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2′-Iodo-2,2″,3,3″,4,4″,5,5″,6,6″-decamethyl-1,1′:3′,1″-terphenyl chloroform monosolvate

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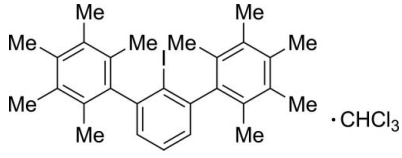
Key indicators: single-crystal X-ray study; T = 297 K; mean σ(C–C) = 0.007 Å; R factor = 0.066; wR factor = 0.137; data-to-parameter ratio = 19.1.

The title compound, $C_{28}H_{33}I{\cdot}CHCl_3$, forms dimers through C—I$\cdots\pi$ interactions. The crystal structure is consolidated by the presence of C—H$\cdots\pi$ interactions between the chloroform solvent and the main molecule.

Related literature

For the synthesis and spectroscopic characterization of 2′-iodo-2,2″,3,3″,4,4″,5,5″,6,6″-decamethyl-1,1′:3′,1″-terphenyl, see: Hino *et al.* (2005); Duttwyler *et al.* (2008). For related *m*-terphenyl iodides, see: Niemeyer (1998); Twamley *et al.* (2000); Zakharov *et al.* (2003). For general background to compounds with *m*-terphenyl substituents, see: Power (2004).



Experimental

Crystal data

$C_{28}H_{33}I{\cdot}CHCl_3$
M_r = 615.81
Monoclinic, $P2_1/n$
a = 12.0294 (10) Å
b = 16.0651 (13) Å
c = 15.3762 (12) Å
β = 103.385 (1)°
V = 2890.8 (4) Å^3
Z = 4
Mo $K\alpha$ radiation
μ = 1.40 mm^{-1}
T = 297 K
0.32 × 0.28 × 0.26 mm

Data collection

Bruker SMART CCD area-detector diffractometer
Absorption correction: multi-scan (*SADABS*; Bruker, 2000)
T_{min} = 0.663, T_{max} = 0.712
22910 measured reflections
5896 independent reflections
4698 reflections with $I > 2\sigma(I)$
R_{int} = 0.056

Refinement

$R[F^2 > 2\sigma(F^2)]$ = 0.066
$wR(F^2)$ = 0.137
S = 1.13
5896 reflections
308 parameters
H-atom parameters constrained
$\Delta\rho_{max}$ = 0.83 e Å^{-3}
$\Delta\rho_{min}$ = −0.68 e Å^{-3}

Table 1
C—H$\cdots\pi$ interactions (Å, °).

*Cg*2 and C3 are the centroids of the C7–C12 and C18–C23 benzene rings, respectively.

D—H$\cdots A$	D—H	H$\cdots A$	$D\cdots A$	D—H$\cdots A$
C26—H26*A*$\cdots$*Cg*2^{i}	0.96	3.86 (1)	2.97	155
C28—H28*A*$\cdots$*Cg*2^{ii}	0.96	3.53 (1)	2.87	127
C29—H29$\cdots$*Cg*3^{iii}	0.98	3.42 (1)	2.44	177

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

Table 2
C—I$\cdots\pi$ interactions (Å, °).

*Cg*3 is the centroid of the C18–C23 benzene ring.

Y—$X\cdots Cg$	Y—X	$X\cdots Cg$	$Y\cdots Cg$	Y—$X\cdots Cg$
C1—I1$\cdots$*Cg*3^{i}	2.099 (4)	3.975 (2)	6.026 (5)	164.67 (13)

Symmetry code: (i) $2-x, -y, 1-z$.

Data collection: *SMART* (Bruker, 2000); cell refinement: *SAINT-Plus* (Bruker, 2001); data reduction: *SAINT-Plus*; program(s) used to solve structure: *SHELXS97* (Sheldrick, 2008); program(s) used to refine structure: *SHELXL97* (Sheldrick, 2008); molecular graphics: *DIAMOND* (Brandenburg, 2009); software used to prepare material for publication: *publCIF* (Westrip, 2010) and *PLATON* (Spek, 2009).

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Supplementary data and figures for this paper are available from the IUCr electronic archives (Reference: PK2291).

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supporting information

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2′-Iodo-2,2″,3,3″,4,4″,5,5″,6,6″-decamethyl-1,1′:3′,1″-terphenyl chloroform monosolvate

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S1. Comment

m-Terphenyl substituents are sterrically crowded ligands used in stabilizing labile bonds and unusual geometries (Power, 2004).

