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## Structure Reports

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## 2-(3-Oxo-3,4-dihydro-2H-1,4-benzothiazin-4-yl)acetohydrazide

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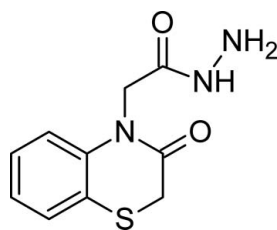
Received 3 August 2010; accepted 4 August 2010

Key indicators: single-crystal X-ray study;  $T = 296$  K; mean  $\sigma(\text{C}-\text{C}) = 0.003$  Å;  $R$  factor = 0.027;  $wR$  factor = 0.076; data-to-parameter ratio = 13.8.

In the title compound,  $\text{C}_{10}\text{H}_{11}\text{N}_3\text{O}_2\text{S}$ , the thiazine ring exists in a conformation intermediate between twist-boat and half-chair. The dihedral angle between the mean plane of the thiazine ring and the hydrazide group is  $89.45(13)^\circ$ . In the crystal,  $\text{N}-\text{H}\cdots\text{O}$  hydrogen bonds link the molecules into (100) sheets and weak  $\text{C}-\text{H}\cdots\text{O}$  interactions further consolidate the packing.

## Related literature

For the biological and medicinal activity of 1,4-benzothiazine compounds, see: Armenise *et al.* (1991); Gupta *et al.* (1993); Vicente *et al.* (2009); Schiaffella *et al.* (2006); Kaneko *et al.* (2002). For the pharmacological properties of hydrazones and their derivatives, see: Sivasankar *et al.* (1995); Satyanarayana *et al.* (2008). For hydrogen-bond motifs, see: Bernstein *et al.* (1995).



## Experimental

## Crystal data

$\text{C}_{10}\text{H}_{11}\text{N}_3\text{O}_2\text{S}$   
 $M_r = 237.28$   
 Monoclinic,  $Cc$   
 $a = 15.3744(10)$  Å  
 $b = 7.5162(5)$  Å  
 $c = 9.6256(7)$  Å  
 $\beta = 95.413(3)^\circ$

$V = 1107.35(13)$  Å<sup>3</sup>  
 $Z = 4$   
 Mo  $K\alpha$  radiation  
 $\mu = 0.28$  mm<sup>-1</sup>  
 $T = 296$  K  
 $0.46 \times 0.23 \times 0.20$  mm

## Data collection

Bruker Kappa APEXII CCD diffractometer  
 Absorption correction: multi-scan (SADABS; Bruker, 2007)  
 $T_{\min} = 0.882$ ,  $T_{\max} = 0.946$

6103 measured reflections  
 2168 independent reflections  
 2077 reflections with  $I > 2\sigma(I)$   
 $R_{\text{int}} = 0.021$

## Refinement

$R[F^2 > 2\sigma(F^2)] = 0.027$   
 $wR(F^2) = 0.076$   
 $S = 1.04$   
 2168 reflections  
 157 parameters  
 2 restraints

H atoms treated by a mixture of independent and constrained refinement  
 $\Delta\rho_{\max} = 0.18$  e Å<sup>-3</sup>  
 $\Delta\rho_{\min} = -0.14$  e Å<sup>-3</sup>  
 Absolute structure: Flack (1983), 792 Friedel pairs  
 Flack parameter: 0.00 (7)

Table 1

Hydrogen-bond geometry (Å, °).

$D-\text{H}\cdots A$	$D-\text{H}$	$\text{H}\cdots A$	$D\cdots A$	$D-\text{H}\cdots A$
$\text{N3}-\text{H3N}\cdots\text{O2}^{\text{i}}$	0.96 (3)	2.35 (3)	3.299 (3)	171 (2)
$\text{N2}-\text{H1N}\cdots\text{O1}^{\text{ii}}$	0.87 (2)	2.07 (2)	2.935 (2)	175.5 (18)
$\text{C3}-\text{H3}\cdots\text{O2}^{\text{iii}}$	0.93	2.48	3.406 (3)	174
$\text{C8}-\text{H8B}\cdots\text{O1}^{\text{iv}}$	0.97	2.57	3.442 (2)	150

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

Data collection: APEX2 (Bruker, 2007); cell refinement: SAINT (Bruker, 2007); data reduction: SAINT; program(s) used to solve structure: SHELXS97 (Sheldrick, 2008); program(s) used to refine structure: SHELXL97 (Sheldrick, 2008); molecular graphics: ORTEP-3 (Farrugia, 1997) and PLATON (Spek, 2009); software used to prepare material for publication: WinGX (Farrugia, 1999) and PLATON.

