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1,2-Bis(3-phenoxybenzylidene)hydrazine

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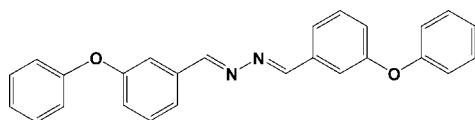
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Key indicators: single-crystal X-ray study; $T = 170$ K; mean $\sigma(C-C) = 0.002$ Å; R factor = 0.050; wR factor = 0.144; data-to-parameter ratio = 17.2.

Molecules of the title compound, $C_{26}H_{20}N_2O_2$, reside on crystallographic centres of inversion located at the mid-point of the N–N bond. The benzylidene ring is essentially coplanar with the central hydrazine group, with an interplanar angle of $4.5(2)^\circ$, whereas the phenyl ring is oriented at $34.0(3)^\circ$ with respect to the mean plane of the central 1,2-dibenzylidenehydrazine group. In the crystal, C–H $\cdots\pi$ (arene)-ring interactions link molecules about inversion centres.

Related literature

For the biological activity of Schiff bases, see: Aydogan *et al.* (2001); Desai *et al.* (2001); El-masry *et al.* (2000); Hodnett & Dunn (1970); Kundu *et al.* (2005); Pandeya *et al.* (1999); Singh & Dash (1988); Taggi *et al.* (2002); Xu *et al.* (1997); For crystallography and coordination chemistry of compounds containing the azine functionality or a diimine linkage, see: Xu *et al.* (1997); Kundu *et al.* (2005); For related structures, see: Liu *et al.* (2007); Odabaşoğlu *et al.* (2007); Zhang & Zheng (2008); Zheng *et al.* (2005a,b). For standard bond lengths, see Allen *et al.* (1987).



Experimental

Crystal data

$C_{26}H_{20}N_2O_2$
 $M_r = 392.44$
 Monoclinic, $C2/c$
 $a = 23.6271(16)$ Å
 $b = 11.2942(6)$ Å
 $c = 8.2359(7)$ Å
 $\beta = 109.538(8)^\circ$

$V = 2071.2(3)$ Å³
 $Z = 4$
 Mo $K\alpha$ radiation
 $\mu = 0.08$ mm⁻¹
 $T = 170$ K
 $0.44 \times 0.20 \times 0.06$ mm

Data collection

Oxford Diffraction Xcalibur Eos Gemini diffractometer
 Absorption correction: multi-scan (*CrysAlis RED*; Oxford Diffraction, 2010)
 $T_{\min} = 0.966$, $T_{\max} = 0.995$
 4228 measured reflections
 2341 independent reflections
 1643 reflections with $I > 2\sigma(I)$
 $R_{\text{int}} = 0.023$

Refinement

$R[F^2 > 2\sigma(F^2)] = 0.050$
 $wR(F^2) = 0.144$
 $S = 1.05$
 2341 reflections
 136 parameters
 H-atom parameters constrained
 $\Delta\rho_{\max} = 0.14$ e Å⁻³
 $\Delta\rho_{\min} = -0.19$ e Å⁻³

Table 1

Hydrogen-bond geometry (Å, °).

Cg is the centroid of the C7–C12 benzylidene ring.

D—H \cdots A	D—H	H \cdots A	D \cdots A	D—H \cdots A
C5—H5A \cdots Cg ⁱ	0.93	2.68	3.5947 (17)	167

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

Data collection: *CrysAlis PRO* (Oxford Diffraction, 2010); cell refinement: *CrysAlis PRO*; data reduction: *CrysAlis RED* (Oxford Diffraction, 2010); program(s) used to solve structure: *SHELXS97* (Sheldrick, 2008); program(s) used to refine structure: *SHELXL97* (Sheldrick, 2008); molecular graphics: *SHELXTL* (Sheldrick, 2008); software used to prepare material for publication: *SHELXTL*.

ASP thanks the University of Mysore for research facilities. JPJ acknowledges the NSF–MRI program (grant No. CHE1039027) for funds to purchase the X-ray diffractometer.

