

{2-Hydroxy-6-[(2-oxidophenyl)imino-methyl- κ^2N,O]phenolato- κO^1]phenyl-boron

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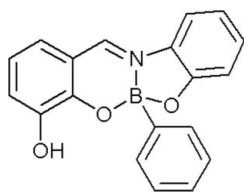
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 Key indicators: single-crystal X-ray study; $T = 293$ K; mean $\sigma(C-C) = 0.004$ Å; R factor = 0.061; wR factor = 0.127; data-to-parameter ratio = 15.1.

The [4.3.0]heterobicyclic title structure, $C_{19}H_{14}BNO_3$, is composed of a five-membered OBNC₂ ring and a six-membered OBNC₃ ring, each of which has an approximate envelope conformation. The coordination geometry of the B atom is distorted tetrahedral. In the crystal structure, centrosymmetrically related molecules are associated through pairs of O—H...O hydrogen bonds.

Related literature

For related boronates, see: Barba *et al.* (2001); Höpfl *et al.* (1998); Lamère *et al.* (2006). For boronates with non-linear optical properties, see: Reyes *et al.* (2005); Muñoz *et al.* (2008). For the use of boronates in organic synthesis, see: Rodríguez *et al.* (2005a,b); López-Ruiz *et al.* (2008). For a definition of the tetrahedral character in boron compounds, see: Höpfl *et al.* (1999).



Experimental

Crystal data

$C_{19}H_{14}BNO_3$
 $M_r = 315.12$
 Orthorhombic, $Pbca$
 $a = 14.7245$ (13) Å
 $b = 11.196$ (1) Å
 $c = 18.4060$ (16) Å

$V = 3034.3$ (5) Å³
 $Z = 8$
 Mo $K\alpha$ radiation
 $\mu = 0.09$ mm⁻¹
 $T = 293$ K
 $0.21 \times 0.20 \times 0.16$ mm

Data collection

Bruker SMART APEX CCD area-detector diffractometer
 Absorption correction: multi-scan (SADABS; Sheldrick, 1996)
 $T_{min} = 0.981$, $T_{max} = 0.985$

31865 measured reflections
 3317 independent reflections
 1942 reflections with $I > 2\sigma(I)$
 $R_{int} = 0.088$

Refinement

$R[F^2 > 2\sigma(F^2)] = 0.061$
 $wR(F^2) = 0.127$
 $S = 1.09$
 3317 reflections
 220 parameters
 1 restraint

H atoms treated by a mixture of independent and constrained refinement
 $\Delta\rho_{max} = 0.14$ e Å⁻³
 $\Delta\rho_{min} = -0.19$ e Å⁻³

Table 1

Hydrogen-bond geometry (Å, °).

$D-H\cdots A$	$D-H$	$H\cdots A$	$D\cdots A$	$D-H\cdots A$
$O3-H3'\cdots O2^i$	0.84 (2)	1.97 (2)	2.740 (2)	152 (3)
$C7-H7\cdots O3^{ii}$	0.93	2.53	3.165 (3)	126

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

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

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Supplementary data and figures for this paper are available from the IUCr electronic archives (Reference: TK2636).

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supplementary materials

Acta Cryst. (2010). E66, o758 [doi:10.1107/S1600536810007609]

{2-Hydroxy-6-[(2-oxidophenyl)iminomethyl- κ^2N,O]phenolato- κO^1 }phenylboron

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Comment

Recently, boronates derived from ligands having electron-donating and -attracting groups have been prepared in order to generate chromophores with nonlinear optical properties (Reyes *et al.*, 2005; Lamère *et al.*, 2006; Muñoz *et al.*, 2008). Boronates derived from Schiff bases have been applied in Diels-Alder reactions and for the stereoselective addition of Grignard reagents to azomethine carbon atoms (Rodríguez *et al.*, 2005a, 2005b; López-Ruiz *et al.*, 2008).