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Supplementary data and figures for this paper are available from the IUCr electronic archives (Reference: HB5598).

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**supplementary materials**

*Acta Cryst.* (2010). E66, o2289-o2290 [ doi:10.1107/S1600536810031272 ]

## 2-(3-Oxo-3,4-dihydro-2H-1,4-benzothiazin-4-yl)acetohydrazide

A. Saeed, Z. Mahmood, S. Yang, S. Ahmad and M. Salim

### Comment

The 4H-benzo(1,4)thiazine compounds exhibit a broad spectrum of biological activities, such as tetramic acids substituted benzothiazine derivatives are potent inhibitors of HCV polymerase (Vicente et al., 2009) and the pyrazino substituted 1,4-benzothiazine derivatives are inhibitors of adhesion molecule-1, (Kaneko et al., 2002). They are also known to have antibacterial (Armenise et al., 1991), anticancer (Gupta et al., 1993), antifungal (Schiaffella et al., 2006) activities. The hydrazone compounds were known for their coordinating capability, pharmacological activity, antibacterial and antifungal properties (Sivasankar et al., 1995) (Satyanarayana, et al., 2008). We paid attention to the preparation of hydrazone derivatives of 2-(3-oxo-2,3-dihydro-2H-1,4-benzothiazin-3-one and we report here the structure of the title compound.

The dihedral angle between the aromatic benzene ring C1–C6 and thiazine ring C1/C6/N1/C7/C8/S1 is  $16.77(0.10)^\circ$  while the hydrazide group C9/C10/N2/N3 is oriented at dihedral angle of  $89.45(0.13)^\circ$  with respect to the thiazine ring. The symmetry related intermolecular N—H $\cdots$ O and C—H $\cdots$ O interaction form the dimer along the b axis which results in a ring motif  $R_2^2(9)$  (Bernstein et al., 1995). The crystal structure is further stabilized through the N—H $\cdots$ O and weak C—H $\cdots$ O interaction to form three dimensional network.

### Experimental

Ethyl 2-(3-oxo-2,3-dihydrobenzo[b][1,4]thiazin-4-yl)acetate (1.26 g, 5mmol) was refluxed in 50 ml ethanol with 2.0 ml of hydrazine for 5 hours. On completion the solution was evaporated under reduce pressure and solid obtained was purified from ethanol. Colourless needles of (I) were obtained by slow evaporation from methanol (m.p. 430 K).

### Refinement

The C-H H-atoms were positioned with idealized geometry with C—H = 0.93 Å and were refined using a riding model with  $U_{\text{iso}}(\text{H}) = 1.2 U_{\text{eq}}(\text{C})$ . The N-H H atoms were located in difference map with N—H = 0.76 (4)—0.83 (4) Å,  $U_{\text{iso}}(\text{H}) = 1.2$  for N atoms.

### Figures

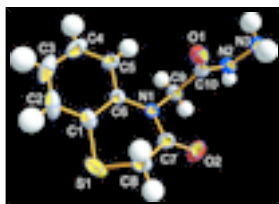


Fig. 1. The molecular structure of the title compound, with displacement ellipsoids drawn at the 50% probability level.

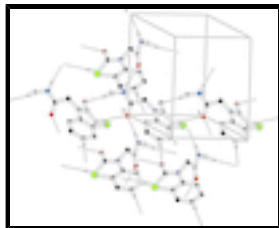


Fig. 2. The crystal packing of the title compound and Intermolecular hydrogen bonds are shown by dashed lines.

## 2-(3-Oxo-3,4-dihydro-2H-1,4-benzothiazin-4-yl)acetohydrazide

### Crystal data

$C_{10}H_{11}N_3O_2S$

$M_r = 237.28$

Monoclinic,  $Cc$

Hall symbol:  $C -2yc$

$a = 15.3744 (10) \text{ \AA}$

$b = 7.5162 (5) \text{ \AA}$

$c = 9.6256 (7) \text{ \AA}$

$\beta = 95.413 (3)^\circ$

$V = 1107.35 (13) \text{ \AA}^3$

$Z = 4$

$F(000) = 496$

$D_x = 1.423 \text{ Mg m}^{-3}$

Melting point: 430 K

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

Cell parameters from 4206 reflections

$\theta = 3.0\text{--}31.2^\circ$

$\mu = 0.28 \text{ mm}^{-1}$

$T = 296 \text{ K}$

Needle, colorless

$0.46 \times 0.23 \times 0.20 \text{ mm}$

### Data collection

Bruker Kappa APEXII CCD  
diffractometer

Radiation source: fine-focus sealed tube  
graphite

$\varphi$  and  $\omega$  scans

Absorption correction: multi-scan  
(*SADABS*; Bruker, 2007)