We report herein the crystal structure of the title compound. The synthesis and the spectroscopic characterization of 2′-iodo-2,2″,3,3″,4,4″,5,5″,6,6″-decamethyl-1,1′:3′,1″-terphenyl were previously reported (Hino *et al.*, 2005; Duttwyler *et al.*, 2008).

The crystal structure (Fig. 1) of the title compound is similar to the structures of other 2,6-diarylphenyliodides. The C—I bond length [2.099 (4) Å] is slightly smaller than those found in 2,6-(2,4,6 - iPr$_3$C$_6$H$_2$)$_2$C$_6$H$_3$I [2.102 (6) Å] (Twamley *et al.*, 2000), 2,6-Ph$_2$C$_6$H$_3$I [2.122 (4) Å] (Niemeyer, 1998), and 2,6-Mes$_2$C$_6$H$_3$I [2.102 (5) Å] (Zakharov *et al.*, 2003).

Similar to the other 2,6-diarylphenyliodides, the dihedral angles between the flanking groups and the central benzene ring are close to 90° (Niemeyer, 1998; Twamley *et al.*, 2000; Zakharov *et al.*, 2003). This arrangement permits the presence of an intramolecular I···*Cg* contact [3.955 (1) Å]. The I···*Cg* interaction is reflected in the difference [Δ = 2.9°] at the C1 bonding angles.

In the crystal structure symmetry related molecules are linked into dimers through C—I···π interactions (Table 1 and Fig. 2). In addition there are C—H···π interactions (Table 2 and Fig. 2) that link the dimeric units and the solvent molecules into a three dimensional network.

S2. Experimental

The synthesis of 2′-iodo-2,2″,3,3″,4,4″,5,5″,6,6″-decamethyl-1,1′:3′,1″-terphenyl was carried out according to a previously described method (Hino *et al.*, 2005). Crystals of the title compound were obtained by slow evaporation of the solvent from a solution of 2′-iodo-2,2″,3,3″,4,4″,5,5″,6,6″-decamethyl-1,1′:3′,1″-terphenyl in chloroform.

S3. Refinement

Hydrogen atoms were placed in calculated positions with isotropic thermal parameters set at 1.2 times the carbon atoms directly attached for aromatic and methine hydrogen atoms and 1.5 for hydrogen atoms of the methyl groups. Methyl hydrogen atoms were allowed to rotate but not to tip.

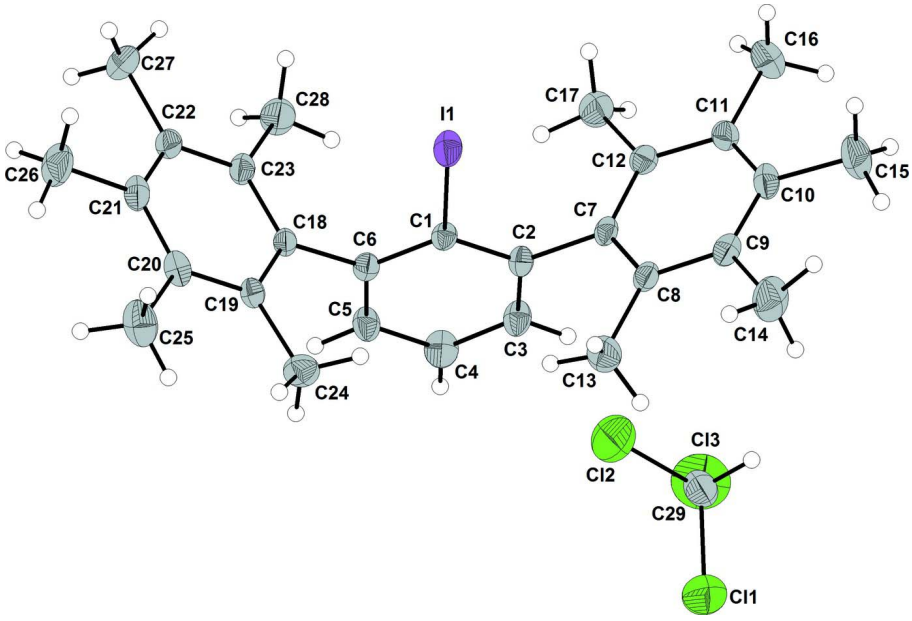

Figure 1

Crystal structure of the title compound with labelling and displacement ellipsoids of C, Cl and I atoms drawn at 25% probability level.