Supplementary data and figures for this paper are available from the IUCr electronic archives (Reference: GG2067).

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supplementary materials

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1,2-Bis(3-phenoxybenzylidene)hydrazine

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Comment

Schiff bases are known to exhibit biological activity such as antimicrobial (El-masry *et al.*, 2000 & Pandeya *et al.*, 1999), antifungal (Singh & Dash, 1988), antitumor (Hodnett & Dunn, 1970; Desai *et al.*, 2001) and as herbicides. Moreover, Schiff bases are used as substrates in the preparation of number of industrial and biologically active compounds *via* ring closure, cycloaddition and replacement reactions. Schiff bases have also been employed as ligands for complexation of metal ions (Aydogan *et al.*, 2001). On the industrial scale, they have wide range of applications such as dyes and pigments (Taggi *et al.*, 2002). Compounds containing an azine functionality or a diimine linkage have been investigated in terms of their crystallography and coordination chemistry (Xu *et al.*, 1997; Kundu *et al.*, 2005).

The crystal structures of some Schiff base hydrazines, *viz.*, 4-fluorobenzaldehyde[(E)-4-fluorobenzylidene]hydrazone (Odabaşoğlu *et al.*, 2007), N,N'-bis(3-nitrobenzylidene)hydrazine (Zheng *et al.*, 2005a), N,N'-Bis(4-chlorobenzylidene)hydrazine (Zheng *et al.*, 2005b), 1,2-bis(2-chlorobenzylidene)hydrazine (Zhang & Zheng, 2008), N,N'-Bis(4-hydroxybenzylidene)hydrazine (Liu *et al.*, 2007) have been reported. In view of the importance of Schiff base hydrazines, the crystal structure of title compound (I) is reported herein.

The complete molecule of the title compound, C₂₆H₂₀N₂O₂, is generated by the application of a centre of inversion (Fig. 1). The benzylidene ring is essentially coplanar with the central hydrazine group [dihedral angle = 4.5 (2)°] The dihedral angle between the mean plane of the 1,2-bis(benzylidene) hydrazine group and the two parallel phenyl rings is 34.0 (3)°. Weak C—H...Cg π -ring intermolecular interactions are observed (Table 1) providing some crystal packing stability (Fig. 2).

Experimental

A mixture of 3-phenoxybenzaldehyde (0.02 mol) and hydrazine hydrate (0.012 mol) was refluxed in 15 ml of absolute alcohol containing 2 drops of sulfuric acid, for about 3 hours. On cooling, the resulting solid was filtered and dried. Single crystals were grown from DMF (dimethylformamide) by the slow evaporation method. Yield: 86%. (m.p.: 404 K).

Refinement

All of the H atoms were placed in their calculated positions and then refined using the riding model with Atom—H lengths of 0.93 Å (C—H). Isotropic displacement parameters for these atoms were set to 1.19–1.20 (CH) times U_{eq} of the parent atom.

Figures

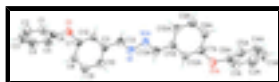


Fig. 1. Molecular structure of the title compound showing the atom labeling scheme and 50% probability displacement ellipsoids.

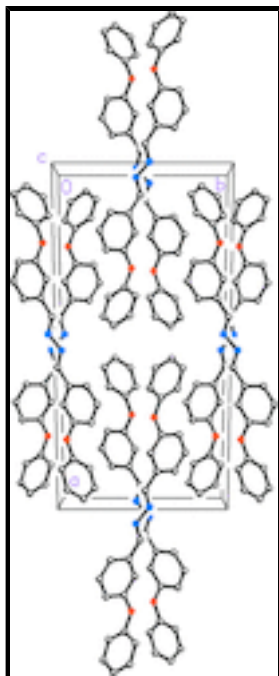


Fig. 2. Packing diagram of the title compound viewed along the *c* axis. The H atoms have been removed for clarity.

