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## Structure Reports

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# (E)-5-[(2-Hydroxy-5-methoxybenzylidene)amino]-1,3,4-thiadiazole-2(3H)-thione

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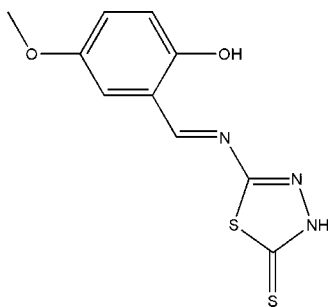
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 Key indicators: single-crystal X-ray study;  $T = 291$  K; mean  $\sigma(\text{C}-\text{C}) = 0.002$  Å;  $R$  factor = 0.040;  $wR$  factor = 0.102; data-to-parameter ratio = 18.7.

In the title thione–Schiff base compound,  $\text{C}_{10}\text{H}_9\text{N}_3\text{O}_2\text{S}_2$ , the dihedral angle between the benzene ring and the five-membered ring is  $6.69$  ( $8^\circ$ ). An intramolecular  $\text{O}-\text{H}\cdots\text{N}$  hydrogen bond forms an  $S_2^2(6)$  ring. In the crystal, inversion dimers linked by pairs of  $\text{N}-\text{H}\cdots\text{S}$  interactions occur, generating  $R_2^2(8)$  ring motifs. The crystal structure features a  $\text{S}\cdots\text{S}$  contact [ $3.3776$  ( $7$ ) Å], which is significantly shorter than the sum of the van der Waals radii ( $3.7$  Å). The crystal structure also features  $\text{C}-\text{H}\cdots\text{O}$  and  $\pi-\pi$  interactions [centroid–centroid distances =  $3.4636$  ( $9$ )– $3.808$  ( $1$ ) Å].

## Related literature

For standard values of bond lengths, see: Allen *et al.* (1987). For hydrogen-bond motifs, see: Bernstein *et al.* (1995). For the biological versatility of thione ligands see, for example: Kumar *et al.* (1988); Yadav *et al.* (1989). For a related structure, see: Zhang (2003). For van der Waals radii, see: Bondi, (1964).



## Experimental

## Crystal data

$\text{C}_{10}\text{H}_9\text{N}_3\text{O}_2\text{S}_2$   
 $M_r = 267.32$   
 Triclinic,  $P\bar{1}$   
 $a = 6.2266$  ( $2$ ) Å  
 $b = 8.0680$  ( $2$ ) Å  
 $c = 11.9695$  ( $3$ ) Å  
 $\alpha = 83.027$  ( $2$ )°  
 $\beta = 77.993$  ( $1$ )°  
 $\gamma = 87.898$  ( $1$ )°  
 $V = 583.76$  ( $3$ ) Å<sup>3</sup>  
 $Z = 2$   
 Mo  $K\alpha$  radiation  
 $\mu = 0.45$  mm<sup>-1</sup>  
 $T = 291$  K  
 $0.11 \times 0.08 \times 0.05$  mm

## Data collection

Bruker SMART APEXII CCD area-detector diffractometer  
 Absorption correction: multi-scan (SADABS; Bruker, 2005)  
 $T_{\min} = 0.952$ ,  $T_{\max} = 0.978$   
 10303 measured reflections  
 2894 independent reflections  
 1990 reflections with  $I > 2\sigma(I)$   
 $R_{\text{int}} = 0.030$

## Refinement

$R[F^2 > 2\sigma(F^2)] = 0.040$   
 $wR(F^2) = 0.102$   
 $S = 1.02$   
 2894 reflections  
 155 parameters  
 H-atom parameters constrained  
 $\Delta\rho_{\text{max}} = 0.30$  e Å<sup>-3</sup>  
 $\Delta\rho_{\text{min}} = -0.25$  e Å<sup>-3</sup>

Table 1

Hydrogen-bond geometry (Å, °).