$T_{\min} = 0.882$ ,  $T_{\max} = 0.946$

6103 measured reflections

2168 independent reflections

2077 reflections with  $I > 2\sigma(I)$

$R_{\text{int}} = 0.021$

$\theta_{\max} = 28.4^\circ$ ,  $\theta_{\min} = 2.7^\circ$

$h = -12 \rightarrow 20$

$k = -8 \rightarrow 10$

$l = -12 \rightarrow 10$

### Refinement

Refinement on  $F^2$

Least-squares matrix: full

$R[F^2 > 2\sigma(F^2)] = 0.027$

$wR(F^2) = 0.076$

$S = 1.04$

2168 reflections

157 parameters

2 restraints

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

$w = 1/[\sigma^2(F_o^2) + (0.0437P)^2 + 0.2667P]$

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

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

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

$\Delta\rho_{\min} = -0.14 \text{ e \AA}^{-3}$

Absolute structure: Flack (1983), 792 Friedel pairs

Primary atom site location: structure-invariant direct methods Flack parameter: 0.00 (7)

*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 ( $\text{\AA}^2$ )*

	x	y	z	$U_{\text{iso}}^*/U_{\text{eq}}$
C1	0.81283 (13)	0.1402 (3)	0.4407 (2)	0.0444 (4)
C2	0.88393 (15)	0.1642 (4)	0.3624 (3)	0.0684 (7)
H2	0.9061	0.0674	0.3170	0.082*
C3	0.92138 (15)	0.3276 (5)	0.3514 (3)	0.0727 (7)
H3	0.9691	0.3412	0.2998	0.087*
C4	0.88855 (14)	0.4717 (4)	0.4166 (3)	0.0621 (6)
H4	0.9145	0.5827	0.4099	0.074*
C5	0.81688 (12)	0.4524 (2)	0.4924 (2)	0.0451 (4)
H5	0.7944	0.5514	0.5347	0.054*
C6	0.77806 (10)	0.2868 (2)	0.50596 (15)	0.0325 (3)
C7	0.64582 (13)	0.1320 (2)	0.56774 (18)	0.0393 (4)
C8	0.65696 (13)	-0.0005 (2)	0.4532 (2)	0.0473 (4)
H8A	0.6398	0.0537	0.3633	0.057*
H8B	0.6194	-0.1024	0.4638	0.057*
C9	0.68864 (12)	0.3985 (2)	0.69169 (16)	0.0371 (4)
H9A	0.7419	0.4638	0.7190	0.044*
H9B	0.6710	0.3388	0.7738	0.044*
C10	0.61760 (11)	0.52828 (19)	0.63748 (15)	0.0311 (3)
N1	0.70588 (9)	0.26545 (17)	0.58681 (13)	0.0328 (3)
N2	0.57596 (11)	0.60513 (19)	0.73773 (16)	0.0391 (3)
N3	0.50598 (12)	0.7263 (3)	0.70912 (19)	0.0491 (4)
O1	0.60169 (9)	0.56431 (16)	0.51324 (12)	0.0441 (3)
O2	0.58443 (11)	0.1213 (2)	0.63853 (16)	0.0602 (4)
S1	0.76839 (4)	-0.07275 (6)	0.45907 (6)	0.06435 (19)
H1N	0.5831 (14)	0.561 (3)	0.822 (3)	0.036 (5)*
H2N	0.4722 (19)	0.678 (4)	0.641 (3)	0.072 (9)*
H3N	0.5321 (18)	0.834 (4)	0.681 (3)	0.065 (8)*

*Atomic displacement parameters ( $\text{\AA}^2$ )*

$U^{11}$                        $U^{22}$                        $U^{33}$                        $U^{12}$                        $U^{13}$                        $U^{23}$