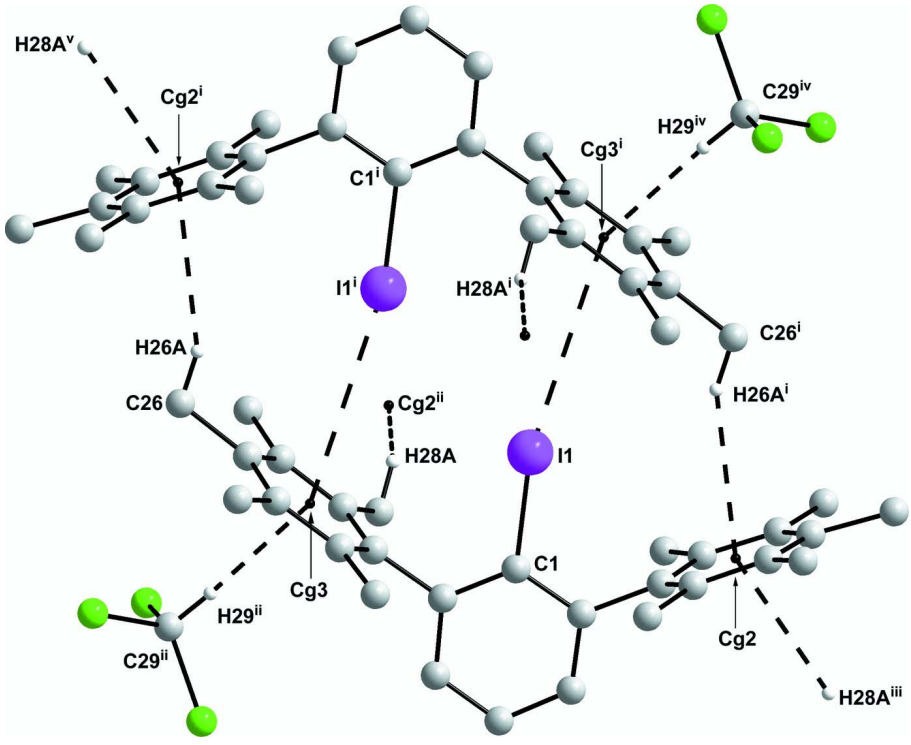

Figure 2

Intermolecular C—H···π and C—I···π interactions in the structure of the title compound. Symmetry codes: (i) 2 - *x*,-*y*,1 - *z*; (ii) 1/2 + *x*,1/2 - *y*,1/2 + *z*; (iii) -1/2 + *x*,1/2 - *y*,-1/2 + *z*; (iv) 3/2 - *x*, -1/2 + *y*, 1/2 - *z*; (v) 5/2 - *x*, -1/2 + *y*, 3/2 - *z*. *Cg*2 and *Cg*3 are the centroids of the benzene rings C7–C12 and C18–C23, respectively.

2′-Iodo-2,2″,3,3″,4,4″,5,5″,6,6″-decamethyl-1,1′:3′,1″-terphenyl chloroform monosolvate

Crystal data

$C_{28}H_{33}I·CHCl_3$
M_r = 615.81
Monoclinic, $P2_1/n$
Hall symbol: -P 2yn
a = 12.0294 (10) Å
b = 16.0651 (13) Å
c = 15.3762 (12) Å
β = 103.385 (1)°
V = 2890.8 (4) Å^3
Z = 4
$F(000)$ = 1248
D_x = 1.415 Mg m^{-3}
Melting point = 498–497 K
Mo $K\alpha$ radiation, λ = 0.71073 Å
Cell parameters from 3032 reflections
θ = 2.3–19.9°
μ = 1.40 mm^{-1}
T = 297 K
Block, colourless
0.32 × 0.28 × 0.26 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, 2000)
T_{min} = 0.663, T_{max} = 0.712
22910 measured reflections
5896 independent reflections
4698 reflections with $I > 2\sigma(I)$
R_{int} = 0.056
θ_{max} = 26.4°, θ_{min} = 1.9°
h = −15→15
k = −20→20
l = −19→19