$D-\text{H}\cdots A$	$D-\text{H}$	$\text{H}\cdots A$	$D\cdots A$	$D-\text{H}\cdots A$
$\text{O1}-\text{H1}\cdots\text{N1}$	0.85	1.84	2.616 (2)	151
$\text{N3}-\text{H3}\cdots\text{S2}^i$	0.80	2.53	3.3163 (15)	169
$\text{C2}-\text{H2A}\cdots\text{O1}^{ii}$	0.93	2.57	3.481 (2)	167
$\text{C3}-\text{H3A}\cdots\text{O2}^{iii}$	0.93	2.52	3.442 (3)	172

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

Data collection: APEX2 (Bruker, 2005); cell refinement: SAINT (Bruker, 2005); data reduction: SAINT; 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: SHELXTL and PLATON (Spek, 2009).

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Supplementary data and figures for this paper are available from the IUCr electronic archives (Reference: FF2041).

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## supporting information

*Acta Cryst.* (2011). E67, o3311 [https://doi.org/10.1107/S1600536811047362]

**(E)-5-[(2-Hydroxy-5-methoxybenzylidene)amino]-1,3,4-thiadiazole-2(3H)-thione****Hadi Kargar, Reza Kia and Muhammad Nawaz Tahir****S1. Comment**

Compounds incorporating a thiadiazole ring have attracted much attention due to their biological activity (Kumar *et al.*, 1988; Yadav *et al.*, 1989). Here we report the crystal structure of a new Schiff base compound containing a thiadiazol ring system.

The asymmetric unit of the title compound, Fig. 1, comprises a thione-Schiff base ligand. The bond lengths (Allen *et al.*, 1987) and angles are within the normal ranges and are comparable to the related structure (Zhang, 2003).

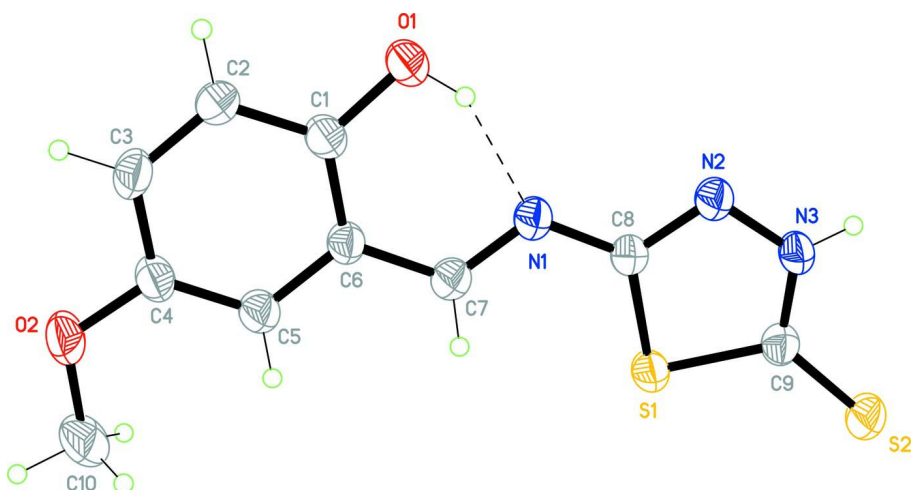
The dihedral angle between the benzene ring and the five-membered ring is 6.69 (8)°. An intramolecular O—H···N hydrogen bond makes S<sub>2</sub><sup>2</sup>(6) ring motif. Intermolecular N—H···S interactions link neighboring molecules into individual dimers with R<sub>2</sub><sup>2</sup>(8) ring motifs (Bernstein *et al.*, 1995). The interesting feature of the crystal structure is a short S(1)···S(1)<sup>i</sup> [3.3776 (7)Å; (i) 1 - x, 1 - y, 1 - z] contact which is significantly shorter than the sum of the Van der Waals radius of S atoms (Bondi, 1964). The crystal structure is stabilized by the intermolecular C—H···O, and  $\pi$ - $\pi$  interactions [Cg1···Cg1<sup>iv</sup> = 3.4636 (9)Å, (iv) -1 - x, -y, 1 - z; Cg1···Cg2<sup>v</sup> = 3.5242 (10)Å, (v) 1 + x, y, z; Cg2···Cg2<sup>vi</sup> = 3.808 (1)Å, (vi) -x, 1 - y, -z Cg1 and Cg2 are centroids of S(1)/C(8)/N(2)/N(3)/C(9) and C1–C6 rings, respectively.