We have synthesized the [4.3.0]heterobicycle (I), Fig. 1, which is composed of a five-membered OBNC₂ ring and a six-membered OBNC₃ ring. The N→B, B—O and B—C bond lengths are comparable to the values determined for the molecular structure of the boronate obtained from the unsubstituted ligand 2-[(2-hydroxyphenyl)imino]methyl]phenol (Höpfl *et al.*, 1998; Barba *et al.*, 2001); however, the sum of bond lengths at the boron atom is significantly shorter for (I) (6.15 versus 6.18 Å), indicating that there is a small, but systematic tendency for bond length shortening.

The bond angles around the boron atom vary from 98.72 (2) ° for O2—B1—N1 to 112.8 (2) ° for O2—B1—C14, thus indicating a significant distortion from ideal tetrahedral geometry, which can be seen also from the value for the tetrahedral character [71 %] (Höpfl, 1999).

In the five-membered ring, the bond angles vary from 98.7 (2) to 114.0 (2) ° giving a mean value of 106.9 °. In the six-membered heterocycle, the bond angles range from 106.6 (2) to 123.4 (2) °, and the mean value is 117.9 °. Both rings have an approximate envelope conformation, in which the boron atom is localized at the tip. The proximity to an ideal envelope can be illustrated by the N1—C8—C9—O2 torsion angle [−2.7 (3) °] in the first case, and the O1—C1—C2—C7 and C2—C7—N1—B1 torsion angles [2.3 (3) and 2.5 (3) °] in the second case.

The degree of π -delocalization between the aromatic rings connected by the imino group can be analyzed by the C—N—C—C torsion angles (Reyes *et al.*, 2005; Lamère *et al.*, 2006; Muñoz *et al.*, 2008). The C7—N1—C8—C13 torsion angle has a value of 26.1 (4) ° indicating that the π -delocalization is distorted. The distortion can be evidenced also by the relatively long N1—C7 bond [1.282 (3) Å].

In the crystal structure, neighboring molecules are associated through O3—H \cdots O2¹ hydrogen bonds [symmetry code: (i), -x, 2-y, 1-z] to form supramolecular dimeric units having crystallographic inversion symmetry (Fig. 2). The crystal packing is further stabilized by C—H \cdots O interactions (Table 1).

Experimental

For the preparation of (I), 2,3-dihydroxysalicylaldehyde (0.250 g, 1.81 mmol), 2-aminophenol (0.198 g, 1.81 mmol) and phenylboronic acid (0.221 g, 1.81 mmol) were dissolved in DMF (35 ml), and the solution was refluxed for 1 h in the presence of a Dean-Stark trap. After cooling, the product precipitated in the form of a yellow crystalline solid, which was filtered and washed with hexane. Yield: 0.102 g (18 %). M.pt. 531 K.

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MS (FAB⁺): m/z (%) = 316 ($[M+H]^+$, 40).

Refinement

H atoms were positioned geometrically and constrained using the riding-model approximation [$C-H = 0.93 \text{ \AA}$, $U_{\text{iso}}(\text{H}) = 1.2 U_{\text{eq}}(\text{C})$]. The hydrogen atom bonded to O3 was located in a difference Fourier map. Its coordinates were refined with a distance restraint: $O-H = 0.84 \pm 0.01 \text{ \AA}$ and [$U_{\text{iso}}(\text{H}) = 1.5 U_{\text{eq}}(\text{O})$].

Figures

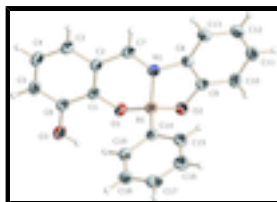


Fig. 1. Perspective view of (I), showing the atom-labelling scheme. Displacement ellipsoids are drawn at the 30% probability level and H atoms are shown as small spheres of arbitrary radii.

