



Crystal structure and Hirshfeld surface analysis of (3*Z*)-7-methoxy-3-(2-phenylhydrazinylidene)-1-benzofuran-2(3*H*)-one

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Received 30 June 2021
Accepted 2 August 2021

Edited by M. Weil, Vienna University of Technology, Austria

Keywords: crystal structure; 2,3-dihydro-1-benzofuran ring system; dimers; hydrogen bonds; Hirshfeld surface analysis.

CCDC reference: 1984938

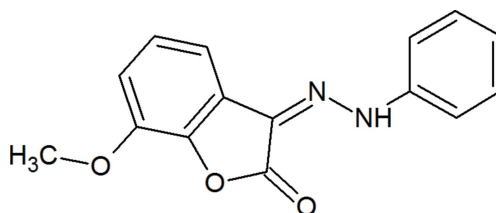
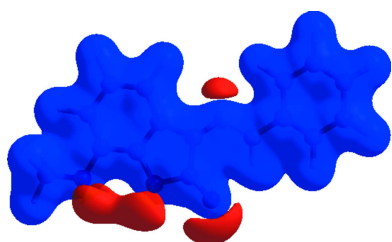
Supporting information: this article has supporting information at journals.iucr.org/e

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In the title compound, C₁₅H₁₂N₂O₃, pairs of molecules are linked into dimers by N—H···O hydrogen bonds, forming an *R*₂²(12) ring motif, with the dimers stacked along the *a* axis. These dimers are connected through π – π stacking interactions between the centroids of the benzene and furan rings of their 2,3-dihydro-1-benzofuran ring systems. Furthermore, there exists a C—H··· π interaction that consolidates the crystal packing. A Hirshfeld surface analysis indicates that the most important contacts are H···H (40.7%), O···H/H···O (24.7%), C···H/H···C (16.1%) and C···C (8.8%).

1. Chemical context

Hydrazones are a versatile class of organic ligands that have extensive applications in synthetic transformations, the synthesis of bioactive compounds, the design of materials and in coordination chemistry (Ma *et al.*, 2017*a,b*; Viswanathan *et al.*, 2019). Moreover, metal complexes of hydrazone ligands have been successfully applied as catalysts in organic synthesis (Gurbanov *et al.*, 2018). The properties of metal-hydrzonates can be regulated by the design of ligands through the involvement of non-covalent-bond donor or acceptor substituents (Ma *et al.*, 2020, 2021; Mahmudov *et al.*, 2013). Supramolecular networks of all dimensions in the crystal structures of hydrazone compounds or metal-hydrzonates, resulting from extensive hydrogen-bonding and other types of intermolecular interactions, have been reported (Gurbanov *et al.*, 2020*a*; Kopylovich *et al.*, 2011). Thus, the attachment of suitable substituents or synthons to hydrazone ligands can improve their functional properties and the catalytic or biological activity of the corresponding coordination compounds (Mizar *et al.*, 2012; Gurbanov *et al.*, 2020*a,b*; Khalilov *et al.*, 2018*a,b*; Maharramov *et al.*, 2018; Shihkaliyev *et al.*, 2019; Shixaliyev *et al.*, 2014).



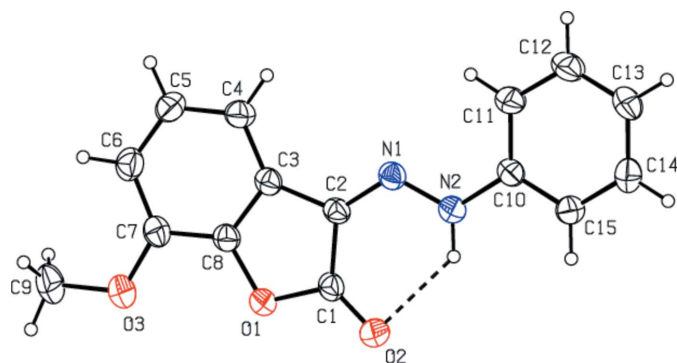


Figure 1
The title molecule with the labelling scheme and displacement ellipsoids drawn at the 30% probability level. The intramolecular N—H···O hydrogen bond is shown as a dashed line.

In a continuation of our work in this context (Atioğlu *et al.*, 2020, 2021), we have synthesized a new hydrazone compound, (3*Z*)-7-methoxy-3-(2-phenylhydrazinylidene)-1-benzofuran-2(3*H*)-one, which shows multiple intermolecular non-covalent interactions.

2. Structural commentary

In the title compound, the molecular conformation is stabilized by an intramolecular N2—H1···O2 hydrogen bond, forming an *S*(6) ring motif (Table 1, Fig. 1; Bernstein *et al.*, 1995). The 2,3-dihydro-1-benzofuran ring system (O1/C1—C8) is essentially planar [maximum deviation of 0.016 (2) Å for O1] and subtends a dihedral angle of 5.32 (14)° with the phenyl ring (C10—C15).