**S2. Experimental**

The title compound was synthesized by adding 5-methoxy-salicylaldehyde (1 mmol) to a solution of 5-aminothiophene-2-thiol (1 mmol) in ethanol (30 ml). The mixture was refluxed with stirring for half an hour. The resultant solution was filtered. Yellow single crystals of the title compound suitable for X-ray structure determination were recrystallized from ethanol by slow evaporation of the solvents at room temperature over several days.

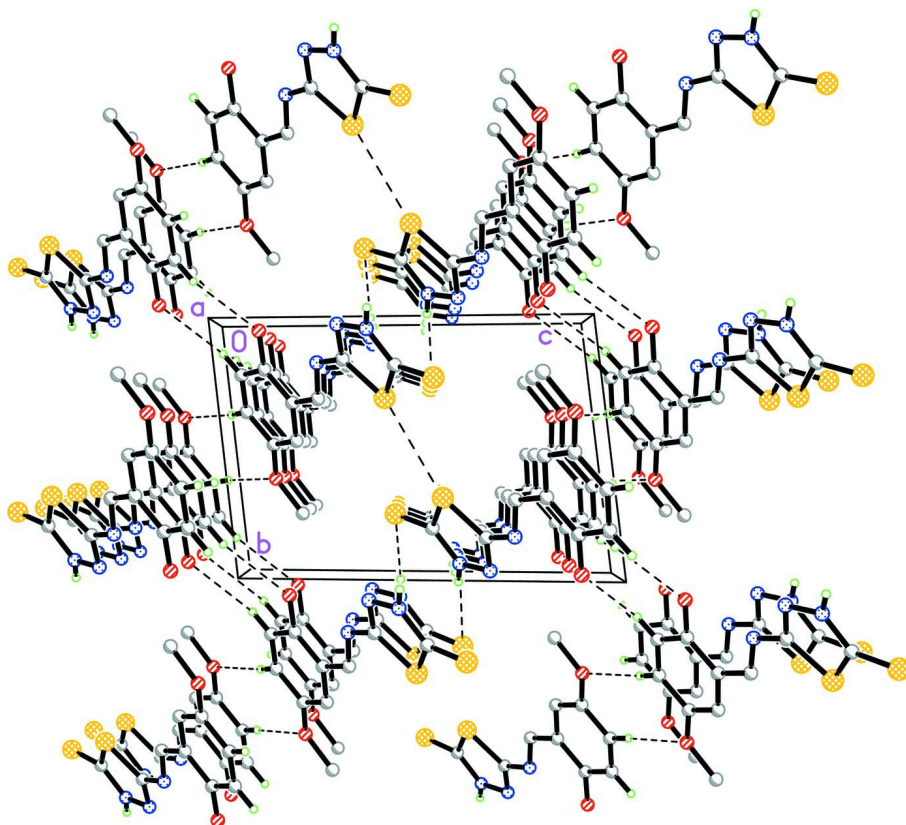
**S3. Refinement**

All hydrogen atoms were positioned geometrically with C—H = 0.93–0.96 Å and included in a riding model approximation with U<sub>iso</sub> (H) = 1.2 or 1.5 U<sub>eq</sub> (C). A rotating group model was applied to the methyl group.



**Figure 1**

ORTEP plot of the title compound, showing 40% probability displacement ellipsoids. The intramolecular hydrogen bond is drawn as dashed line.



**Figure 2**

Packing diagram of the title compound viewed down the *a*-axis. Hydrogen bonds and S...S contacts are drawn as dashed lines.