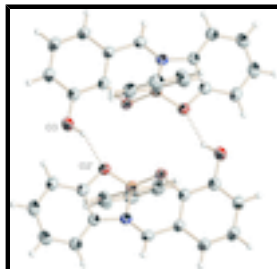


Fig. 2. Fragment of the crystal structure of (I), showing the supramolecular aggregate formed through O—H...O hydrogen bonding interactions. Displacement ellipsoids are drawn at the 30% probability level and H atoms are shown as small spheres of arbitrary radii.

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Crystal data

$C_{19}H_{14}BNO_3$

$M_r = 315.12$

Orthorhombic, *Pbca*

Hall symbol: -P 2ac 2ab

$a = 14.7245 (13) \text{ \AA}$

$b = 11.196 (1) \text{ \AA}$

$c = 18.4060 (16) \text{ \AA}$

$V = 3034.3 (5) \text{ \AA}^3$

$Z = 8$

$F(000) = 1312$

$D_x = 1.380 \text{ Mg m}^{-3}$

Melting point: 531 K

Mo $K\alpha$ radiation, $\lambda = 0.71073 \text{ \AA}$

Cell parameters from 3128 reflections

$\theta = 2.2\text{--}21.4^\circ$

$\mu = 0.09 \text{ mm}^{-1}$

$T = 293 \text{ K}$

Rectangular prism, orange

$0.21 \times 0.20 \times 0.16 \text{ mm}$

Data collection

Bruker SMART APEX CCD area-detector diffractometer

Radiation source: fine-focus sealed tube graphite

3317 independent reflections

1942 reflections with $I > 2\sigma(I)$

$R_{\text{int}} = 0.088$

Detector resolution: 8.3 pixels mm⁻¹
 phi and ω scans
 Absorption correction: multi-scan
 (SADABS; Sheldrick, 1996)
 $T_{\min} = 0.981$, $T_{\max} = 0.985$
 31865 measured reflections

$\theta_{\max} = 27.0^\circ$, $\theta_{\min} = 2.2^\circ$
 $h = -18 \rightarrow 18$
 $k = -14 \rightarrow 14$
 $l = -23 \rightarrow 23$

Refinement

Refinement on F^2
 Least-squares matrix: full
 $R[F^2 > 2\sigma(F^2)] = 0.061$
 $wR(F^2) = 0.127$
 $S = 1.09$
 3317 reflections
 220 parameters
 1 restraint

Primary atom site location: structure-invariant direct methods
 Secondary atom site location: difference Fourier map
 Hydrogen site location: inferred from neighbouring sites
 H atoms treated by a mixture of independent and constrained refinement
 $w = 1/[\sigma^2(F_o^2) + (0.0408P)^2 + 0.7703P]$
 where $P = (F_o^2 + 2F_c^2)/3$
 $(\Delta/\sigma)_{\max} = 0.001$
 $\Delta\rho_{\max} = 0.14 \text{ e } \text{Å}^{-3}$
 $\Delta\rho_{\min} = -0.19 \text{ e } \text{Å}^{-3}$

Special details

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

	<i>x</i>	<i>y</i>	<i>z</i>	$U_{\text{iso}}^*/U_{\text{eq}}$
B1	0.10076 (17)	0.8654 (3)	0.51225 (15)	0.0453 (7)
N1	0.20017 (12)	0.80938 (17)	0.50322 (10)	0.0460 (5)
O1	0.11329 (10)	0.98838 (14)	0.53706 (8)	0.0486 (4)
O2	0.07078 (10)	0.85831 (14)	0.43509 (8)	0.0514 (4)
O3	0.09544 (12)	1.17706 (17)	0.62833 (11)	0.0672 (6)
H3'	0.0545 (15)	1.147 (3)	0.6024 (14)	0.101*
C1	0.17986 (15)	1.0082 (2)	0.58667 (12)	0.0467 (6)
C2	0.25781 (16)	0.9374 (2)	0.59210 (13)	0.0479 (6)
C3	0.32673 (17)	0.9695 (2)	0.64063 (14)	0.0579 (7)
H3	0.3794	0.9238	0.6432	0.069*
C4	0.31745 (18)	1.0667 (3)	0.68403 (14)	0.0613 (7)