1,2-Bis(3-phenoxybenzylidene)hydrazine

Crystal data

$C_{26}H_{20}N_2O_2$

$M_r = 392.44$

Monoclinic, $C2/c$

Hall symbol: $-C\ 2yc$

$a = 23.6271$ (16) Å

$b = 11.2942$ (6) Å

$c = 8.2359$ (7) Å

$\beta = 109.538$ (8)°

$V = 2071.2$ (3) Å³

$Z = 4$

$F(000) = 824$

$D_x = 1.259$ Mg m⁻³

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

Cell parameters from 1146 reflections

$\theta = 3.1$ – 28.6 °

$\mu = 0.08$ mm⁻¹

$T = 170$ K

Plate, yellow

$0.44 \times 0.20 \times 0.06$ mm

Data collection

Oxford Diffraction Xcalibur Eos Gemini diffractometer

Radiation source: Enhance (Mo) X-ray Source graphite

Detector resolution: 16.1500 pixels mm⁻¹

ω scans

Absorption correction: multi-scan (*Crys.Alis RED*; Oxford Diffraction, 2010)

$T_{\min} = 0.966$, $T_{\max} = 0.995$

4228 measured reflections

2341 independent reflections

1643 reflections with $I > 2\sigma(I)$

$R_{\text{int}} = 0.023$

$\theta_{\max} = 28.7$ °, $\theta_{\min} = 3.1$ °

$h = -31 \rightarrow 19$

$k = -12 \rightarrow 14$

$l = -10 \rightarrow 10$

Refinement

Refinement on F^2	Primary atom site location: structure-invariant direct methods
Least-squares matrix: full	Secondary atom site location: difference Fourier map
$R[F^2 > 2\sigma(F^2)] = 0.050$	Hydrogen site location: inferred from neighbouring sites
$wR(F^2) = 0.144$	H-atom parameters constrained
$S = 1.05$	$w = 1/[\sigma^2(F_o^2) + (0.0591P)^2 + 0.4457P]$
2341 reflections	where $P = (F_o^2 + 2F_c^2)/3$
136 parameters	$(\Delta/\sigma)_{\max} < 0.001$
0 restraints	$\Delta\rho_{\max} = 0.14 \text{ e } \text{\AA}^{-3}$
	$\Delta\rho_{\min} = -0.19 \text{ e } \text{\AA}^{-3}$

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 (\AA^2)

	x	y	z	$U_{\text{iso}}^*/U_{\text{eq}}$
O1	0.28415 (5)	0.56492 (12)	0.14723 (17)	0.0634 (4)
N1	0.02329 (6)	0.53855 (12)	0.04610 (19)	0.0533 (4)
C1	0.36654 (9)	0.54126 (16)	0.4051 (2)	0.0582 (5)
H1B	0.3467	0.4787	0.4361	0.070*
C2	0.42338 (10)	0.5731 (2)	0.5077 (3)	0.0769 (7)
H2A	0.4421	0.5321	0.6096	0.092*
C3	0.45254 (9)	0.6644 (2)	0.4612 (3)	0.0818 (7)
H3A	0.4911	0.6854	0.5310	0.098*
C4	0.42510 (10)	0.7249 (2)	0.3120 (3)	0.0760 (7)
H4A	0.4450	0.7870	0.2802	0.091*
C5	0.36825 (9)	0.69433 (15)	0.2091 (2)	0.0584 (5)
H5A	0.3493	0.7355	0.1076	0.070*
C6	0.33985 (7)	0.60286 (14)	0.2574 (2)	0.0432 (4)
C7	0.23306 (7)	0.60721 (15)	0.1737 (2)	0.0442 (4)
C8	0.23209 (7)	0.70730 (14)	0.2691 (2)	0.0483 (4)
H8A	0.2672	0.7497	0.3218	0.058*
C9	0.17840 (8)	0.74360 (15)	0.2851 (2)	0.0491 (4)
H9A	0.1776	0.8111	0.3491	0.059*

