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Racemic tricarbonyl[7-methoxy-2-(η^6 -phenyl)chromane]chromium(0)

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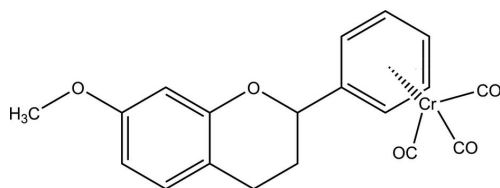
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Key indicators: single-crystal X-ray study; $T = 100$ K; mean $\sigma(\text{C}-\text{C}) = 0.002$ Å; R factor = 0.034; wR factor = 0.090; data-to-parameter ratio = 18.3.

In the title compound, $[\text{Cr}(\text{C}_{16}\text{H}_{16}\text{O}_2)(\text{CO})_3]$, the Cr^0 atom of the $\text{Cr}(\text{CO})_3$ unit is coordinated to the phenyl ring of the flavan ligand in an η^6 mode, with a normal arene-to-metal distance. The $\text{Cr}(\text{CO})_3$ unit exhibits a three-legged piano-stool conformation, while the dihydropyran ring displays a distorted envelope configuration. The phenyl ring is twisted away from the fused ring system by 25.5 (2)°. The methoxy group is almost coplanar with the phenyl ring [$\text{C}_{\text{Me}}-\text{O}-\text{C}_{\text{ar}}-\text{C}_{\text{ar}}$ torsion angle = 8.46 (2)°]. The crystal packing is stabilized by intermolecular $\text{C}-\text{H}\cdots\text{O}$ interactions.

Related literature

For similar structures, see: van Tonder *et al.* (2010*a,b*) and for other related structures, see: van Tonder *et al.* (2009*a,b*). For the synthesis of the title compound, see: Müller *et al.* (1999) and for the synthesis of 7-methoxyflavan-4-one, see: Sato *et al.* (2006). For standard bond lengths, see: Allen *et al.* (1987). For the importance of flavonoids in biological investigations, see: Rice-Evans & Packer (2003). For the use of tricarbonyl-(arene)chromium complexes in regioselective organic synthesis, see: Muschalek *et al.* (2007).



Experimental

Crystal data

$[\text{Cr}(\text{C}_{16}\text{H}_{16}\text{O}_2)(\text{CO})_3]$

$M_r = 376.32$

Monoclinic, $P2_1/c$

$a = 9.7703$ (5) Å

$b = 19.1820$ (9) Å

$c = 8.8049$ (4) Å

$\beta = 97.494$ (2)°

$V = 1636.07$ (14) Å³

$Z = 4$
Mo $K\alpha$ radiation
 $\mu = 0.73$ mm⁻¹

$T = 100$ K
 $0.34 \times 0.23 \times 0.08$ mm

Data collection

Bruker X8 APEXII 4K KappaCCD diffractometer
Absorption correction: multi-scan (SADABS; Bruker, 2008)
 $T_{\text{min}} = 0.818$, $T_{\text{max}} = 0.942$

29057 measured reflections
4069 independent reflections
3526 reflections with $I > 2\sigma(I)$
 $R_{\text{int}} = 0.033$

Refinement

$R[F^2 > 2\sigma(F^2)] = 0.034$
 $wR(F^2) = 0.090$
 $S = 1.00$
4069 reflections

222 parameters
H-atom parameters constrained
 $\Delta\rho_{\text{max}} = 0.73$ e Å⁻³
 $\Delta\rho_{\text{min}} = -0.52$ e Å⁻³

Table 1

Hydrogen-bond geometry (Å, °).

$D-H\cdots A$	$D-H$	$H\cdots A$	$D\cdots A$	$D-H\cdots A$
$\text{C4}'-\text{H4}'\cdots\text{O2}^{\text{i}}$	0.93	2.54	3.459 (2)	169
$\text{C2}'-\text{H2}'\cdots\text{O4}^{\text{ii}}$	0.93	2.46	3.153 (2)	132
$\text{C1}-\text{H1C}\cdots\text{O4}^{\text{iii}}$	0.96	2.57	3.314 (2)	134

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

Data collection: APEX2 (Bruker, 2008); cell refinement: SAINT-Plus (Bruker, 2008); 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 & Putz, 2005); software used to prepare material for publication: WinGX (Farrugia, 1999).