*(E)*-5-[(2-Hydroxy-5-methoxybenzylidene)amino]-1,3,4-thiadiazole- 2(3*H*)-thione*Crystal data*C<sub>10</sub>H<sub>9</sub>N<sub>3</sub>O<sub>2</sub>S<sub>2</sub> $M_r = 267.32$ Triclinic,  $P\bar{1}$ 

Hall symbol: -P 1

 $a = 6.2266$  (2) Å $b = 8.0680$  (2) Å $c = 11.9695$  (3) Å $\alpha = 83.027$  (2)° $\beta = 77.993$  (1)° $\gamma = 87.898$  (1)° $V = 583.76$  (3) Å<sup>3</sup> $Z = 2$  $F(000) = 276$  $D_x = 1.521$  Mg m<sup>-3</sup>Mo  $K\alpha$  radiation,  $\lambda = 0.71073$  Å

Cell parameters from 2525 reflections

 $\theta = 2.5$ – $27.4$ ° $\mu = 0.45$  mm<sup>-1</sup> $T = 291$  K

Block, yellow

 $0.11 \times 0.08 \times 0.05$  mm*Data collection*Bruker SMART APEXII CCD area-detector  
diffractometer

Radiation source: fine-focus sealed tube

Graphite monochromator

 $\varphi$  and  $\omega$  scans

Absorption correction: multi-scan

(SADABS; Bruker, 2005)

 $T_{\min} = 0.952$ ,  $T_{\max} = 0.978$ 

10303 measured reflections

2894 independent reflections

1990 reflections with  $I > 2\sigma(I)$  $R_{\text{int}} = 0.030$  $\theta_{\text{max}} = 28.4$ °,  $\theta_{\text{min}} = 2.9$ ° $h = -8 \rightarrow 5$  $k = -10 \rightarrow 10$  $l = -15 \rightarrow 15$ *Refinement*Refinement on  $F^2$ 

Least-squares matrix: full

 $R[F^2 > 2\sigma(F^2)] = 0.040$  $wR(F^2) = 0.102$  $S = 1.02$ 

2894 reflections

155 parameters

0 restraints

Primary atom site location: structure-invariant

direct methods

Secondary atom site location: difference Fourier  
mapHydrogen site location: inferred from  
neighbouring sites

H-atom parameters constrained

 $w = 1/[\sigma^2(F_o^2) + (0.0525P)^2]$ where  $P = (F_o^2 + 2F_c^2)/3$  $(\Delta/\sigma)_{\text{max}} = 0.001$  $\Delta\rho_{\text{max}} = 0.30$  e Å<sup>-3</sup> $\Delta\rho_{\text{min}} = -0.25$  e Å<sup>-3</sup>*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  $F^2 > 2\sigma(F^2)$  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<sup>2</sup>)*

	<i>x</i>	<i>y</i>	<i>z</i>	$U_{\text{iso}}^*/U_{\text{eq}}$
S1	0.51725 (8)	0.31804 (6)	0.44207 (4)	0.04448 (16)
S2	0.89337 (8)	0.25182 (7)	0.56684 (5)	0.05370 (18)
O1	0.1438 (3)	0.05675 (17)	0.13460 (13)	0.0639 (4)

H1	0.2279	0.0646	0.1813	0.096*
O2	-0.3796 (3)	0.62067 (18)	0.12072 (13)	0.0651 (4)
N1	0.3359 (2)	0.18322 (18)	0.27908 (12)	0.0401 (4)
N2	0.6290 (2)	0.05119 (18)	0.34526 (12)	0.0409 (4)
N3	0.7659 (2)	0.07198 (18)	0.41793 (12)	0.0402 (4)
H3	0.8576	0.0018	0.4258	0.048*
C1	0.0238 (3)	0.2001 (2)	0.13354 (15)	0.0421 (4)
C2	-0.1342 (3)	0.2186 (2)	0.06674 (16)	0.0457 (5)
H2A	-0.1529	0.1347	0.0229	0.055*
C3	-0.2630 (3)	0.3596 (2)	0.06470 (15)	0.0449 (5)
H3A	-0.3691	0.3701	0.0198	0.054*
C4	-0.2373 (3)	0.4871 (2)	0.12883 (15)	0.0418 (4)
C5	-0.0803 (3)	0.4727 (2)	0.19435 (14)	0.0404 (4)
H5	-0.0620	0.5587	0.2366	0.049*
C6	0.0536 (3)	0.3286 (2)	0.19832 (14)	0.0352 (4)
C7	0.2144 (3)	0.3154 (2)	0.26892 (14)	0.0385 (4)
H7A	0.2320	0.4045	0.3086	0.046*
C8	0.4892 (3)	0.1745 (2)	0.34711 (14)	0.0364 (4)
C9	0.7411 (3)	0.2037 (2)	0.47685 (15)	0.0391 (4)
C10	-0.3578 (4)	0.7562 (3)	0.1826 (2)	0.0667 (6)
H10A	-0.3879	0.7189	0.2635	0.100*
H10B	-0.4599	0.8434	0.1666	0.100*
H10C	-0.2109	0.7979	0.1596	0.100*