3. Supramolecular features

In the crystal, pairs of molecules are linked into dimers by intermolecular N—H···O hydrogen bonds, forming an *R*₂²(12)

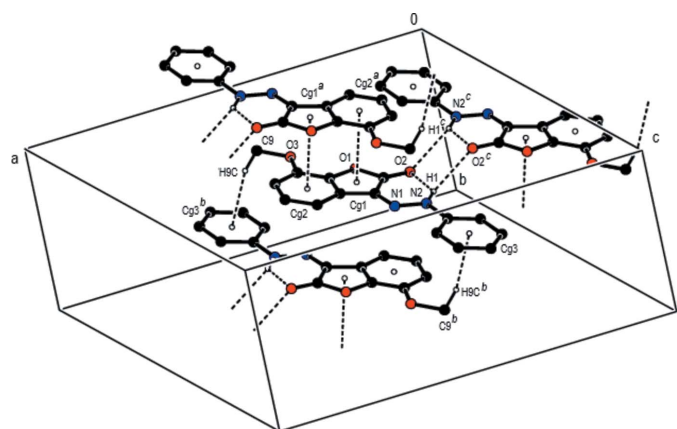


Figure 2
Intermolecular N—H···O hydrogen bonds, C—H··· π interactions and π — π stacking interactions (shown as dashed lines) in the title compound. [Symmetry codes: (a) $1 - x, -y, 1 - z$; (b) $1 - x, 1 - y, 1 - z$; (c) $\frac{1}{2} - x, \frac{1}{2} - y, 1 - z$].

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

Cg3 is the centroid of the C10—C15 phenyl ring.

<i>D</i> —H··· <i>A</i>	<i>D</i> —H	H··· <i>A</i>	<i>D</i> ··· <i>A</i>	<i>D</i> —H··· <i>A</i>
N2—H1···O2	0.92 (4)	2.14 (3)	2.843 (3)	133 (3)
N2—H1···O2 ⁱ	0.92 (4)	2.44 (4)	3.181 (4)	138 (3)
C9—H9C···Cg3 ⁱⁱ	0.96	2.70	3.555 (4)	149

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

ring motif (Table 1). These dimers are stacked along the *a* axis and connected by π — π stacking interactions between the centroids of the benzene and furan rings of their 2,3-dihydro-1-benzofuran ring systems [*Cg*1···*Cg*2($1 - x, -y, 1 - z$) = 3.5316 (19) Å, slippage = 0.352 Å, where *Cg*1 and *Cg*2 are the centroids of the benzene (C3—C8) and furan (O1/C1—C3/C8) rings, respectively] (Figs. 2, 3 and 4). Furthermore, there exists a C—H··· π interaction between the H9C atom of the methyl group C9 and the centroid of the phenyl ring (C10—C15).

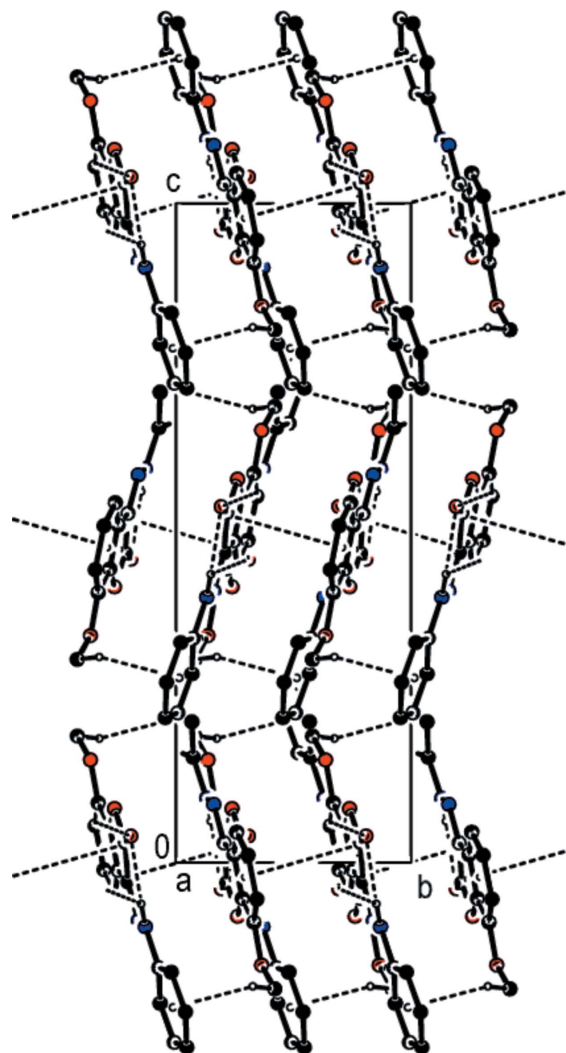


Figure 3
A view of the molecular packing of the title compound along the *a*-axis direction. Intermolecular interactions are depicted as in Fig. 2.

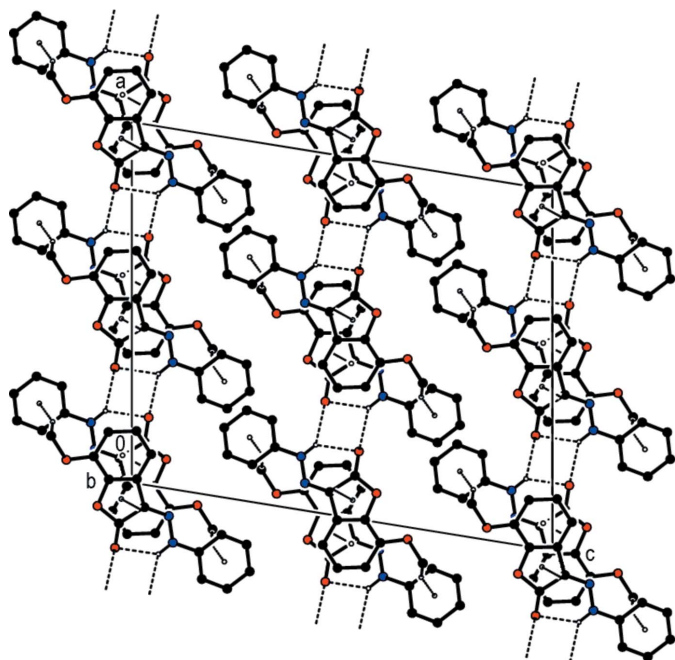


Figure 4
A view of the molecular packing of the title compound along the *b*-axis direction. Intermolecular interactions are depicted as in Fig. 2.