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Supplementary data and figures for this paper are available from the IUCr electronic archives (Reference: HP2002).

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Tonder, J. H. van, Bezuidenhout, B. C. B. & Janse van Rensburg, J. M.
(2010*b*). *Acta Cryst.* **E66**, m1086.

supporting information

Acta Cryst. (2011). E67, m438–m439 [doi:10.1107/S1600536811008683]

Racemic tricarbonyl[7-methoxy-2-(η^6 -phenyl)chromane]chromium(0)**Mukut Gohain, Theunis J. Muller and Barend C. B. Bezuidenhout****S1. Comment**

The title compound, $[\text{Cr}(\text{CO})_3(\text{C}_{16}\text{H}_{16}\text{O}_2)]$ forms part of a series of chromium(0) complexes of the type $[\text{Cr}(\text{flav})(\text{CO})_3]$ (flav = flavan, flavone or isoflavone ligand) (van Tonder *et al.*, 2009*a,b* and 2010*a,b*). Our interest in this type of chromium(0) complexes is partly due to regioselective organic synthesis (Muschalek *et al.*, 2007) and to the general biological activity of flavanoids (Rice-Evans & Packer, 2003).

The title compound crystallized in the monoclinic space group $P2(1)/c$, with $Z = 4$. For the title compound the molecular structure displays the $\text{Cr}(\text{CO})_3$ moiety complexed to the flavone C-ring, exhibiting the known three-legged piano-stool conformation. All bond distances and angles are within range (Allen *et al.*, 1987). The distance between the Cr^0 atom and the centroid of the A - η^6 -coordinated phenyl ring is 1.7119 (3) Å. The plane through C1'-C6' (r.m.s = 0.0038 fitted atoms C1'-C6') of the phenyl ring is at an angle of 42.762 (43) ° to the plane formed between C4—C10 and O1 (r.m.s = 0.00824 fitted atoms C4, C5, C6, C7, C8, C9, C10 and O1). The dihydropyran ring displays a distorted envelope configuration by displacement of atoms C2 and C3 from the fused ring system plane, with distances of 0.105 (2) 0.618 (2) Å respectively. The methoxy group at the C7 position is nearly coplanar to the plane created by the fused ring system. The methoxy forms a dihedral angle of -8.464 (23) ° between C1—O2—C7—C6. The molecules form chains through intermolecular C4—H4 \cdots O2ⁱ, C2—H2 \cdots O4ⁱⁱ and C1—H1C \cdots O4ⁱⁱⁱ hydrogen interactions (Table 1).

S2. Experimental

7-methoxyflavan-4-one was synthesized as described by Sato *et al.* (2006). 7-Methoxyflavan-4-one (1.00 g; 3.9 mmol), 10% Pd/C (0.10 g), 3 M H_2SO_4 (aq) (1 ml), ethanol (30 ml). Purification by means of flash column-chromatography yielded 7-methoxyflavan (0.67 g; 70.6%) as a colourless oil as described by Sato *et al.* (2006) R_f 0.65 (hexane:DCM:ethyl acetate; 50:50:1); ^1H NMR (600 MHz, CDCl_3) δ 7.44–7.41 (m, H-2' and H-6', 2H), 7.40–7.37 (m, H-3' and H-5', 2H), 7.34–7.31 (m, H-4', 1H), 6.99–6.97 (m, H-5, 1H), 6.50–6.47 (m, H-6 and H-8, 2H), 5.05 (dd, $J = 10.19, 2.37$ Hz, H-2 1H), 3.77 (s, -OCH₃, 3H), 2.92 (ddd, $J = 16.08, 10.92, 6.02$ Hz, H-4(*a*) 1H), 2.74 (ddd, $J = 16.08, 5.12, 3.40$ Hz, H-4(*e*) 1H), 2.22–2.18 (m, H-3 1H), 2.11–2.04 (m, H-3 1H); ^{13}C NMR (151 MHz, CDCl_3) δ p.p.m. 24.47 (C-4), 30.19 (C-3), 55.38 (-OCH₃), 77.98 (C-2), 101.71 (C-6/8), 107.54 (C-6/8), 114.01, 126.11, 127.93, 128.61, 130.05, 141.79, 155.91, 159.23.