*Atomic displacement parameters (Å<sup>2</sup>)*

	$U^{11}$	$U^{22}$	$U^{33}$	$U^{12}$	$U^{13}$	$U^{23}$
S1	0.0469 (3)	0.0408 (3)	0.0532 (3)	0.0106 (2)	-0.0243 (2)	-0.0142 (2)
S2	0.0522 (3)	0.0568 (3)	0.0638 (3)	0.0094 (2)	-0.0324 (3)	-0.0208 (3)
O1	0.0781 (10)	0.0504 (9)	0.0822 (10)	0.0294 (8)	-0.0512 (8)	-0.0319 (8)
O2	0.0684 (10)	0.0569 (9)	0.0849 (11)	0.0316 (7)	-0.0458 (8)	-0.0255 (8)
N1	0.0404 (8)	0.0411 (9)	0.0433 (8)	0.0049 (7)	-0.0180 (7)	-0.0081 (7)
N2	0.0412 (8)	0.0420 (9)	0.0451 (8)	0.0065 (7)	-0.0191 (7)	-0.0108 (7)
N3	0.0392 (8)	0.0391 (8)	0.0469 (8)	0.0090 (6)	-0.0198 (7)	-0.0073 (7)
C1	0.0491 (11)	0.0382 (10)	0.0424 (10)	0.0097 (8)	-0.0166 (8)	-0.0085 (8)
C2	0.0554 (12)	0.0430 (11)	0.0461 (10)	0.0054 (9)	-0.0239 (9)	-0.0134 (9)
C3	0.0453 (10)	0.0530 (12)	0.0418 (10)	0.0037 (9)	-0.0218 (8)	-0.0057 (9)
C4	0.0440 (10)	0.0409 (10)	0.0422 (10)	0.0109 (8)	-0.0139 (8)	-0.0062 (8)
C5	0.0442 (10)	0.0392 (10)	0.0406 (9)	0.0063 (8)	-0.0137 (8)	-0.0091 (8)
C6	0.0361 (9)	0.0372 (9)	0.0342 (8)	0.0036 (7)	-0.0119 (7)	-0.0049 (7)
C7	0.0395 (10)	0.0398 (10)	0.0385 (9)	0.0012 (8)	-0.0125 (8)	-0.0068 (8)
C8	0.0361 (9)	0.0357 (9)	0.0397 (9)	0.0022 (8)	-0.0135 (8)	-0.0039 (8)
C9	0.0367 (9)	0.0401 (10)	0.0415 (9)	0.0011 (8)	-0.0119 (8)	-0.0028 (8)
C10	0.0740 (15)	0.0496 (13)	0.0833 (16)	0.0266 (11)	-0.0277 (13)	-0.0228 (12)