Refinement

Refinement on F^2
Least-squares matrix: full
$R[F^2 > 2\sigma(F^2)]$ = 0.066
$wR(F^2)$ = 0.137
S = 1.13
5896 reflections
308 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
$w = 1/[\sigma^2(F_o^2) + (0.0465P)^2 + 3.6212P]$ where $P = (F_o^2 + 2F_c^2)/3$
$(\Delta/\sigma)_{max}$ = 0.031
$\Delta\rho_{max}$ = 0.83 e Å^{-3}
$\Delta\rho_{min}$ = −0.68 e Å^{-3}

Special details

Geometry. All s.u.'s (except the s.u. in the dihedral angle between two l.s. planes) are estimated using the full covariance matrix. The cell s.u.'s are taken into account individually in the estimation of s.u.'s in distances, angles and torsion angles; correlations between s.u.'s in cell parameters are only used when they are defined by crystal symmetry. An approximate (isotropic) treatment of cell s.u.'s is used for estimating s.u.'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 (Å²)

	x	y	z	U_{iso}*/U_{eq}
C1	0.7505 (4)	0.1344 (3)	0.4780 (3)	0.0364 (11)
C2	0.6846 (4)	0.1920 (3)	0.4197 (3)	0.0383 (11)
C3	0.6011 (5)	0.2346 (3)	0.4498 (3)	0.0499 (13)
H3	0.5556	0.273	0.4122	0.06*

C4	0.5835 (5)	0.2216 (4)	0.5341 (4)	0.0570 (15)
H4	0.5278	0.2518	0.5535	0.068*
C5	0.6487 (5)	0.1637 (3)	0.5897 (3)	0.0490 (13)
H5	0.6354	0.1544	0.6461	0.059*
C6	0.7338 (4)	0.1191 (3)	0.5631 (3)	0.0377 (11)
C7	0.6986 (4)	0.2074 (3)	0.3265 (3)	0.0377 (11)
C8	0.6370 (4)	0.1577 (3)	0.2572 (3)	0.0422 (12)
C9	0.6476 (4)	0.1740 (3)	0.1698 (3)	0.0476 (13)
C10	0.7155 (5)	0.2388 (3)	0.1525 (3)	0.0493 (13)
C11	0.7780 (4)	0.2865 (3)	0.2222 (3)	0.0473 (12)
C12	0.7713 (4)	0.2701 (3)	0.3103 (3)	0.0422 (12)
C13	0.5589 (6)	0.0903 (4)	0.2764 (4)	0.0680 (17)
H13A	0.4823	0.1009	0.2431	0.102*
H13B	0.5607	0.0898	0.3391	0.102*
H13C	0.5839	0.0374	0.2592	0.102*
C14	0.5810 (7)	0.1203 (5)	0.0945 (4)	0.084 (2)
H14A	0.6145	0.1244	0.0438	0.125*
H14B	0.5031	0.139	0.0782	0.125*
H14C	0.5831	0.0634	0.1139	0.125*
C15	0.7196 (6)	0.2594 (5)	0.0568 (4)	0.084 (2)
H15A	0.6584	0.2314	0.0163	0.127*
H15B	0.7914	0.2415	0.046	0.127*
H15C	0.7117	0.3184	0.0476	0.127*
C16	0.8538 (6)	0.3562 (4)	0.2035 (5)	0.079 (2)
H16A	0.813	0.3886	0.1538	0.118*
H16B	0.9211	0.3331	0.1895	0.118*
H16C	0.8753	0.3911	0.2553	0.118*
C17	0.8413 (5)	0.3199 (4)	0.3880 (4)	0.0681 (17)
H17A	0.8268	0.2997	0.443	0.102*
H17B	0.8204	0.3776	0.3807	0.102*
H17C	0.9211	0.3139	0.3893	0.102*
C18	0.8017 (4)	0.0550 (3)	0.6229 (3)	0.0368 (11)
C19	0.7593 (4)	−0.0266 (3)	0.6220 (3)	0.0396 (11)
C20	0.8242 (5)	−0.0875 (3)	0.6762 (3)	0.0468 (13)
C21	0.9318 (5)	−0.0657 (3)	0.7299 (3)	0.0489 (13)
C22	0.9726 (4)	0.0152 (3)	0.7312 (3)	0.0459 (12)
C23	0.9068 (4)	0.0763 (3)	0.6789 (3)	0.0447 (12)
C24	0.6463 (5)	−0.0464 (4)	0.5600 (4)	0.0608 (16)
H24A	0.6358	−0.1056	0.5565	0.091*
H24B	0.645	−0.0248	0.5015	0.091*
H24C	0.5859	−0.0214	0.5822	0.091*
C25	0.7790 (6)	−0.1748 (3)	0.6782 (4)	0.076 (2)
H25A	0.6995	−0.176	0.6484	0.114*
H25B	0.7879	−0.1924	0.7391	0.114*
H25C	0.8208	−0.2117	0.6483	0.114*
C26	1.0042 (6)	−0.1318 (4)	0.7865 (4)	0.078 (2)
H26A	1.0682	−0.1451	0.7616	0.117*
H26B	0.9591	−0.1809	0.7876	0.117*