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H4	0.3633	1.0874	0.7164	0.074*
C5	0.23854 (18)	1.1354 (2)	0.67962 (13)	0.0582 (7)
H5	0.2319	1.2017	0.7096	0.070*
C6	0.17066 (16)	1.1066 (2)	0.63186 (13)	0.0518 (6)
C7	0.26874 (16)	0.8426 (2)	0.54104 (13)	0.0506 (6)
H7	0.3248	0.8055	0.5353	0.061*
C8	0.19835 (16)	0.7394 (2)	0.43960 (12)	0.0466 (6)
C9	0.12224 (16)	0.7743 (2)	0.40089 (13)	0.0482 (6)
C10	0.10560 (18)	0.7249 (2)	0.33315 (14)	0.0576 (7)
H10	0.0555	0.7478	0.3057	0.069*
C11	0.1662 (2)	0.6406 (3)	0.30787 (14)	0.0622 (7)
H11	0.1565	0.6067	0.2624	0.075*
C12	0.24025 (19)	0.6050 (3)	0.34753 (14)	0.0619 (7)
H12	0.2788	0.5465	0.3291	0.074*
C13	0.25807 (17)	0.6548 (2)	0.41423 (14)	0.0565 (7)
H13	0.3086	0.6321	0.4412	0.068*
C14	0.03782 (15)	0.7892 (2)	0.56575 (12)	0.0444 (6)
C15	0.00739 (17)	0.6756 (2)	0.54791 (15)	0.0599 (7)
H15	0.0247	0.6426	0.5036	0.072*
C16	-0.04753 (19)	0.6099 (2)	0.59356 (17)	0.0707 (8)
H16	-0.0665	0.5338	0.5800	0.085*
C17	-0.07412 (18)	0.6570 (3)	0.65887 (16)	0.0649 (8)
H17	-0.1107	0.6128	0.6901	0.078*
C18	-0.04662 (17)	0.7693 (3)	0.67788 (14)	0.0590 (7)
H18	-0.0651	0.8021	0.7219	0.071*
C19	0.00840 (16)	0.8338 (2)	0.63199 (13)	0.0523 (6)
H19	0.0265	0.9100	0.6459	0.063*

Atomic displacement parameters (\AA^2)

	U^{11}	U^{22}	U^{33}	U^{12}	U^{13}	U^{23}
B1	0.0384 (14)	0.0458 (16)	0.0517 (17)	0.0035 (13)	-0.0048 (12)	-0.0004 (14)
N1	0.0408 (11)	0.0516 (12)	0.0455 (12)	-0.0010 (9)	0.0017 (9)	0.0026 (10)
O1	0.0430 (9)	0.0478 (10)	0.0551 (10)	-0.0032 (8)	-0.0085 (8)	0.0019 (8)
O2	0.0503 (10)	0.0558 (11)	0.0482 (10)	0.0042 (8)	-0.0050 (8)	0.0005 (8)
O3	0.0517 (11)	0.0594 (12)	0.0905 (15)	-0.0025 (10)	-0.0096 (10)	-0.0189 (10)
C1	0.0441 (14)	0.0494 (15)	0.0466 (14)	-0.0100 (12)	-0.0008 (12)	0.0056 (12)
C2	0.0442 (14)	0.0526 (15)	0.0469 (14)	-0.0062 (12)	-0.0031 (12)	0.0021 (12)
C3	0.0456 (15)	0.0660 (19)	0.0619 (17)	-0.0046 (13)	-0.0079 (13)	0.0010 (15)
C4	0.0540 (17)	0.0685 (19)	0.0614 (17)	-0.0169 (15)	-0.0099 (13)	0.0012 (15)
C5	0.0594 (17)	0.0547 (17)	0.0605 (17)	-0.0124 (14)	-0.0017 (14)	-0.0047 (14)
C6	0.0471 (15)	0.0507 (16)	0.0577 (16)	-0.0095 (12)	0.0020 (12)	0.0026 (13)
C7	0.0409 (14)	0.0577 (17)	0.0533 (15)	0.0020 (12)	-0.0038 (12)	0.0076 (13)
C8	0.0448 (13)	0.0532 (15)	0.0418 (13)	-0.0043 (12)	0.0058 (11)	0.0010 (12)
C9	0.0461 (14)	0.0488 (15)	0.0496 (15)	-0.0032 (12)	0.0024 (12)	0.0041 (13)
C10	0.0598 (16)	0.0647 (18)	0.0483 (16)	-0.0095 (14)	-0.0046 (13)	0.0020 (14)
C11	0.0699 (19)	0.0683 (19)	0.0484 (15)	-0.0157 (16)	0.0126 (14)	-0.0058 (14)
C12	0.0578 (17)	0.0641 (19)	0.0637 (18)	-0.0029 (14)	0.0166 (15)	-0.0065 (15)