*Geometric parameters (Å, °)*

S1—C9	1.7383 (18)	C1—C6	1.407 (2)
S1—C8	1.7550 (17)	C2—C3	1.368 (3)
S2—C9	1.6624 (19)	C2—H2A	0.9300
O1—C1	1.354 (2)	C3—C4	1.389 (3)
O1—H1	0.8513	C3—H3A	0.9300
O2—C4	1.377 (2)	C4—C5	1.368 (2)
O2—C10	1.418 (3)	C5—C6	1.408 (2)
N1—C7	1.293 (2)	C5—H5	0.9300
N1—C8	1.373 (2)	C6—C7	1.432 (2)
N2—C8	1.297 (2)	C7—H7A	0.9300
N2—N3	1.366 (2)	C10—H10A	0.9600
N3—C9	1.332 (2)	C10—H10B	0.9600
N3—H3	0.8004	C10—H10C	0.9600
C1—C2	1.385 (3)		
C9—S1—C8	89.57 (8)	C4—C5—H5	119.8
C1—O1—H1	105.8	C6—C5—H5	119.8
C4—O2—C10	117.48 (15)	C1—C6—C5	118.89 (16)
C7—N1—C8	120.57 (15)	C1—C6—C7	121.70 (16)
C8—N2—N3	109.05 (15)	C5—C6—C7	119.40 (16)
C9—N3—N2	120.21 (15)	N1—C7—C6	121.91 (17)
C9—N3—H3	121.3	N1—C7—H7A	119.0
N2—N3—H3	118.5	C6—C7—H7A	119.0
O1—C1—C2	118.24 (16)	N2—C8—N1	120.02 (16)
O1—C1—C6	122.23 (17)	N2—C8—S1	114.13 (13)
C2—C1—C6	119.53 (17)	N1—C8—S1	125.85 (13)
C3—C2—C1	120.48 (17)	N3—C9—S2	127.15 (14)
C3—C2—H2A	119.8	N3—C9—S1	107.00 (13)
C1—C2—H2A	119.8	S2—C9—S1	125.86 (11)
C2—C3—C4	120.82 (17)	O2—C10—H10A	109.5
C2—C3—H3A	119.6	O2—C10—H10B	109.5
C4—C3—H3A	119.6	H10A—C10—H10B	109.5
C5—C4—O2	125.16 (16)	O2—C10—H10C	109.5
C5—C4—C3	119.82 (17)	H10A—C10—H10C	109.5
O2—C4—C3	115.02 (16)	H10B—C10—H10C	109.5
C4—C5—C6	120.46 (17)		
C8—N2—N3—C9	0.3 (2)	C4—C5—C6—C7	-179.03 (16)
O1—C1—C2—C3	178.58 (18)	C8—N1—C7—C6	179.75 (15)
C6—C1—C2—C3	-0.9 (3)	C1—C6—C7—N1	-1.9 (3)
C1—C2—C3—C4	0.4 (3)	C5—C6—C7—N1	177.34 (16)
C10—O2—C4—C5	1.9 (3)	N3—N2—C8—N1	178.84 (14)
C10—O2—C4—C3	-178.74 (19)	N3—N2—C8—S1	-1.70 (19)
C2—C3—C4—C5	0.5 (3)	C7—N1—C8—N2	-171.14 (17)
C2—C3—C4—O2	-178.95 (18)	C7—N1—C8—S1	9.5 (2)
O2—C4—C5—C6	178.60 (17)	C9—S1—C8—N2	2.05 (14)

C3—C4—C5—C6	-0.8 (3)	C9—S1—C8—N1	-178.53 (16)
O1—C1—C6—C5	-178.85 (17)	N2—N3—C9—S2	-178.78 (13)
C2—C1—C6—C5	0.6 (3)	N2—N3—C9—S1	1.2 (2)
O1—C1—C6—C7	0.4 (3)	C8—S1—C9—N3	-1.69 (13)
C2—C1—C6—C7	179.87 (17)	C8—S1—C9—S2	178.29 (13)
C4—C5—C6—C1	0.2 (3)		

*Hydrogen-bond geometry (Å, °)*

<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>
O1—H1...N1	0.85	1.84	2.616 (2)	151
N3—H3...S2 <sup>i</sup>	0.80	2.53	3.3163 (15)	169
C2—H2 <i>A</i> ...O1 <sup>ii</sup>	0.93	2.57	3.481 (2)	167
C3—H3 <i>A</i> ...O2 <sup>iii</sup>	0.93	2.52	3.442 (3)	172

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