4. Hirshfeld surface analysis

Crystal Explorer 17.5 (Turner *et al.*, 2017) was used to calculate the Hirshfeld surfaces and generate the two-dimensional fingerprint plots. Hirshfeld surfaces allow for the display of intermolecular interactions by using distinct colours and intensities to indicate short and long contacts, as well as the

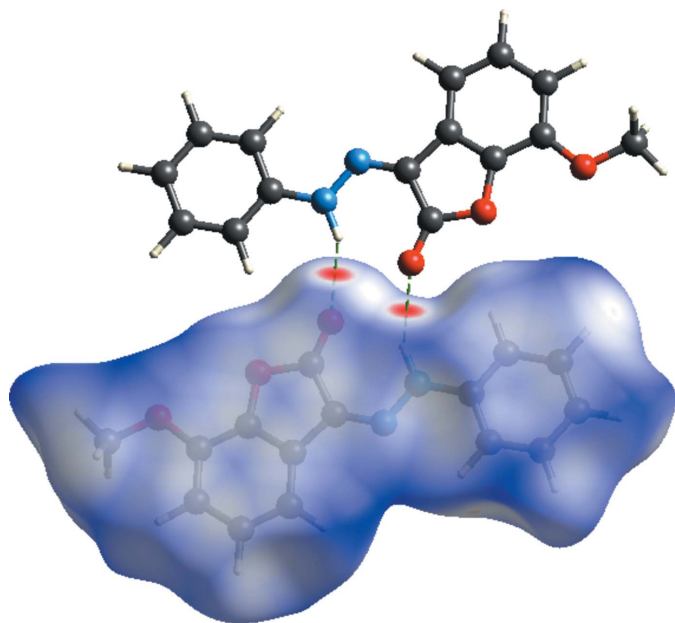


Figure 5
View of the three-dimensional Hirshfeld surface of the title compound plotted over d_{norm} in the range -0.1718 to 1.3843 a.u. The two $\text{N}-\text{H}\cdots\text{O}$ hydrogen bonds forming the dimer are depicted as dashed lines.

Table 2
Interatomic contacts of the title compound (\AA).

Contact	Distance	Symmetry operation
$\text{H1}\cdots\text{O2}$	2.44	$\frac{1}{2} - x, \frac{1}{2} - y, 1 - z$
$\text{H9B}\cdots\text{N2}$	2.91	$1 - x, -y, 1 - z$
$\text{H9C}\cdots\text{C11}$	2.93	$1 - x, 1 - y, 1 - z$
$\text{H5A}\cdots\text{H15A}$	2.51	$\frac{1}{2} + x, -\frac{1}{2} + y, z$
$\text{C9}\cdots\text{H14A}$	2.85	$\frac{1}{2} + x, \frac{1}{2} - y, -\frac{1}{2} + z$
$\text{H11A}\cdots\text{H11A}$	2.31	$1 - x, y, \frac{3}{2} - z$
$\text{C15}\cdots\text{H13A}$	3.07	$\frac{1}{2} - x, -\frac{1}{2} + y, \frac{3}{2} - z$

relative strength of the interactions. The three-dimensional Hirshfeld surface of the title compound plotted over d_{norm} in the range -0.1718 to 1.3843 a.u. is shown in Fig. 5. The $\text{N2}-\text{H1}\cdots\text{O2}$ interactions, which play a key role in the molecular packing of the title compound, are responsible for the red spot that occurs around O2. The bright-red spots appearing near O2 and hydrogen atom H1 indicate their roles as donors and/or acceptors in hydrogen-bonding; they also appear as blue and red regions corresponding to positive and negative potentials on the Hirshfeld surface mapped over electrostatic potential (Spackman *et al.*, 2008) shown in Fig. 6. Here the blue regions indicate positive electrostatic potential (hydrogen-bond donors), while the red regions indicate negative electrostatic potential (hydrogen-bond acceptors).

The overall two-dimensional fingerprint plot for the title compound is given in Fig. 7*a*, and those delineated into $\text{H}\cdots\text{H}$, $\text{O}\cdots\text{H}/\text{H}\cdots\text{O}$, $\text{C}\cdots\text{H}/\text{H}\cdots\text{C}$ and $\text{C}\cdots\text{C}$ contacts are shown in Fig. 7*b–e*, while numerical details of the different contacts are given in Table 2. The percentage contributions to the Hirshfeld surfaces from the various interatomic contacts are as follows: $\text{H}\cdots\text{H}$ (Fig. 7*b*; 40.7%), $\text{O}\cdots\text{H}/\text{H}\cdots\text{O}$ (Fig. 7*c*; 24.7%), $\text{C}\cdots\text{H}/\text{H}\cdots\text{C}$ (Fig. 7*d*; 16.1%) and $\text{C}\cdots\text{C}$ (Fig. 7*e*; 8.8%). Other minor contributions to the Hirshfeld surface are from $\text{N}\cdots\text{C}/\text{C}\cdots\text{N}$ (3.8%), $\text{N}\cdots\text{H}/\text{H}\cdots\text{N}$ (3.5%), $\text{O}\cdots\text{C}/\text{C}\cdots\text{O}$ (1.9%), $\text{O}\cdots\text{N}/\text{N}\cdots\text{O}$ (0.4%) and $\text{O}\cdots\text{O}$ (0.2%) contacts.