H26C	1.0314	−0.1113	0.8463	0.117*
C27	1.0888 (5)	0.0364 (5)	0.7901 (4)	0.0730 (18)
H27A	1.1433	−0.0051	0.7828	0.109*
H27B	1.0837	0.038	0.8515	0.109*
H27C	1.1128	0.0899	0.7733	0.109*
C28	0.9492 (6)	0.1650 (4)	0.6799 (4)	0.0728 (19)
H28A	1.0236	0.1655	0.6671	0.109*
H28B	0.9536	0.189	0.7378	0.109*
H28C	0.8973	0.1969	0.6354	0.109*
I1	0.87935 (3)	0.06968 (3)	0.43456 (2)	0.05850 (16)
C29	0.2774 (6)	0.4708 (4)	0.3696 (4)	0.0684 (17)
H29	0.2996	0.4802	0.313	0.082*
Cl1	0.14519 (19)	0.41957 (14)	0.34585 (16)	0.1030 (7)
Cl2	0.3802 (2)	0.40834 (19)	0.43583 (18)	0.1351 (10)
Cl3	0.2678 (3)	0.56550 (15)	0.4179 (2)	0.1352 (10)

Atomic displacement parameters (Å²)

	U^{11}	U^{22}	U^{33}	U^{12}	U^{13}	U^{23}
C1	0.037 (3)	0.036 (3)	0.036 (2)	0.008 (2)	0.008 (2)	0.002 (2)
C2	0.046 (3)	0.038 (3)	0.030 (2)	0.002 (2)	0.006 (2)	0.000 (2)
C3	0.057 (3)	0.048 (3)	0.044 (3)	0.021 (3)	0.012 (3)	0.006 (2)
C4	0.065 (4)	0.056 (4)	0.055 (3)	0.024 (3)	0.024 (3)	−0.005 (3)
C5	0.059 (3)	0.056 (3)	0.035 (3)	0.008 (3)	0.016 (2)	0.002 (2)
C6	0.040 (3)	0.036 (3)	0.036 (3)	0.000 (2)	0.006 (2)	0.001 (2)
C7	0.037 (3)	0.041 (3)	0.033 (2)	0.014 (2)	0.005 (2)	0.007 (2)
C8	0.039 (3)	0.047 (3)	0.037 (3)	0.007 (2)	0.002 (2)	0.006 (2)
C9	0.047 (3)	0.053 (3)	0.039 (3)	0.011 (3)	0.004 (2)	0.000 (2)
C10	0.060 (3)	0.054 (3)	0.036 (3)	0.016 (3)	0.016 (3)	0.008 (2)
C11	0.044 (3)	0.049 (3)	0.049 (3)	0.006 (2)	0.012 (2)	0.015 (3)
C12	0.040 (3)	0.041 (3)	0.044 (3)	0.006 (2)	0.007 (2)	0.003 (2)
C13	0.072 (4)	0.064 (4)	0.065 (4)	−0.013 (3)	0.009 (3)	0.006 (3)
C14	0.104 (6)	0.096 (5)	0.044 (3)	−0.010 (4)	0.005 (4)	−0.015 (4)
C15	0.104 (6)	0.109 (6)	0.046 (4)	0.002 (5)	0.028 (4)	0.015 (4)
C16	0.084 (5)	0.080 (5)	0.077 (5)	−0.008 (4)	0.028 (4)	0.025 (4)
C17	0.067 (4)	0.072 (4)	0.061 (4)	−0.011 (3)	0.006 (3)	0.003 (3)
C18	0.039 (3)	0.040 (3)	0.031 (2)	0.004 (2)	0.008 (2)	0.007 (2)
C19	0.047 (3)	0.044 (3)	0.029 (2)	0.000 (2)	0.011 (2)	0.002 (2)
C20	0.067 (4)	0.038 (3)	0.039 (3)	0.003 (2)	0.018 (3)	0.002 (2)
C21	0.057 (3)	0.055 (3)	0.038 (3)	0.020 (3)	0.017 (2)	0.011 (2)
C22	0.042 (3)	0.062 (4)	0.035 (3)	0.004 (2)	0.010 (2)	0.005 (2)
C23	0.049 (3)	0.049 (3)	0.033 (2)	−0.003 (2)	0.004 (2)	0.004 (2)
C24	0.057 (4)	0.060 (4)	0.063 (4)	−0.012 (3)	0.010 (3)	−0.006 (3)
C25	0.119 (6)	0.037 (3)	0.070 (4)	−0.006 (3)	0.019 (4)	0.005 (3)
C26	0.079 (5)	0.078 (5)	0.073 (4)	0.032 (4)	0.008 (4)	0.028 (4)
C27	0.049 (4)	0.103 (5)	0.059 (4)	0.005 (3)	−0.003 (3)	0.011 (4)
C28	0.074 (4)	0.065 (4)	0.070 (4)	−0.023 (3)	−0.002 (3)	0.008 (3)
I1	0.0572 (2)	0.0742 (3)	0.0485 (2)	0.03317 (19)	0.02111 (17)	0.01532 (19)