C13	0.0487 (15)	0.0663 (18)	0.0545 (16)	0.0001 (13)	0.0073 (13)	-0.0024 (14)
C14	0.0351 (12)	0.0475 (15)	0.0505 (15)	0.0023 (11)	-0.0046 (11)	0.0014 (12)
C15	0.0570 (16)	0.0528 (17)	0.0700 (18)	-0.0022 (14)	0.0112 (14)	-0.0056 (14)
C16	0.0662 (18)	0.0510 (17)	0.095 (2)	-0.0083 (14)	0.0173 (17)	0.0024 (16)
C17	0.0549 (16)	0.070 (2)	0.0702 (19)	-0.0027 (15)	0.0131 (14)	0.0159 (16)
C18	0.0501 (15)	0.074 (2)	0.0530 (16)	-0.0008 (14)	0.0025 (13)	0.0036 (14)
C19	0.0456 (14)	0.0590 (17)	0.0523 (16)	-0.0047 (12)	-0.0013 (12)	-0.0008 (13)

Geometric parameters (Å, °)

B1—O1	1.463 (3)	C8—C13	1.375 (3)
B1—O2	1.489 (3)	C8—C9	1.384 (3)
B1—C14	1.599 (4)	C9—C10	1.386 (3)
B1—N1	1.601 (3)	C10—C11	1.380 (4)
N1—C7	1.282 (3)	C10—H10	0.93
N1—C8	1.409 (3)	C11—C12	1.371 (4)
O1—C1	1.358 (3)	C11—H11	0.93
O2—C9	1.362 (3)	C12—C13	1.374 (3)
O3—C6	1.362 (3)	C12—H12	0.93
O3—H3'	0.84 (2)	C13—H13	0.93
C1—C6	1.387 (3)	C14—C19	1.387 (3)
C1—C2	1.399 (3)	C14—C15	1.388 (3)
C2—C3	1.399 (3)	C15—C16	1.379 (3)
C2—C7	1.426 (3)	C15—H15	0.93
C3—C4	1.357 (3)	C16—C17	1.370 (4)
C3—H3	0.93	C16—H16	0.93
C4—C5	1.396 (4)	C17—C18	1.367 (4)
C4—H4	0.93	C17—H17	0.93
C5—C6	1.370 (3)	C18—C19	1.375 (3)
C5—H5	0.93	C18—H18	0.93
C7—H7	0.93	C19—H19	0.93
O1—B1—O2	112.7 (2)	C9—C8—N1	106.6 (2)
O1—B1—C14	112.5 (2)	O2—C9—C8	114.0 (2)
O2—B1—C14	112.78 (19)	O2—C9—C10	126.4 (2)
O1—B1—N1	106.61 (18)	C8—C9—C10	119.6 (2)
O2—B1—N1	98.70 (18)	C11—C10—C9	117.5 (3)
C14—B1—N1	112.6 (2)	C11—C10—H10	121.2
C7—N1—C8	128.9 (2)	C9—C10—H10	121.2
C7—N1—B1	123.4 (2)	C12—C11—C10	122.3 (3)
C8—N1—B1	106.65 (18)	C12—C11—H11	118.9
C1—O1—B1	117.06 (18)	C10—C11—H11	118.9
C9—O2—B1	108.21 (18)	C11—C12—C13	120.6 (3)
C6—O3—H3'	112 (2)	C11—C12—H12	119.7
O1—C1—C6	117.6 (2)	C13—C12—H12	119.7
O1—C1—C2	123.2 (2)	C12—C13—C8	117.5 (3)
C6—C1—C2	119.2 (2)	C12—C13—H13	121.3
C1—C2—C3	119.7 (2)	C8—C13—H13	121.3
C1—C2—C7	117.9 (2)	C19—C14—C15	115.9 (2)
C3—C2—C7	122.0 (2)	C19—C14—B1	122.1 (2)