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C10	0.12623 (7)	0.68200 (14)	0.2084 (2)	0.0462 (4)
H10A	0.0905	0.7071	0.2213	0.055*
C11	0.12717 (7)	0.58123 (13)	0.1108 (2)	0.0416 (4)
C12	0.18101 (7)	0.54480 (14)	0.0936 (2)	0.0441 (4)
H12A	0.1820	0.4783	0.0281	0.053*
C13	0.07271 (7)	0.51346 (15)	0.0257 (2)	0.0469 (4)
H13A	0.0743	0.4504	-0.0452	0.056*

Atomic displacement parameters (\AA^2)

	U^{11}	U^{22}	U^{33}	U^{12}	U^{13}	U^{23}
O1	0.0376 (6)	0.0815 (9)	0.0687 (9)	-0.0038 (6)	0.0148 (6)	-0.0324 (7)
N1	0.0396 (8)	0.0516 (9)	0.0670 (10)	-0.0047 (6)	0.0156 (7)	-0.0052 (7)
C1	0.0663 (12)	0.0553 (11)	0.0548 (11)	-0.0063 (9)	0.0226 (9)	0.0036 (9)
C2	0.0680 (14)	0.0838 (16)	0.0605 (13)	0.0133 (13)	-0.0029 (11)	0.0005 (11)
C3	0.0425 (11)	0.0966 (18)	0.0949 (18)	-0.0106 (12)	0.0079 (11)	-0.0309 (15)
C4	0.0707 (14)	0.0690 (14)	0.0975 (18)	-0.0299 (12)	0.0403 (14)	-0.0178 (13)
C5	0.0670 (12)	0.0496 (10)	0.0595 (11)	-0.0038 (9)	0.0225 (10)	0.0045 (9)
C6	0.0354 (8)	0.0473 (9)	0.0479 (9)	-0.0026 (7)	0.0152 (7)	-0.0100 (7)
C7	0.0383 (8)	0.0501 (9)	0.0433 (9)	0.0021 (7)	0.0123 (7)	-0.0041 (7)
C8	0.0431 (9)	0.0499 (10)	0.0466 (9)	-0.0039 (8)	0.0080 (7)	-0.0081 (8)
C9	0.0516 (10)	0.0447 (9)	0.0482 (10)	0.0061 (8)	0.0131 (8)	-0.0054 (8)
C10	0.0434 (9)	0.0477 (9)	0.0479 (9)	0.0082 (7)	0.0157 (7)	0.0030 (7)
C11	0.0407 (8)	0.0412 (9)	0.0425 (9)	0.0005 (7)	0.0134 (7)	0.0047 (7)
C12	0.0423 (9)	0.0431 (9)	0.0459 (9)	-0.0014 (7)	0.0133 (7)	-0.0063 (7)
C13	0.0436 (9)	0.0453 (9)	0.0527 (10)	-0.0013 (7)	0.0175 (8)	-0.0007 (8)

Geometric parameters (\AA , $^\circ$)

O1—C7	1.3814 (18)	C5—H5A	0.9300
O1—C6	1.3930 (18)	C7—C12	1.379 (2)
N1—C13	1.266 (2)	C7—C8	1.381 (2)
N1—N1 ⁱ	1.410 (3)	C8—C9	1.381 (2)
C1—C6	1.359 (2)	C8—H8A	0.9300
C1—C2	1.374 (3)	C9—C10	1.372 (2)
C1—H1B	0.9300	C9—H9A	0.9300
C2—C3	1.364 (3)	C10—C11	1.398 (2)
C2—H2A	0.9300	C10—H10A	0.9300
C3—C4	1.365 (3)	C11—C12	1.388 (2)
C3—H3A	0.9300	C11—C13	1.459 (2)
C4—C5	1.371 (3)	C12—H12A	0.9300
C4—H4A	0.9300	C13—H13A	0.9300
C5—C6	1.362 (2)		
C7—O1—C6	118.43 (12)	C12—C7—O1	115.75 (14)
C13—N1—N1 ⁱ	112.25 (17)	C8—C7—O1	123.66 (14)
C6—C1—C2	118.82 (18)	C9—C8—C7	119.21 (15)
C6—C1—H1B	120.6	C9—C8—H8A	120.4
C2—C1—H1B	120.6	C7—C8—H8A	120.4