Preparation of the title compound was based on a method described by Müller *et al.* (1999). A solution of 7-methoxyflavan (0.27 g, 1.1 mmol) and $\text{Cr}(\text{CO})_6$ (0.25 g, 1.1 mmol, 1 eq) in butylether:THF (9:1; 25 ml) was degassed with argon, using standard Schlenk techniques, and refluxed (70 hr) under an oxygen free atmosphere. The reaction mixture was cooled to room temperature and evaporated *in vacuo*. Purification through flash column-chromatography yielded the title compound (0.10 g; 24%) as a yellow solid. Recrystallization from hexane:dichloromethane (6:1) yielded yellow plate like crystals suitable for X-ray analysis.

R_f 0.30 (hexane:ethyl acetate; 3:5); Mp 145–147 °C IR $\nu(\text{CO}) = 1952, 1894$ and 1844 cm^{-1} ; ^1H NMR (600 MHz, CDCl_3) δ 6.96 (d, $J = 8.3$ Hz, 1H), 6.49 (dd, $J = 8.3, 2.5$ Hz, 1H), 6.48 (d, $J = 2.5$ Hz, 1H), 5.56 (d, $J = 6.2$ Hz, 1H), 5.43–5.41 (m, 1H), 5.37–5.32 (m, 3H), 4.67 (dd, $J = 10.4, 2.3$ Hz, 1H), 3.77 (s, 3H), 2.94–2.87 (m, 1H), 2.76 (m, 1H), 2.25 (m, 1H), 1.99 (m, 1H). ^{13}C NMR (151 MHz, CDCl_3) δ p.p.m. 232.53, 159.27, 155.02, 129.85, 113.31, 111.38, 108.09, 101.51, 92.22, 91.62, 91.49, 91.22, 90.68, 75.37, 55.37, 29.89, 24.11.

S3. Refinement

All H atoms were positioned geometrically and allowed to ride on their parent atoms, with $U_{\text{iso}}(\text{H}) = 1.2U_{\text{eq}}(\text{parent})$ of the parent atom with a C—H distance of 0.93. The methine H atoms were placed in geometrically idealized positions and constrained to ride on its parent atoms with $U_{\text{iso}}(\text{H}) = 1.5U_{\text{eq}}(\text{C})$ and at a distance of 0.97 Å. The methyl H atoms were placed in geometrically idealized positions and constrained to ride on its parent atoms with $U_{\text{iso}}(\text{H}) = 1.5U_{\text{eq}}(\text{C})$ and at a distance of 0.96 Å.

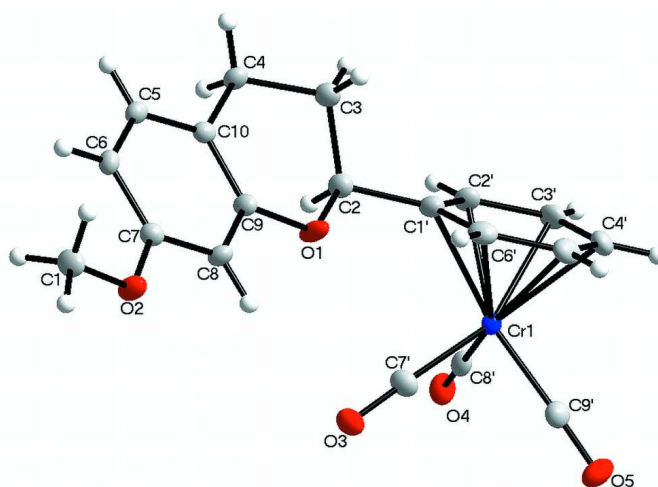


Figure 1

Diamond representation of the title compound, showing the numbering scheme and displacement ellipsoids (50% probability).

tricarbonyl[7-methoxy-2-(η^6 -phenyl)chromane]chromium(0)