supplementary materials

C4—C3—C2	120.6 (3)	C15—C14—B1	122.0 (2)
C4—C3—H3	119.7	C16—C15—C14	122.3 (3)
C2—C3—H3	119.7	C16—C15—H15	118.9
C3—C4—C5	119.4 (2)	C14—C15—H15	118.9
C3—C4—H4	120.3	C17—C16—C15	119.8 (3)
C5—C4—H4	120.3	C17—C16—H16	120.1
C6—C5—C4	121.0 (3)	C15—C16—H16	120.1
C6—C5—H5	119.5	C18—C17—C16	119.6 (3)
C4—C5—H5	119.5	C18—C17—H17	120.2
O3—C6—C5	119.2 (2)	C16—C17—H17	120.2
O3—C6—C1	120.7 (2)	C17—C18—C19	120.0 (3)
C5—C6—C1	120.1 (2)	C17—C18—H18	120.0
N1—C7—C2	119.0 (2)	C19—C18—H18	120.0
N1—C7—H7	120.5	C18—C19—C14	122.4 (2)
C2—C7—H7	120.5	C18—C19—H19	118.8
C13—C8—C9	122.5 (2)	C14—C19—H19	118.8
C13—C8—N1	130.8 (2)		
O1—B1—N1—C7	29.0 (3)	C7—N1—C8—C13	26.0 (4)
O2—B1—N1—C7	145.9 (2)	B1—N1—C8—C13	-165.7 (3)
C14—B1—N1—C7	-94.8 (3)	C7—N1—C8—C9	-151.6 (2)
O1—B1—N1—C8	-140.00 (19)	B1—N1—C8—C9	16.7 (2)
O2—B1—N1—C8	-23.1 (2)	B1—O2—C9—C8	-13.6 (3)
C14—B1—N1—C8	96.1 (2)	B1—O2—C9—C10	167.3 (2)
O2—B1—O1—C1	-146.93 (18)	C13—C8—C9—O2	179.4 (2)
C14—B1—O1—C1	84.2 (2)	N1—C8—C9—O2	-2.7 (3)
N1—B1—O1—C1	-39.7 (3)	C13—C8—C9—C10	-1.5 (4)
O1—B1—O2—C9	133.55 (19)	N1—C8—C9—C10	176.3 (2)
C14—B1—O2—C9	-97.7 (2)	O2—C9—C10—C11	180.0 (2)
N1—B1—O2—C9	21.4 (2)	C8—C9—C10—C11	1.0 (4)
B1—O1—C1—C6	-154.4 (2)	C9—C10—C11—C12	0.4 (4)
B1—O1—C1—C2	28.1 (3)	C10—C11—C12—C13	-1.5 (4)
O1—C1—C2—C3	174.7 (2)	C11—C12—C13—C8	1.0 (4)
C6—C1—C2—C3	-2.7 (3)	C9—C8—C13—C12	0.5 (4)
O1—C1—C2—C7	2.3 (3)	N1—C8—C13—C12	-176.8 (2)
C6—C1—C2—C7	-175.1 (2)	O1—B1—C14—C19	-6.9 (3)
C1—C2—C3—C4	2.0 (4)	O2—B1—C14—C19	-135.7 (2)
C7—C2—C3—C4	174.1 (2)	N1—B1—C14—C19	113.6 (2)
C2—C3—C4—C5	-0.4 (4)	O1—B1—C14—C15	171.4 (2)
C3—C4—C5—C6	-0.4 (4)	O2—B1—C14—C15	42.6 (3)
C4—C5—C6—O3	-178.9 (2)	N1—B1—C14—C15	-68.0 (3)
C4—C5—C6—C1	-0.3 (4)	C19—C14—C15—C16	-1.0 (4)
O1—C1—C6—O3	2.8 (3)	B1—C14—C15—C16	-179.4 (2)
C2—C1—C6—O3	-179.6 (2)	C14—C15—C16—C17	0.3 (4)
O1—C1—C6—C5	-175.7 (2)	C15—C16—C17—C18	0.6 (4)
C2—C1—C6—C5	1.8 (3)	C16—C17—C18—C19	-0.9 (4)
C8—N1—C7—C2	164.0 (2)	C17—C18—C19—C14	0.1 (4)
B1—N1—C7—C2	-2.5 (3)	C15—C14—C19—C18	0.8 (3)
C1—C2—C7—N1	-14.9 (3)	B1—C14—C19—C18	179.2 (2)
C3—C2—C7—N1	172.9 (2)		