C3—C2—C1	120.4 (2)	C10—C9—C8	121.21 (15)
C3—C2—H2A	119.8	C10—C9—H9A	119.4
C1—C2—H2A	119.8	C8—C9—H9A	119.4
C2—C3—C4	119.92 (19)	C9—C10—C11	119.53 (15)
C2—C3—H3A	120.0	C9—C10—H10A	120.2
C4—C3—H3A	120.0	C11—C10—H10A	120.2
C3—C4—C5	120.2 (2)	C12—C11—C10	119.42 (14)
C3—C4—H4A	119.9	C12—C11—C13	119.11 (14)
C5—C4—H4A	119.9	C10—C11—C13	121.47 (14)
C6—C5—C4	119.14 (19)	C7—C12—C11	120.06 (15)
C6—C5—H5A	120.4	C7—C12—H12A	120.0
C4—C5—H5A	120.4	C11—C12—H12A	120.0
C1—C6—C5	121.55 (17)	N1—C13—C11	121.53 (15)
C1—C6—O1	118.67 (15)	N1—C13—H13A	119.2
C5—C6—O1	119.58 (16)	C11—C13—H13A	119.2
C12—C7—C8	120.56 (15)		
C6—C1—C2—C3	-0.5 (3)	O1—C7—C8—C9	178.76 (16)
C1—C2—C3—C4	0.2 (3)	C7—C8—C9—C10	0.1 (3)
C2—C3—C4—C5	0.1 (3)	C8—C9—C10—C11	-0.6 (2)
C3—C4—C5—C6	-0.2 (3)	C9—C10—C11—C12	0.3 (2)
C2—C1—C6—C5	0.4 (3)	C9—C10—C11—C13	-179.48 (15)
C2—C1—C6—O1	175.27 (16)	C8—C7—C12—C11	-1.1 (3)
C4—C5—C6—C1	0.0 (3)	O1—C7—C12—C11	-179.24 (14)
C4—C5—C6—O1	-174.87 (16)	C10—C11—C12—C7	0.5 (2)
C7—O1—C6—C1	88.2 (2)	C13—C11—C12—C7	-179.68 (14)
C7—O1—C6—C5	-96.80 (19)	N1 ⁱ —N1—C13—C11	-179.40 (15)
C6—O1—C7—C12	-163.42 (14)	C12—C11—C13—N1	175.49 (16)
C6—O1—C7—C8	18.5 (2)	C10—C11—C13—N1	-4.7 (3)
C12—C7—C8—C9	0.7 (3)		

Symmetry codes: (i) $-x, -y+1, -z$.

Hydrogen-bond geometry ($\text{\AA}, ^\circ$)

C_g is the centroid of the C7—C12 benzylidene ring.

$D-H\cdots A$	$D-H$	$H\cdots A$	$D\cdots A$	$D-H\cdots A$
C5—H5A \cdots Cg ⁱⁱ	0.93	2.68	3.5947 (17)	167

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

Fig. 1

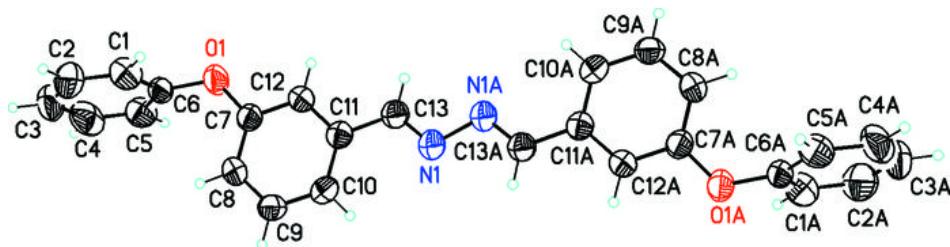


Fig. 2