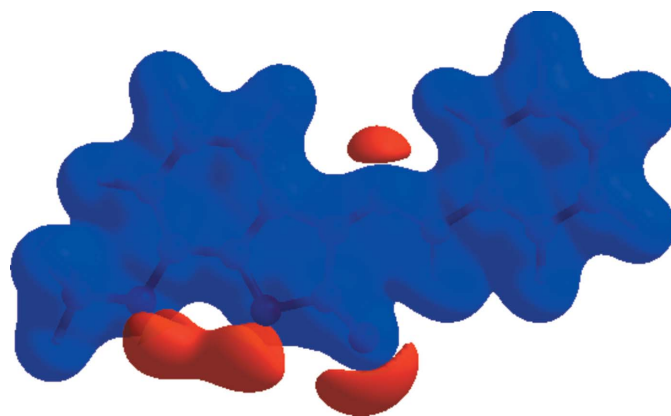
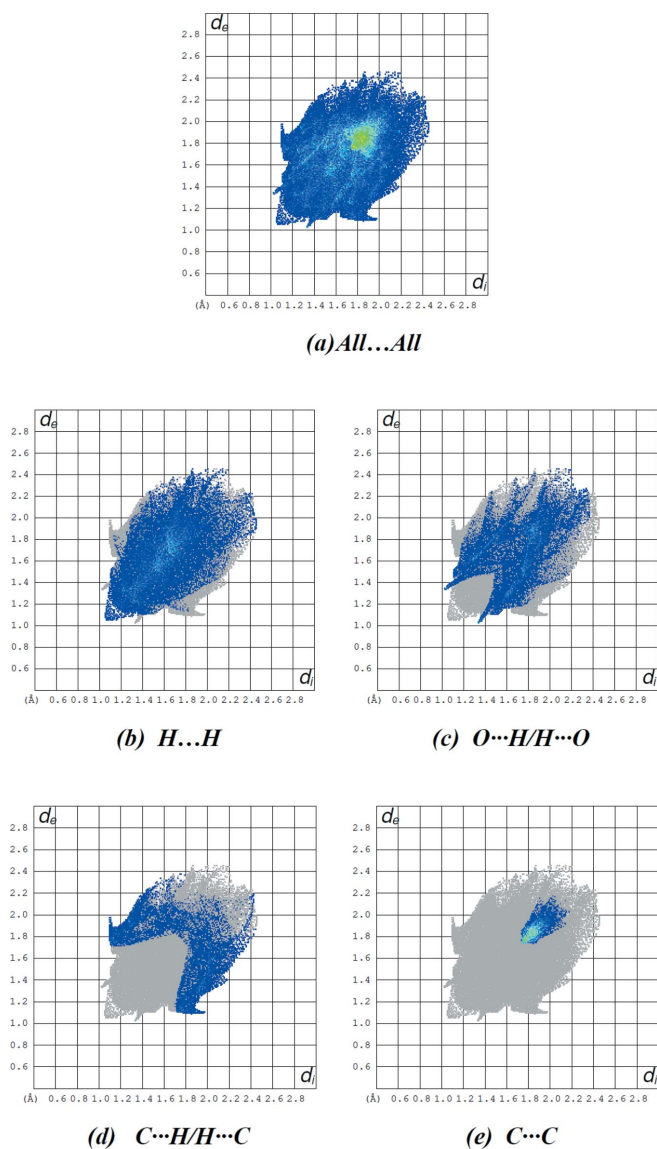


Figure 6
View of the three-dimensional Hirshfeld surface of the title compound plotted over electrostatic potential in the range -0.0500 to 0.0500 a.u. using the STO-3 G basis set at the Hartree–Fock level of theory. The hydrogen-bond donors and acceptors are viewed as blue and red regions, respectively, around atoms, corresponding to positive and negative potentials.


Figure 7

The full two-dimensional fingerprint plots for the title compound, showing (a) all interactions, and delineated into (b) H...H, (c) O...H/H...O, (d) C...H/H...C and (e) C...C interactions. The d_i and d_e values are the closest internal and external distances (in Å) from given points on the Hirshfeld surface contacts.

5. Database survey

A search of the Cambridge Crystallographic Database (CSD version 5.40, update of September 2019; Groom *et al.*, 2016) gave 763 hits for structures with a hydrazone moiety. Five structures that are closely related to the title compound are: 2-(4-nitro-1*H*-imidazol-1-yl)-*N'*-[1-(pyridin-2-yl)ethylidene]acetohydrazide (TODMEH; Oliveira *et al.*, 2019); 2-(2-nitro-1*H*-imidazol-1-yl)-*N'*-[1-(pyridin-2-yl)ethylidene]acetohydrazide (TODMIL; Oliveira *et al.*, 2019); 2-(4-nitro-1*H*-imidazol-1-yl)-*N'*-[phenyl(pyridin-2-yl)methylidene]acetohydrazide (TODMOR; Oliveira *et al.*, 2019); 2-(4-nitro-1*H*-imidazol-1-yl)-*N'*-[phenyl(pyridin-2-yl)methylidene]acetohydrazide (TODMUX; Oliveira *et al.*, 2019) and 1,1'-[1,3-phenyl-

enebis(2,2-dichloroethene-1,1-diyl)]bis(phenyldiazene) (EXIWOA; Shikhaliyev *et al.*, 2021).