C29	0.072 (4)	0.068 (4)	0.070 (4)	0.006 (3)	0.027 (3)	0.014 (3)
Cl1	0.0842 (14)	0.1114 (17)	0.1078 (16)	−0.0161 (12)	0.0108 (12)	0.0079 (13)
Cl2	0.0936 (17)	0.165 (3)	0.144 (2)	0.0425 (16)	0.0228 (16)	0.0602 (19)
Cl3	0.160 (3)	0.1012 (18)	0.150 (2)	−0.0052 (16)	0.048 (2)	−0.0496 (16)

Geometric parameters (Å, º)

C1—C6	1.392 (6)	C16—H16C	0.96
C1—C2	1.400 (6)	C17—H17A	0.96
C1—I1	2.099 (4)	C17—H17B	0.96
C2—C3	1.381 (6)	C17—H17C	0.96
C2—C7	1.501 (6)	C18—C23	1.398 (7)
C3—C4	1.377 (7)	C18—C19	1.406 (7)
C3—H3	0.93	C19—C20	1.400 (7)
C4—C5	1.379 (7)	C19—C24	1.503 (7)
C4—H4	0.93	C20—C21	1.408 (8)
C5—C6	1.386 (7)	C20—C25	1.507 (7)
C5—H5	0.93	C21—C22	1.389 (7)
C6—C18	1.492 (6)	C21—C26	1.514 (7)
C7—C12	1.394 (7)	C22—C23	1.393 (7)
C7—C8	1.398 (7)	C22—C27	1.518 (8)
C8—C9	1.403 (7)	C23—C28	1.513 (7)
C8—C13	1.507 (7)	C24—H24A	0.96
C9—C10	1.386 (7)	C24—H24B	0.96
C9—C14	1.516 (8)	C24—H24C	0.96
C10—C11	1.389 (7)	C25—H25A	0.96
C10—C15	1.521 (7)	C25—H25B	0.96
C11—C12	1.401 (7)	C25—H25C	0.96
C11—C16	1.512 (8)	C26—H26A	0.96
C12—C17	1.520 (7)	C26—H26B	0.96
C13—H13A	0.96	C26—H26C	0.96
C13—H13B	0.96	C27—H27A	0.96
C13—H13C	0.96	C27—H27B	0.96
C14—H14A	0.96	C27—H27C	0.96
C14—H14B	0.96	C28—H28A	0.96
C14—H14C	0.96	C28—H28B	0.96
C15—H15A	0.96	C28—H28C	0.96
C15—H15B	0.96	C29—Cl3	1.708 (7)
C15—H15C	0.96	C29—Cl2	1.729 (7)
C16—H16A	0.96	C29—Cl1	1.753 (7)
C16—H16B	0.96	C29—H29	0.98
C6—C1—C2	122.3 (4)	C12—C17—H17A	109.5
C6—C1—I1	119.4 (3)	C12—C17—H17B	109.5
C2—C1—I1	118.3 (3)	H17A—C17—H17B	109.5
C3—C2—C1	117.5 (4)	C12—C17—H17C	109.5
C3—C2—C7	119.3 (4)	H17A—C17—H17C	109.5
C1—C2—C7	123.2 (4)	H17B—C17—H17C	109.5