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>
O3—H3' \cdots O2 ⁱ	0.84 (2)	1.97 (2)	2.740 (2)	152 (3)
C7—H7 \cdots O3 ⁱⁱ	0.93	2.53	3.165 (3)	126

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

Fig. 1

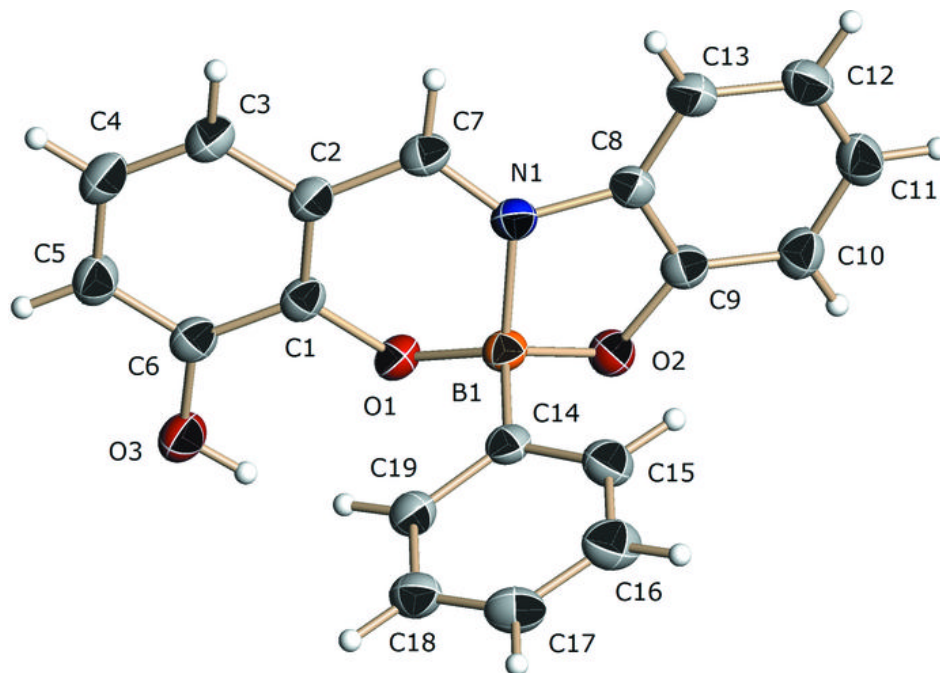


Fig. 2