TODMEH and TODMOR crystallize in the monoclinic space group $P2_1/c$ with $Z = 4$. TODMIL crystallizes in the monoclinic space group $I2/a$ with $Z = 8$ and TODMUX crystallizes in the triclinic space group $P\bar{1}$ with $Z = 2$. EXIWOA crystallizes in the monoclinic space group $P2_1/c$ with $Z = 4$. The *E* conformation in TODMEH, TODMIL and TODMUX is stabilized by a strong intermolecular N—H...O interaction. These interactions lead to the formation of dimeric structural arrangements. In the crystal packing of TODMOR, an intermolecular N—H...N interaction results in a zigzag structural arrangement, with the formation of chains along the crystallographic *b* axis. Non-classical intermolecular C—H...N and C—H...O interactions are also observed in the crystal structures of TODMEH, TODMIL, TODMOR and TODMUX. In EXIWOA, molecules are linked by C—H... π , C—Cl... π , Cl...Cl and Cl...H interactions, forming a three-dimensional supramolecular network.

6. Synthesis and crystallization

A 20 ml screw-neck vial was charged with dimethyl sulfoxide (DMSO; 10 ml), (*E*)-2-[[2-(3,5-dimethylphenyl)hydrazinylidene]methyl]phenol (240 mg, 1 mmol), tetramethylethylenediamine (TMEDA; 295 mg, 2.5 mmol), CuCl (2 mg, 0.02 mmol) and CCl₄ (20 mmol, 10 equiv). After 1–3 h (until TLC analysis showed complete consumption of the corresponding Schiff base), the reaction mixture was poured into a 0.01 *M* solution of HCl (100 mL, pH = 2–3), and extracted with dichloromethane (3 × 20 ml). The combined organic phase was washed with water (3 × 50 ml), brine (30 ml), dried over anhydrous Na₂SO₄ and concentrated *in vacuo* in a rotary evaporator. The residue was purified by column chromatography on silica gel using appropriate mixtures of hexane and dichloromethane ($v/v = 3/1$ – $1/1$). Colourless solid (yield 65%); m.p. 475 K. Analysis calculated for C₁₅H₁₂N₂O₃ ($M = 268.27$): C 67.16, H 4.51, N 10.44; found: C 67.11, H 4.47, N 10.35%. ¹H NMR (300 MHz, CDCl₃) δ 12.13 (*s*, 1H, NH), 6.91–7.43 (8H, Ar), 3.99 (*s*, 3H, OCH₃). ¹³C NMR (75 MHz, CDCl₃) δ 186.20, 161.87, 150.65, 141.76, 129.60, 125.09, 124.44, 123.87, 114.90, 112.74, 111.44, 108.76, 56.46. ESI-MS: m/z : 269.26 [$M + H$]⁺. Crystals suitable for X-ray analysis were obtained by slow evaporation of a dichloromethane solution.

7. Refinement details

Crystal data, data collection and structure refinement details are summarized in Table 3. The H atom of the NH group was located in a difference-Fourier map and refined freely [$N2-H1 = 0.92$ (4) Å]. H atoms bonded to C atoms were positioned geometrically and refined using a riding model, with C—H = 0.93 or 0.96 Å, and with $U_{iso}(H) = 1.2U_{eq}(C)$ for aromatic or $1.5U_{eq}(C)$ for methyl H atoms. Owing to poor agreement between observed and calculated intensities, seven outliers, ($\bar{1}\bar{3}$ 7 1), ($\bar{8}$ 6 13), (13 7 0), ($\bar{9}$ 5 19), ($\bar{7}$ 5 20), ($\bar{1}\bar{5}$ 5 12) and (0 6 16), were omitted in the final cycles of refinement.

Table 3

Experimental details.

Crystal data	
Chemical formula	C ₁₅ H ₁₂ N ₂ O ₃
<i>M_r</i>	268.27
Crystal system, space group	Monoclinic, <i>C2/c</i>
Temperature (K)	296
<i>a</i> , <i>b</i> , <i>c</i> (Å)	17.436 (2), 7.2485 (7), 20.595 (2)
β (°)	99.181 (4)
<i>V</i> (Å ³)	2569.6 (5)
<i>Z</i>	8
Radiation type	Mo <i>K</i> α
μ (mm ⁻¹)	0.10
Crystal size (mm)	0.49 × 0.15 × 0.06
Data collection	
Diffractometer	Bruker APEXII CCD
Absorption correction	Multi-scan (<i>SADABS</i> ; Krause <i>et al.</i> , 2015)
<i>T_{min}</i> , <i>T_{max}</i>	0.629, 0.745
No. of measured, independent and observed [<i>I</i> > 2 σ (<i>I</i>)] reflections	12800, 2427, 1463
<i>R_{int}</i>	0.085
(<i>sin</i> θ / λ) _{max} (Å ⁻¹)	0.616
Refinement	
<i>R</i> [<i>F</i> ² > 2 σ (<i>F</i> ²)], <i>wR</i> (<i>F</i> ²), <i>S</i>	0.073, 0.152, 1.01
No. of reflections	2427
No. of parameters	187
H-atom treatment	H atoms treated by a mixture of independent and constrained refinement
$\Delta\rho_{\text{max}}$, $\Delta\rho_{\text{min}}$ (e Å ⁻³)	0.18, -0.18

Computer programs: *APEX3* and *SAINT* (Bruker, 2017), *SHELXS* (Sheldrick, 2008), *SHELXL* (Sheldrick, 2015), *ORTEP-3 for Windows* (Farrugia, 2012) and *PLATON* (Spek, 2020).