C4—C3—C2	121.5 (5)	C23—C18—C19	120.6 (4)
C4—C3—H3	119.3	C23—C18—C6	119.9 (4)
C2—C3—H3	119.3	C19—C18—C6	119.5 (4)
C3—C4—C5	119.8 (5)	C20—C19—C18	119.7 (5)
C3—C4—H4	120.1	C20—C19—C24	121.9 (5)
C5—C4—H4	120.1	C18—C19—C24	118.5 (4)
C4—C5—C6	121.2 (5)	C19—C20—C21	119.0 (5)
C4—C5—H5	119.4	C19—C20—C25	120.5 (5)
C6—C5—H5	119.4	C21—C20—C25	120.4 (5)
C5—C6—C1	117.7 (4)	C22—C21—C20	121.0 (5)
C5—C6—C18	120.8 (4)	C22—C21—C26	119.6 (5)
C1—C6—C18	121.5 (4)	C20—C21—C26	119.4 (5)
C12—C7—C8	121.3 (4)	C21—C22—C23	120.0 (5)
C12—C7—C2	120.2 (4)	C21—C22—C27	119.5 (5)
C8—C7—C2	118.5 (4)	C23—C22—C27	120.5 (5)
C7—C8—C9	118.3 (5)	C22—C23—C18	119.7 (5)
C7—C8—C13	120.3 (5)	C22—C23—C28	120.9 (5)
C9—C8—C13	121.3 (5)	C18—C23—C28	119.5 (5)
C10—C9—C8	120.8 (5)	C19—C24—H24A	109.5
C10—C9—C14	120.8 (5)	C19—C24—H24B	109.5
C8—C9—C14	118.4 (5)	H24A—C24—H24B	109.5
C9—C10—C11	120.3 (5)	C19—C24—H24C	109.5
C9—C10—C15	120.2 (5)	H24A—C24—H24C	109.5
C11—C10—C15	119.6 (5)	H24B—C24—H24C	109.5
C10—C11—C12	120.0 (5)	C20—C25—H25A	109.5
C10—C11—C16	120.3 (5)	C20—C25—H25B	109.5
C12—C11—C16	119.7 (5)	H25A—C25—H25B	109.5
C7—C12—C11	119.2 (5)	C20—C25—H25C	109.5
C7—C12—C17	119.8 (5)	H25A—C25—H25C	109.5
C11—C12—C17	121.0 (5)	H25B—C25—H25C	109.5
C8—C13—H13A	109.5	C21—C26—H26A	109.5
C8—C13—H13B	109.5	C21—C26—H26B	109.5
H13A—C13—H13B	109.5	H26A—C26—H26B	109.5
C8—C13—H13C	109.5	C21—C26—H26C	109.5
H13A—C13—H13C	109.5	H26A—C26—H26C	109.5
H13B—C13—H13C	109.5	H26B—C26—H26C	109.5
C9—C14—H14A	109.5	C22—C27—H27A	109.5
C9—C14—H14B	109.5	C22—C27—H27B	109.5
H14A—C14—H14B	109.5	H27A—C27—H27B	109.5
C9—C14—H14C	109.5	C22—C27—H27C	109.5
H14A—C14—H14C	109.5	H27A—C27—H27C	109.5
H14B—C14—H14C	109.5	H27B—C27—H27C	109.5
C10—C15—H15A	109.5	C23—C28—H28A	109.5
C10—C15—H15B	109.5	C23—C28—H28B	109.5
H15A—C15—H15B	109.5	H28A—C28—H28B	109.5
C10—C15—H15C	109.5	C23—C28—H28C	109.5
H15A—C15—H15C	109.5	H28A—C28—H28C	109.5
H15B—C15—H15C	109.5	H28B—C28—H28C	109.5

C11—C16—H16A	109.5	Cl3—C29—Cl2	111.9 (4)
C11—C16—H16B	109.5	Cl3—C29—Cl1	111.1 (4)
H16A—C16—H16B	109.5	Cl2—C29—Cl1	110.0 (4)
C11—C16—H16C	109.5	Cl3—C29—H29	107.9
H16A—C16—H16C	109.5	Cl2—C29—H29	107.9
H16B—C16—H16C	109.5	Cl1—C29—H29	107.9

