.0046 (9)
C9	0.0272 (8)	0.0777 (15)	0.0488 (10)	0.0032 (9)	−0.0062 (7)	0.0007 (10)
C10	0.0311 (9)	0.0559 (12)	0.0537 (11)	−0.0020 (8)	−0.0039 (8)	−0.0007 (9)
C11	0.0259 (8)	0.0483 (11)	0.0383 (8)	−0.0064 (7)	−0.0018 (6)	0.0044 (8)
C12	0.0402 (9)	0.0508 (12)	0.0537 (11)	−0.0071 (8)	−0.0072 (8)	0.0019 (9)
C13	0.0452 (10)	0.0584 (13)	0.0481 (10)	−0.0179 (9)	−0.0011 (9)	−0.0028 (9)
C14	0.0264 (8)	0.0454 (11)	0.0444 (9)	−0.0010 (7)	−0.0031 (7)	0.0028 (8)
C15	0.0211 (7)	0.0401 (9)	0.0307 (7)	−0.0023 (6)	−0.0026 (6)	0.0010 (7)
C16	0.0287 (8)	0.0352 (10)	0.0518 (10)	−0.0024 (7)	−0.0098 (7)	−0.0023 (7)
C17	0.0260 (8)	0.0401 (10)	0.0526 (10)	−0.0063 (7)	−0.0106 (7)	−0.0001 (8)
C18	0.0246 (7)	0.0408 (10)	0.0353 (8)	−0.0009 (7)	−0.0033 (6)	0.0021 (7)
C19	0.0325 (8)	0.0379 (10)	0.0412 (9)	0.0015 (7)	−0.0065 (7)	0.0004 (7)
C20	0.0289 (8)	0.0396 (10)	0.0394 (8)	−0.0070 (7)	−0.0065 (7)	−0.0021 (7)
C21	0.0265 (8)	0.0438 (11)	0.0406 (9)	−0.0007 (7)	−0.0043 (7)	−0.0004 (7)

Geometric parameters (Å, º)

O1—C1	1.3811 (18)	C8—C13	1.373 (3)
O1—C8	1.395 (2)	C8—C9	1.376 (3)
O2—C15	1.3810 (17)	C9—C10	1.391 (2)
O2—C3	1.3949 (18)	C9—H3A	0.9300
O3—C7	1.3179 (17)	C10—C11	1.385 (2)
O3—H6	0.8200	C10—H2A	0.9300
O4—C7	1.2199 (19)	C11—C12	1.387 (3)
O5—C14	1.250 (2)	C11—C14	1.481 (2)
O6—C14	1.281 (2)	C12—C13	1.384 (2)
O6—H3	0.8200	C12—H6A	0.9300
O7—C21	1.286 (2)	C13—H5A	0.9300
O7—H8	0.8200	C15—C20	1.383 (2)
O8—C21	1.254 (2)	C15—C16	1.392 (2)
C1—C6	1.385 (2)	C16—C17	1.379 (2)
C1—C2	1.385 (2)	C16—H17A	0.9300
C2—C3	1.376 (2)	C17—C18	1.391 (2)
C2—H11A	0.9300	C17—H16A	0.9300
C3—C4	1.385 (2)	C18—C19	1.392 (2)
C4—C5	1.384 (2)	C18—C21	1.476 (2)
C4—H9A	0.9300	C19—C20	1.384 (2)
C5—C6	1.396 (2)	C19—H20A	0.9300
C5—C7	1.493 (2)	C20—H19A	0.9300
C6—H13A	0.9300		
C1—O1—C8	119.01 (12)	C9—C10—H2A	119.9
C15—O2—C3	119.45 (12)	C10—C11—C12	119.59 (16)

C7—O3—H6	109.5	C10—C11—C14	120.24 (17)
C14—O6—H3	109.5	C12—C11—C14	120.17 (15)
C21—O7—H8	109.5	C13—C12—C11	120.33 (18)
O1—C1—C6	123.82 (14)	C13—C12—H6A	119.8
O1—C1—C2	114.92 (13)	C11—C12—H6A	119.8
C6—C1—C2	121.25 (14)	C8—C13—C12	119.21 (19)
C3—C2—C1	119.18 (14)	C8—C13—H5A	120.4
C3—C2—H11A	120.4	C12—C13—H5A	120.4
C1—C2—H11A	120.4	O5—C14—O6	123.49 (16)
C2—C3—C4	121.42 (14)	O5—C14—C11	120.18 (15)
C2—C3—O2	117.55 (13)	O6—C14—C11	116.32 (16)
C4—C3—O2	120.84 (13)	O2—C15—C20	123.41 (14)
C5—C4—C3	118.45 (13)	O2—C15—C16	115.32 (14)
C5—C4—H9A	120.8	C20—C15—C16	121.14 (14)
C3—C4—H9A	120.8	C17—C16—C15	119.03 (16)
C4—C5—C6	121.50 (14)	C17—C16—H17A	120.5
C4—C5—C7	120.94 (13)	C15—C16—H17A	120.5
C6—C5—C7	117.54 (14)	C16—C17—C18	120.92 (15)
C1—C6—C5	118.17 (14)	C16—C17—H16A	119.5
C1—C6—H13A	120.9	C18—C17—H16A	119.5
C5—C6—H13A	120.9	C17—C18—C19	119.00 (14)
O4—C7—O3	123.42 (14)	C17—C18—C21	121.08 (14)
O4—C7—C5	122.71 (13)	C19—C18—C21	119.84 (15)
O3—C7—C5	113.86 (13)	C20—C19—C18	120.88 (16)
C13—C8—C9	121.61 (16)	C20—C19—H20A	119.6
C13—C8—O1	118.08 (19)	C18—C19—H20A	119.6
C9—C8—O1	120.15 (18)	C15—C20—C19	119.02 (15)
C8—C9—C10	118.97 (17)	C15—C20—H19A	120.5
C8—C9—H3A	120.5	C19—C20—H19A	120.5
C10—C9—H3A	120.5	O8—C21—O7	122.91 (15)
C11—C10—C9	120.25 (19)	O8—C21—C18	120.30 (14)
C11—C10—H2A	119.9	O7—C21—C18	116.78 (15)

Hydrogen-bond geometry (Å, º)

D—H···*A*	*D*—H	H···*A*	*D*···*A*	*D*—H···*A*
O3—H6···O4^{i}	0.82	1.87	2.6919 (16)	176.4
O6—H3···O8^{ii}	0.82	1.85	2.6615 (18)	169.9
O7—H8···O5^{iii}	0.82	1.82	2.6307 (18)	167.2

Symmetry codes: (i) −*x*, *y*, −*z*+1/2; (ii) *x*+1, −*y*+2, *z*+1/2; (iii) *x*−1, −*y*+2, *z*−1/2.