Crystal data

$[\text{Cr}(\text{C}_{16}\text{H}_{16}\text{O}_2)(\text{CO})_3]$

$M_r = 376.32$

Monoclinic, $P2_1/c$

$a = 9.7703$ (5) Å

$b = 19.1820$ (9) Å

$c = 8.8049$ (4) Å

$\beta = 97.494$ (2)°

$V = 1636.07$ (14) Å³

$Z = 4$

$F(000) = 776$

$D_x = 1.528 \text{ Mg m}^{-3}$

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

Cell parameters from 9920 reflections

$\theta = 3.0$ – 28.2 °

$\mu = 0.73 \text{ mm}^{-1}$

$T = 100$ K

Plate, yellow

$0.34 \times 0.23 \times 0.08$ mm

Data collection

Bruker X8 APEXII 4K KappaCCD
diffractometer
Graphite monochromator
Detector resolution: 8.5 pixels mm⁻¹
 φ and ω scans
Absorption correction: multi-scan
(SADABS; Bruker, 2008)
 $T_{\min} = 0.818$, $T_{\max} = 0.942$

29057 measured reflections
4069 independent reflections
3526 reflections with $I > 2\sigma(I)$
 $R_{\text{int}} = 0.033$
 $\theta_{\max} = 28.3^\circ$, $\theta_{\min} = 3.1^\circ$
 $h = -13 \rightarrow 9$
 $k = -25 \rightarrow 25$
 $l = -11 \rightarrow 11$

Refinement

Refinement on F^2
Least-squares matrix: full
 $R[F^2 > 2\sigma(F^2)] = 0.034$
 $wR(F^2) = 0.090$
 $S = 1.00$
4069 reflections
222 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
 $w = 1/[\sigma^2(F_o^2) + (0.0434P)^2 + 1.6967P]$
where $P = (F_o^2 + 2F_c^2)/3$
 $(\Delta/\sigma)_{\max} = 0.015$
 $\Delta\rho_{\max} = 0.73 \text{ e } \text{\AA}^{-3}$
 $\Delta\rho_{\min} = -0.52 \text{ e } \text{\AA}^{-3}$

Special details

Experimental. The intensity data was collected on a Bruker X8 ApexII 4 K Kappa CCD diffractometer using an exposure time of 15 s/frame. A total of 1821 frames were collected with a frame width of 0.5° covering up to $\theta = 28.18^\circ$ with 99.7% completeness accomplished.

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 (\AA^2)

	x	y	z	$U_{\text{iso}}^*/U_{\text{eq}}$
C1	0.3380 (2)	0.29331 (9)	0.6478 (2)	0.0208 (4)
H1A	0.291	0.2854	0.7355	0.031*
H1B	0.3066	0.3363	0.5995	0.031*
H1C	0.4356	0.2959	0.6798	0.031*
C1'	0.17467 (19)	-0.11113 (9)	0.36846 (19)	0.0178 (3)
C2	0.28858 (19)	-0.07158 (9)	0.4658 (2)	0.0190 (3)
H2	0.3777	-0.0837	0.4332	0.023*
C2'	0.19981 (19)	-0.17979 (9)	0.3254 (2)	0.0196 (3)
H2'	0.2859	-0.1997	0.355	0.023*
C3'	0.0952 (2)	-0.21918 (9)	0.2369 (2)	0.0213 (4)
H3'	0.1123	-0.265	0.2106	0.026*
C3	0.29216 (19)	-0.08786 (9)	0.6350 (2)	0.0202 (4)
H3A	0.3106	-0.1371	0.6528	0.024*
H3B	0.2034	-0.0771	0.6676	0.024*