Acknowledgements

The authors' contributions are as follows. Conceptualization, MA and UFA; methodology, ZA and SHM; investigation, SHM, RKA, and ZA; writing (original draft), MA and SM; writing (review and editing of the manuscript), MA and UFA; visualization, RKA, ZA and MA; funding acquisition, UFA, SHM and RKA; resources, RKA, ZA and SHM; supervision, MA and SM.

Funding information

This work was performed under the support of the Science Development Foundation under the President of the Republic of Azerbaijan (grant No. EIF-BGM-4- RFTF-1/2017-21/13/4).

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supporting information

Acta Cryst. (2021). E77, 907-911 [https://doi.org/10.1107/S2056989021007891]

Crystal structure and Hirshfeld surface analysis of (3*Z*)-7-methoxy-3-(2-phenylhydrazinylidene)-1-benzofuran-2(3*H*)-one

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Computing details

Data collection: *APEX3* (Bruker, 2017); cell refinement: *SAINTE* (Bruker, 2017); data reduction: *SAINTE* (Bruker, 2017); program(s) used to solve structure: *SHELXS* (Sheldrick, 2008); program(s) used to refine structure: *SHELXL* (Sheldrick, 2015); molecular graphics: *ORTEP-3 for Windows* (Farrugia, 2012); software used to prepare material for publication: *PLATON* (Spek, 2020).

(3*Z*)-7-Methoxy-3-(2-phenylhydrazinylidene)-1-benzofuran-2(3*H*)-one

Crystal data

$C_{15}H_{12}N_2O_3$

$M_r = 268.27$

Monoclinic, *C2/c*

$a = 17.436$ (2) Å

$b = 7.2485$ (7) Å

$c = 20.595$ (2) Å

$\beta = 99.181$ (4)°

$V = 2569.6$ (5) Å³

$Z = 8$

$F(000) = 1120$

$D_x = 1.387$ Mg m⁻³

Mo *K* α radiation, $\lambda = 0.71073$ Å

Cell parameters from 2240 reflections

$\theta = 2.4$ – 26.3 °

$\mu = 0.10$ mm⁻¹

$T = 296$ K

Prism, colourless

$0.49 \times 0.15 \times 0.06$ mm

Data collection

Bruker APEXII CCD
diffractometer

φ and ω scans

Absorption correction: multi-scan
(*SADABS*; Krause *et al.*, 2015)

$T_{\min} = 0.629$, $T_{\max} = 0.745$

12800 measured reflections

2427 independent reflections

1463 reflections with $I > 2\sigma(I)$

$R_{\text{int}} = 0.085$

$\theta_{\max} = 26.0$ °, $\theta_{\min} = 2.0$ °

$h = -21 \rightarrow 21$

$k = -8 \rightarrow 8$

$l = -25 \rightarrow 25$

Refinement

Refinement on F^2

Least-squares matrix: full

$R[F^2 > 2\sigma(F^2)] = 0.073$

$wR(F^2) = 0.152$

$S = 1.01$

2427 reflections

187 parameters

0 restraints

Primary atom site location: difference Fourier
map

Secondary atom site location: difference Fourier
map

Hydrogen site location: mixed

H atoms treated by a mixture of independent
and constrained refinement

$w = 1/[\sigma^2(F_o^2) + (0.0517P)^2 + 2.9831P]$

where $P = (F_o^2 + 2F_c^2)/3$

$$(\Delta/\sigma)_{\max} < 0.001$$

$$\Delta\rho_{\max} = 0.18 \text{ e } \text{\AA}^{-3}$$

$$\Delta\rho_{\min} = -0.18 \text{ e } \text{\AA}^{-3}$$

Extinction correction: SHELXL2016/6
(Sheldrick 2015),
 $Fc^* = kFc[1 + 0.001x\text{Fc}^2\lambda^3/\sin(2\theta)]^{-1/4}$
Extinction coefficient: 0.0015 (4)

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.

Fractional atomic coordinates and isotropic or equivalent isotropic displacement parameters (\AA^2)