C6—C1—C2—C3	0.3 (7)	C2—C7—C12—C11	−176.4 (4)
I1—C1—C2—C3	−179.4 (4)	C8—C7—C12—C17	−176.9 (5)
C6—C1—C2—C7	−177.9 (4)	C2—C7—C12—C17	3.3 (7)
I1—C1—C2—C7	2.5 (6)	C10—C11—C12—C7	−2.0 (7)
C1—C2—C3—C4	0.5 (8)	C16—C11—C12—C7	178.1 (5)
C7—C2—C3—C4	178.8 (5)	C10—C11—C12—C17	178.3 (5)
C2—C3—C4—C5	−1.3 (9)	C16—C11—C12—C17	−1.5 (8)
C3—C4—C5—C6	1.3 (9)	C5—C6—C18—C23	−94.7 (6)
C4—C5—C6—C1	−0.5 (8)	C1—C6—C18—C23	87.5 (6)
C4—C5—C6—C18	−178.3 (5)	C5—C6—C18—C19	85.6 (6)
C2—C1—C6—C5	−0.3 (7)	C1—C6—C18—C19	−92.2 (6)
I1—C1—C6—C5	179.3 (4)	C23—C18—C19—C20	−1.4 (7)
C2—C1—C6—C18	177.5 (4)	C6—C18—C19—C20	178.3 (4)
I1—C1—C6—C18	−2.8 (6)	C23—C18—C19—C24	−179.5 (5)
C3—C2—C7—C12	89.3 (6)	C6—C18—C19—C24	0.2 (7)
C1—C2—C7—C12	−92.5 (6)	C18—C19—C20—C21	−0.8 (7)
C3—C2—C7—C8	−90.5 (6)	C24—C19—C20—C21	177.2 (5)
C1—C2—C7—C8	87.7 (6)	C18—C19—C20—C25	178.3 (5)
C12—C7—C8—C9	−1.7 (7)	C24—C19—C20—C25	−3.7 (8)
C2—C7—C8—C9	178.1 (4)	C19—C20—C21—C22	1.5 (7)
C12—C7—C8—C13	−179.9 (5)	C25—C20—C21—C22	−177.6 (5)
C2—C7—C8—C13	−0.2 (7)	C19—C20—C21—C26	−178.1 (5)
C7—C8—C9—C10	−1.4 (7)	C25—C20—C21—C26	2.8 (8)
C13—C8—C9—C10	176.8 (5)	C20—C21—C22—C23	0.1 (7)
C7—C8—C9—C14	−180.0 (5)	C26—C21—C22—C23	179.7 (5)
C13—C8—C9—C14	−1.8 (8)	C20—C21—C22—C27	179.8 (5)
C8—C9—C10—C11	2.7 (8)	C26—C21—C22—C27	−0.6 (8)
C14—C9—C10—C11	−178.7 (5)	C21—C22—C23—C18	−2.3 (7)
C8—C9—C10—C15	−175.7 (5)	C27—C22—C23—C18	178.0 (5)
C14—C9—C10—C15	2.8 (8)	C21—C22—C23—C28	179.5 (5)
C9—C10—C11—C12	−1.0 (8)	C27—C22—C23—C28	−0.2 (8)
C15—C10—C11—C12	177.5 (5)	C19—C18—C23—C22	3.0 (7)
C9—C10—C11—C16	178.8 (5)	C6—C18—C23—C22	−176.7 (4)
C15—C10—C11—C16	−2.7 (8)	C19—C18—C23—C28	−178.7 (5)
C8—C7—C12—C11	3.4 (7)	C6—C18—C23—C28	1.5 (7)

Hydrogen-bond geometry (Å, º)

Cg2 and C3 are the centroids of the C7–C12 and C18–C23 benzene rings, respectively.

D—H···*A*	*D*—H	H···*A*	*D*···*A*	*D*—H···*A*
C26—H26*A*···*Cg2*^{i}	0.96	3.86 (1)	2.97	155

C28—H28*A*···*Cg*2^{ii}	0.96	3.53 (1)	2.87	127
C29—H29···*Cg*3^{iii}	0.98	3.42 (1)	2.44	177

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