C4'	-0.03356 (19)	-0.18914 (10)	0.1893 (2)	0.0219 (4)
H4'	-0.1018	-0.2147	0.1301	0.026*
C4	0.40480 (18)	-0.04451 (8)	0.72666 (19)	0.0167 (3)
H4A	0.4003	-0.0501	0.8354	0.02*
H4B	0.4948	-0.0602	0.7056	0.02*
C5	0.43124 (17)	0.08604 (9)	0.77979 (19)	0.0159 (3)
H5	0.4782	0.0757	0.8759	0.019*
C5'	-0.05972 (19)	-0.11962 (10)	0.2316 (2)	0.0220 (4)
H5'	-0.1454	-0.0995	0.2003	0.026*
C6	0.41013 (17)	0.15570 (9)	0.73907 (19)	0.0159 (3)
H6	0.4428	0.1912	0.8062	0.019*
C6'	0.04282 (19)	-0.08107 (9)	0.3203 (2)	0.0197 (3)
H6'	0.0248	-0.0355	0.348	0.024*
C7	0.33885 (17)	0.17112 (9)	0.59536 (19)	0.0159 (3)
C7'	0.20844 (19)	-0.03456 (9)	0.0866 (2)	0.0210 (2)
C8	0.29047 (18)	0.11775 (9)	0.49617 (19)	0.0176 (3)
H8	0.2422	0.1281	0.4007	0.021*
C8'	0.26596 (17)	-0.16022 (8)	0.02184 (19)	0.0168 (3)
C9	0.31439 (18)	0.04906 (9)	0.53999 (19)	0.0163 (3)
C9'	0.02989 (18)	-0.11148 (9)	-0.0765 (2)	0.0185 (3)
C10	0.38501 (17)	0.03117 (8)	0.68272 (19)	0.0148 (3)
O1	0.25988 (15)	0.00078 (6)	0.43410 (14)	0.0232 (3)
O2	0.30939 (14)	0.23734 (6)	0.54184 (14)	0.0210 (3)
O3	0.26044 (13)	0.01758 (6)	0.06402 (15)	0.0210 (2)
O4	0.35317 (13)	-0.18513 (7)	-0.03763 (15)	0.0218 (3)
O5	-0.02741 (14)	-0.10672 (8)	-0.19953 (15)	0.0280 (3)
Cr1	0.12859 (3)	-0.120499 (13)	0.11540 (3)	0.01295 (8)

Atomic displacement parameters (Å²)

	U^{11}	U^{22}	U^{33}	U^{12}	U^{13}	U^{23}
C1	0.0271 (9)	0.0133 (8)	0.0223 (9)	-0.0016 (6)	0.0046 (7)	-0.0035 (6)
C1'	0.0238 (9)	0.0148 (8)	0.0143 (8)	-0.0025 (6)	0.0010 (6)	0.0018 (6)
C2	0.0226 (8)	0.0155 (8)	0.0185 (8)	-0.0028 (6)	0.0008 (7)	0.0002 (6)
C2'	0.0249 (9)	0.0141 (8)	0.0184 (8)	-0.0011 (6)	-0.0022 (7)	0.0037 (6)
C3'	0.0295 (10)	0.0125 (7)	0.0214 (9)	-0.0055 (7)	0.0017 (7)	0.0026 (6)
C3	0.0240 (9)	0.0170 (8)	0.0189 (8)	-0.0025 (7)	0.0007 (7)	0.0026 (6)
C4'	0.0204 (9)	0.0238 (9)	0.0215 (8)	-0.0092 (7)	0.0030 (7)	0.0029 (7)
C4	0.0190 (8)	0.0152 (8)	0.0149 (7)	0.0005 (6)	-0.0017 (6)	0.0014 (6)
C5	0.0134 (7)	0.0194 (8)	0.0145 (7)	-0.0003 (6)	-0.0003 (6)	0.0010 (6)
C5'	0.0157 (8)	0.0282 (9)	0.0231 (9)	-0.0005 (7)	0.0062 (7)	0.0040 (7)
C6	0.0146 (8)	0.0168 (8)	0.0160 (8)	-0.0024 (6)	0.0013 (6)	-0.0034 (6)
C6'	0.0243 (9)	0.0188 (8)	0.0170 (8)	0.0011 (7)	0.0072 (7)	-0.0003 (6)
C7	0.0168 (8)	0.0143 (7)	0.0169 (8)	0.0001 (6)	0.0029 (6)	0.0005 (6)
C7'	0.0240 (5)	0.0159 (4)	0.0236 (5)	-0.0015 (4)	0.0056 (4)	0.0004 (4)
C8	0.0218 (8)	0.0169 (8)	0.0130 (7)	-0.0012 (6)	-0.0016 (6)	0.0016 (6)
C8'	0.0179 (8)	0.0128 (7)	0.0184 (8)	-0.0028 (6)	-0.0019 (6)	-0.0013 (6)
C9	0.0215 (8)	0.0140 (7)	0.0134 (7)	-0.0029 (6)	0.0019 (6)	-0.0017 (6)