	x	y	z	$U_{\text{iso}}^*/U_{\text{eq}}$
O1	0.42001 (12)	0.2382 (3)	0.41831 (10)	0.0546 (6)
O2	0.31175 (13)	0.3077 (4)	0.45997 (10)	0.0648 (7)
O3	0.53382 (14)	0.1359 (4)	0.34426 (11)	0.0728 (8)
N1	0.42389 (15)	0.3396 (4)	0.58811 (12)	0.0477 (7)
N2	0.35184 (16)	0.3699 (4)	0.59750 (13)	0.0500 (7)
C1	0.38107 (19)	0.2841 (5)	0.46932 (14)	0.0483 (8)
C2	0.43742 (17)	0.2955 (4)	0.52959 (13)	0.0434 (8)
C3	0.51208 (17)	0.2536 (4)	0.51198 (14)	0.0450 (8)
C4	0.58718 (18)	0.2419 (5)	0.54600 (16)	0.0591 (10)
H4A	0.598100	0.265759	0.590895	0.071*
C5	0.64493 (19)	0.1937 (5)	0.51096 (18)	0.0653 (10)
H5A	0.695811	0.185900	0.532762	0.078*
C6	0.6297 (2)	0.1564 (5)	0.44427 (17)	0.0615 (10)
H6A	0.670362	0.122386	0.422508	0.074*
C7	0.55574 (19)	0.1688 (5)	0.40953 (15)	0.0521 (9)
C8	0.49847 (17)	0.2183 (4)	0.44538 (14)	0.0466 (8)
C9	0.5936 (2)	0.0804 (6)	0.30818 (17)	0.0778 (12)
H9A	0.571774	0.063916	0.262729	0.117*
H9B	0.615969	-0.033713	0.325700	0.117*
H9C	0.633113	0.173711	0.311913	0.117*
C10	0.33773 (18)	0.4270 (4)	0.65966 (14)	0.0467 (8)
C11	0.3957 (2)	0.4326 (5)	0.71354 (14)	0.0594 (10)
H11A	0.445790	0.395598	0.709579	0.071*
C12	0.3796 (2)	0.4926 (5)	0.77285 (16)	0.0671 (11)
H12A	0.419310	0.497293	0.808808	0.081*
C13	0.3062 (2)	0.5458 (5)	0.78029 (16)	0.0640 (10)
H13A	0.295701	0.585028	0.820967	0.077*
C14	0.2482 (2)	0.5402 (5)	0.72659 (16)	0.0604 (10)
H14A	0.198194	0.577300	0.730814	0.073*
C15	0.26373 (19)	0.4799 (5)	0.66640 (15)	0.0538 (9)
H15A	0.224071	0.475136	0.630425	0.065*
H1	0.313 (2)	0.352 (5)	0.5627 (17)	0.082 (13)*

Atomic displacement parameters (Å²)

	U^{11}	U^{22}	U^{33}	U^{12}	U^{13}	U^{23}
O1	0.0484 (13)	0.0750 (18)	0.0406 (11)	-0.0036 (12)	0.0072 (10)	-0.0031 (11)
O2	0.0484 (15)	0.092 (2)	0.0533 (14)	-0.0061 (13)	0.0049 (11)	-0.0059 (12)
O3	0.0659 (16)	0.103 (2)	0.0527 (14)	0.0041 (14)	0.0200 (12)	-0.0074 (14)
N1	0.0488 (16)	0.0505 (18)	0.0441 (14)	-0.0068 (13)	0.0082 (12)	0.0010 (12)
N2	0.0471 (17)	0.060 (2)	0.0427 (15)	-0.0017 (14)	0.0062 (13)	-0.0044 (13)
C1	0.049 (2)	0.054 (2)	0.0428 (17)	-0.0112 (17)	0.0118 (15)	0.0010 (15)
C2	0.0476 (18)	0.044 (2)	0.0375 (16)	-0.0077 (15)	0.0050 (14)	0.0020 (14)
C3	0.0498 (19)	0.042 (2)	0.0431 (16)	-0.0085 (15)	0.0082 (14)	0.0052 (14)
C4	0.053 (2)	0.073 (3)	0.0489 (18)	-0.0091 (18)	0.0014 (16)	0.0038 (17)
C5	0.044 (2)	0.083 (3)	0.068 (2)	-0.0021 (18)	0.0070 (18)	0.012 (2)
C6	0.054 (2)	0.067 (3)	0.067 (2)	-0.0014 (18)	0.0194 (18)	0.0100 (19)
C7	0.058 (2)	0.051 (2)	0.0499 (19)	-0.0049 (17)	0.0159 (17)	0.0025 (16)
C8	0.0481 (19)	0.047 (2)	0.0447 (17)	-0.0073 (15)	0.0070 (15)	0.0063 (15)
C9	0.092 (3)	0.084 (3)	0.066 (2)	0.011 (2)	0.038 (2)	-0.006 (2)
C10	0.052 (2)	0.047 (2)	0.0404 (16)	-0.0068 (15)	0.0080 (15)	-0.0010 (15)
C11	0.055 (2)	0.076 (3)	0.0456 (18)	0.0031 (18)	0.0016 (16)	-0.0101 (17)
C12	0.074 (3)	0.080 (3)	0.0446 (19)	0.003 (2)	0.0019 (18)	-0.0106 (18)
C13	0.078 (3)	0.069 (3)	0.047 (2)	-0.002 (2)	0.0174 (19)	-0.0081 (17)
C14	0.060 (2)	0.061 (3)	0.065 (2)	0.0041 (18)	0.0248 (19)	0.0021 (18)
C15	0.050 (2)	0.060 (2)	0.0511 (19)	-0.0028 (17)	0.0061 (16)	0.0037 (16)

Geometric parameters (Å, °)

O1—C1	1.380 (3)	C6—C7	1.375 (5)
O1—C8	1.400 (3)	C6—H6A	0.9300
O2—C1	1.205 (3)	C7—C8	1.381 (4)
O3—C7	1.359 (4)	C9—H9A	0.9600
O3—C9	1.431 (4)	C9—H9B	0.9600
N1—C2	1.304 (3)	C9—H9C	0.9600
N1—N2	1.320 (3)	C10—C15	1.374 (4)
N2—C10	1.404 (4)	C10—C11	1.377 (4)
N2—H1	0.92 (4)	C11—C12	1.367 (4)
C1—C2	1.457 (4)	C11—H11A	0.9300
C2—C3	1.438 (4)	C12—C13	1.369 (5)
C3—C8	1.378 (4)	C12—H12A	0.9300
C3—C4	1.386 (4)	C13—C14	1.375 (5)
C4—C5	1.374 (4)	C13—H13A	0.9300
C4—H4A	0.9300	C14—C15	1.381 (4)
C5—C6	1.383 (5)	C14—H14A	0.9300
C5—H5A	0.9300	C15—H15A	0.9300
C1—O1—C8	106.9 (2)	C3—C8—C7	123.8 (3)
C7—O3—C9	116.7 (3)	C3—C8—O1	112.3 (3)
C2—N1—N2	119.5 (3)	C7—C8—O1	123.9 (3)
N1—N2—C10	119.5 (3)	O3—C9—H9A	109.5