C9'	0.0143 (8)	0.0202 (8)	0.0213 (8)	0.0005 (6)	0.0036 (6)	-0.0005 (6)
C10	0.0134 (7)	0.0159 (7)	0.0151 (7)	0.0004 (6)	0.0025 (6)	0.0016 (6)
O1	0.0419 (8)	0.0117 (6)	0.0135 (6)	-0.0046 (5)	-0.0057 (5)	0.0009 (4)
O2	0.0314 (7)	0.0122 (6)	0.0177 (6)	0.0000 (5)	-0.0027 (5)	0.0002 (5)
O3	0.0240 (5)	0.0159 (4)	0.0236 (5)	-0.0015 (4)	0.0056 (4)	0.0004 (4)
O4	0.0190 (6)	0.0206 (6)	0.0258 (7)	0.0003 (5)	0.0032 (5)	-0.0063 (5)
O5	0.0236 (7)	0.0394 (8)	0.0195 (7)	0.0032 (6)	-0.0032 (5)	0.0001 (6)
Cr1	0.01308 (14)	0.01116 (13)	0.01423 (14)	-0.00064 (9)	0.00033 (9)	-0.00028 (9)

Geometric parameters (Å, °)

C1—O2	1.426 (2)	C4—H4A	0.97
C1—H1A	0.96	C4—H4B	0.97
C1—H1B	0.96	C5—C6	1.392 (2)
C1—H1C	0.96	C5—C10	1.393 (2)
C1'—C2'	1.401 (2)	C5—H5	0.93
C1'—C6'	1.424 (3)	C5'—C6'	1.399 (3)
C1'—C2	1.517 (2)	C5'—Cr1	2.2187 (18)
C1'—Cr1	2.2221 (17)	C5'—H5'	0.93
C2—O1	1.436 (2)	C6—C7	1.394 (2)
C2—C3	1.518 (2)	C6—H6	0.93
C2—H2	0.98	C6'—Cr1	2.2188 (17)
C2'—C3'	1.419 (2)	C6'—H6'	0.93
C2'—Cr1	2.2044 (17)	C7—O2	1.372 (2)
C2'—H2'	0.93	C7—C8	1.388 (2)
C3'—C4'	1.397 (3)	C7'—O3	1.151 (2)
C3'—Cr1	2.2194 (17)	C7'—Cr1	1.8552 (18)
C3'—H3'	0.93	C8—C9	1.384 (2)
C3—C4	1.524 (2)	C8—H8	0.93
C3—H3A	0.97	C8'—O4	1.160 (2)
C3—H3B	0.97	C8'—Cr1	1.8300 (18)
C4'—C5'	1.417 (3)	C9—O1	1.3716 (19)
C4'—Cr1	2.2216 (17)	C9—C10	1.395 (2)
C4'—H4'	0.93	C9'—O5	1.156 (2)
C4—C10	1.508 (2)	C9'—Cr1	1.8413 (18)
O2—C1—H1A	109.5	C5'—C6'—C1'	120.42 (16)
O2—C1—H1B	109.5	C5'—C6'—Cr1	71.62 (10)
H1A—C1—H1B	109.5	C1'—C6'—Cr1	71.42 (10)
O2—C1—H1C	109.5	C5'—C6'—H6'	119.8
H1A—C1—H1C	109.5	C1'—C6'—H6'	119.8
H1B—C1—H1C	109.5	Cr1—C6'—H6'	129.6
C2'—C1'—C6'	118.91 (16)	O2—C7—C8	115.32 (14)
C2'—C1'—C2	118.79 (16)	O2—C7—C6	124.44 (15)
C6'—C1'—C2	122.29 (15)	C8—C7—C6	120.24 (15)
C2'—C1'—Cr1	70.86 (10)	O3—C7'—Cr1	177.22 (16)
C6'—C1'—Cr1	71.16 (10)	C9—C8—C7	119.69 (15)
C2—C1'—Cr1	130.16 (12)	C9—C8—H8	120.2