N1—N2—H1	118 (2)	O3—C9—H9B	109.5
C10—N2—H1	123 (2)	H9A—C9—H9B	109.5
O2—C1—O1	121.0 (3)	O3—C9—H9C	109.5
O2—C1—C2	130.6 (3)	H9A—C9—H9C	109.5
O1—C1—C2	108.4 (3)	H9B—C9—H9C	109.5
N1—C2—C3	126.1 (3)	C15—C10—C11	119.4 (3)
N1—C2—C1	127.2 (3)	C15—C10—N2	118.6 (3)
C3—C2—C1	106.7 (2)	C11—C10—N2	122.1 (3)
C8—C3—C4	119.4 (3)	C12—C11—C10	120.1 (3)
C8—C3—C2	105.7 (3)	C12—C11—H11A	120.0
C4—C3—C2	134.9 (3)	C10—C11—H11A	120.0
C5—C4—C3	117.6 (3)	C11—C12—C13	121.2 (3)
C5—C4—H4A	121.2	C11—C12—H12A	119.4
C3—C4—H4A	121.2	C13—C12—H12A	119.4
C4—C5—C6	122.1 (3)	C12—C13—C14	118.8 (3)
C4—C5—H5A	119.0	C12—C13—H13A	120.6
C6—C5—H5A	119.0	C14—C13—H13A	120.6
C7—C6—C5	121.3 (3)	C13—C14—C15	120.5 (3)
C7—C6—H6A	119.4	C13—C14—H14A	119.8
C5—C6—H6A	119.4	C15—C14—H14A	119.8
O3—C7—C6	126.6 (3)	C10—C15—C14	120.0 (3)
O3—C7—C8	117.5 (3)	C10—C15—H15A	120.0
C6—C7—C8	115.9 (3)	C14—C15—H15A	120.0
C2—N1—N2—C10	-176.6 (3)	C4—C3—C8—C7	0.8 (5)
C8—O1—C1—O2	-179.2 (3)	C2—C3—C8—C7	-178.7 (3)
C8—O1—C1—C2	0.8 (3)	C4—C3—C8—O1	-179.3 (3)
N2—N1—C2—C3	-178.2 (3)	C2—C3—C8—O1	1.2 (4)
N2—N1—C2—C1	3.7 (5)	O3—C7—C8—C3	179.2 (3)
O2—C1—C2—N1	-1.8 (6)	C6—C7—C8—C3	-0.3 (5)
O1—C1—C2—N1	178.3 (3)	O3—C7—C8—O1	-0.6 (5)
O2—C1—C2—C3	179.9 (4)	C6—C7—C8—O1	179.8 (3)
O1—C1—C2—C3	-0.1 (3)	C1—O1—C8—C3	-1.3 (3)
N1—C2—C3—C8	-179.0 (3)	C1—O1—C8—C7	178.6 (3)
C1—C2—C3—C8	-0.7 (3)	N1—N2—C10—C15	172.0 (3)
N1—C2—C3—C4	1.5 (6)	N1—N2—C10—C11	-7.6 (5)
C1—C2—C3—C4	179.9 (4)	C15—C10—C11—C12	-0.8 (5)
C8—C3—C4—C5	-0.4 (5)	N2—C10—C11—C12	178.7 (3)
C2—C3—C4—C5	178.9 (4)	C10—C11—C12—C13	0.8 (6)
C3—C4—C5—C6	-0.5 (6)	C11—C12—C13—C14	-0.7 (6)
C4—C5—C6—C7	1.0 (6)	C12—C13—C14—C15	0.7 (5)
C9—O3—C7—C6	0.9 (5)	C11—C10—C15—C14	0.8 (5)
C9—O3—C7—C8	-178.5 (3)	N2—C10—C15—C14	-178.8 (3)
C5—C6—C7—O3	179.9 (3)	C13—C14—C15—C10	-0.8 (5)
C5—C6—C7—C8	-0.6 (5)		

Hydrogen-bond geometry (Å, °)

Cg3 is the centroid of the C10–C15 phenyl ring.

<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>
N2—H1 \cdots O2	0.92 (4)	2.14 (3)	2.843 (3)	133 (3)
N2—H1 \cdots O2 ⁱ	0.92 (4)	2.44 (4)	3.181 (4)	138 (3)
C9—H9C \cdots Cg3 ⁱⁱ	0.96	2.70	3.555 (4)	149

Symmetry codes: (i) $-x+1/2, -y+1/2, -z+1$; (ii) $-x+1, -y+1, -z+1$.*Percentage contributions of interatomic contacts to the Hirshfeld surface for the title compound*

Contact	Percentage contribution
H \cdots H	40.7
O \cdots H/H \cdots O	24.7
C \cdots H/H \cdots C	16.1
C \cdots C	8.8
N \cdots C/C \cdots N	3.8
N \cdots H/H \cdots N	3.5
O \cdots C/C \cdots O	1.9
O \cdots N/N \cdots O	0.4
O \cdots O	0.2