O1—C2—C1'	105.27 (14)	C7—C8—H8	120.2
O1—C2—C3	111.62 (14)	O4—C8'—Cr1	179.72 (16)
C1'—C2—C3	111.90 (14)	O1—C9—C8	114.63 (14)
O1—C2—H2	109.3	O1—C9—C10	123.27 (15)
C1'—C2—H2	109.3	C8—C9—C10	122.07 (15)
C3—C2—H2	109.3	O5—C9'—Cr1	177.22 (16)
C1'—C2'—C3'	120.66 (17)	C5—C10—C9	116.69 (15)
C1'—C2'—Cr1	72.24 (10)	C5—C10—C4	123.31 (15)
C3'—C2'—Cr1	71.86 (10)	C9—C10—C4	119.97 (15)
C1'—C2'—H2'	119.7	C9—O1—C2	118.15 (13)
C3'—C2'—H2'	119.7	C7—O2—C1	117.25 (13)
Cr1—C2'—H2'	128.5	C8'—Cr1—C9'	87.31 (8)
C4'—C3'—C2'	120.04 (17)	C8'—Cr1—C7'	87.84 (8)
C4'—C3'—Cr1	71.75 (10)	C9'—Cr1—C7'	88.31 (8)
C2'—C3'—Cr1	70.71 (10)	C8'—Cr1—C2'	89.59 (7)
C4'—C3'—H3'	120	C9'—Cr1—C2'	152.67 (7)
C2'—C3'—H3'	120	C7'—Cr1—C2'	118.72 (7)
Cr1—C3'—H3'	130.1	C8'—Cr1—C5'	155.74 (7)
C2—C3—C4	109.27 (14)	C9'—Cr1—C5'	93.20 (7)
C2—C3—H3A	109.8	C7'—Cr1—C5'	116.42 (8)
C4—C3—H3A	109.8	C2'—Cr1—C5'	78.89 (7)
C2—C3—H3B	109.8	C8'—Cr1—C6'	152.24 (7)
C4—C3—H3B	109.8	C9'—Cr1—C6'	120.40 (7)
H3A—C3—H3B	108.3	C7'—Cr1—C6'	91.05 (7)
C3'—C4'—C5'	119.71 (16)	C2'—Cr1—C6'	66.76 (7)
C3'—C4'—Cr1	71.57 (10)	C5'—Cr1—C6'	36.75 (7)
C5'—C4'—Cr1	71.28 (10)	C8'—Cr1—C3'	91.40 (7)
C3'—C4'—H4'	120.1	C9'—Cr1—C3'	115.48 (7)
C5'—C4'—H4'	120.1	C7'—Cr1—C3'	156.15 (8)
Cr1—C4'—H4'	129.4	C2'—Cr1—C3'	37.43 (6)
C10—C4—C3	109.26 (13)	C5'—Cr1—C3'	66.50 (7)
C10—C4—H4A	109.8	C6'—Cr1—C3'	78.69 (7)
C3—C4—H4A	109.8	C8'—Cr1—C1'	114.87 (7)
C10—C4—H4B	109.8	C9'—Cr1—C1'	157.81 (7)
C3—C4—H4B	109.8	C7'—Cr1—C1'	91.79 (7)
H4A—C4—H4B	108.3	C2'—Cr1—C1'	36.90 (6)
C6—C5—C10	122.81 (15)	C5'—Cr1—C1'	66.97 (7)
C6—C5—H5	118.6	C6'—Cr1—C1'	37.42 (7)
C10—C5—H5	118.6	C3'—Cr1—C1'	66.98 (6)
C6'—C5'—C4'	120.25 (17)	C8'—Cr1—C4'	118.55 (7)
C6'—C5'—Cr1	71.63 (10)	C9'—Cr1—C4'	90.93 (7)
C4'—C5'—Cr1	71.51 (10)	C7'—Cr1—C4'	153.54 (8)
C6'—C5'—H5'	119.9	C2'—Cr1—C4'	66.91 (7)
C4'—C5'—H5'	119.9	C5'—Cr1—C4'	37.21 (7)
Cr1—C5'—H5'	129.4	C6'—Cr1—C4'	66.71 (7)
C5—C6—C7	118.50 (15)	C3'—Cr1—C4'	36.68 (7)
C5—C6—H6	120.8	C1'—Cr1—C4'	79.24 (7)
C7—C6—H6	120.8		

Hydrogen-bond geometry (Å, °)

<i>D</i> —H \cdots <i>A</i>	<i>D</i> —H	H \cdots <i>A</i>	<i>D</i> \cdots <i>A</i>	<i>D</i> —H \cdots <i>A</i>
C4'—H4' \cdots O2 ⁱ	0.93	2.54	3.459 (2)	169
C2'—H2' \cdots O4 ⁱⁱ	0.93	2.46	3.153 (2)	132
C1—H1C \cdots O4 ⁱⁱⁱ	0.96	2.57	3.314 (2)	134

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