## supplementary materials

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C1	0.0391 (9)	0.0492 (9)	0.0446 (10)	0.0071 (7)	0.0020 (8)	-0.0098 (8)
C2	0.0384 (11)	0.108 (2)	0.0598 (13)	0.0113 (12)	0.0099 (10)	-0.0256 (14)
C3	0.0360 (10)	0.126 (2)	0.0580 (14)	-0.0063 (14)	0.0133 (9)	0.0076 (15)
C4	0.0409 (10)	0.0773 (15)	0.0673 (14)	-0.0139 (10)	0.0019 (10)	0.0272 (12)
C5	0.0411 (10)	0.0407 (9)	0.0531 (12)	-0.0021 (8)	0.0022 (8)	0.0107 (8)
C6	0.0334 (8)	0.0347 (7)	0.0288 (8)	0.0042 (6)	0.0004 (6)	0.0033 (6)
C7	0.0462 (10)	0.0359 (8)	0.0360 (9)	-0.0013 (7)	0.0057 (7)	0.0027 (7)
C8	0.0532 (11)	0.0358 (9)	0.0529 (11)	-0.0081 (7)	0.0049 (8)	-0.0080 (7)
C9	0.0463 (9)	0.0382 (8)	0.0266 (8)	0.0068 (7)	0.0026 (7)	-0.0034 (6)
C10	0.0391 (8)	0.0292 (7)	0.0253 (7)	-0.0012 (6)	0.0052 (6)	-0.0015 (5)
N1	0.0393 (7)	0.0293 (6)	0.0304 (7)	0.0025 (5)	0.0068 (6)	-0.0012 (5)
N2	0.0516 (9)	0.0405 (7)	0.0261 (7)	0.0102 (6)	0.0081 (6)	0.0003 (6)
N3	0.0518 (10)	0.0517 (10)	0.0455 (9)	0.0147 (8)	0.0134 (8)	0.0016 (8)
O1	0.0585 (8)	0.0504 (7)	0.0239 (6)	0.0137 (6)	0.0059 (5)	0.0007 (5)
O2	0.0559 (9)	0.0728 (9)	0.0553 (9)	-0.0161 (7)	0.0223 (7)	-0.0008 (8)
S1	0.0661 (3)	0.0354 (2)	0.0898 (4)	0.0116 (2)	-0.0018 (3)	-0.0191 (3)

### *Geometric parameters (Å, °)*

C1—C2	1.397 (3)	C7—C8	1.508 (3)
C1—C6	1.400 (2)	C8—S1	1.793 (2)
C1—S1	1.756 (2)	C8—H8A	0.9700
C2—C3	1.364 (4)	C8—H8B	0.9700
C2—H2	0.9300	C9—N1	1.463 (2)
C3—C4	1.372 (4)	C9—C10	1.520 (2)
C3—H3	0.9300	C9—H9A	0.9700
C4—C5	1.385 (3)	C9—H9B	0.9700
C4—H4	0.9300	C10—O1	1.228 (2)
C5—C6	1.392 (2)	C10—N2	1.338 (2)
C5—H5	0.9300	N2—N3	1.417 (2)
C6—N1	1.423 (2)	N2—H1N	0.87 (2)
C7—O2	1.218 (2)	N3—H2N	0.87 (3)
C7—N1	1.364 (2)	N3—H3N	0.96 (3)
C2—C1—C6	119.5 (2)	S1—C8—H8A	109.5
C2—C1—S1	120.28 (17)	C7—C8—H8B	109.5
C6—C1—S1	120.24 (15)	S1—C8—H8B	109.5
C3—C2—C1	121.0 (2)	H8A—C8—H8B	108.1
C3—C2—H2	119.5	N1—C9—C10	111.90 (13)
C1—C2—H2	119.5	N1—C9—H9A	109.2
C2—C3—C4	119.9 (2)	C10—C9—H9A	109.2
C2—C3—H3	120.0	N1—C9—H9B	109.2
C4—C3—H3	120.0	C10—C9—H9B	109.2
C3—C4—C5	120.3 (2)	H9A—C9—H9B	107.9
C3—C4—H4	119.9	O1—C10—N2	122.83 (15)
C5—C4—H4	119.9	O1—C10—C9	123.14 (14)
C4—C5—C6	120.8 (2)	N2—C10—C9	113.99 (13)
C4—C5—H5	119.6	C7—N1—C6	124.21 (13)
C6—C5—H5	119.6	C7—N1—C9	115.53 (14)
C5—C6—C1	118.46 (16)	C6—N1—C9	120.12 (13)

C5—C6—N1	121.03 (15)	C10—N2—N3	122.91 (15)
C1—C6—N1	120.49 (15)	C10—N2—H1N	118.4 (13)
O2—C7—N1	121.60 (17)	N3—N2—H1N	117.0 (14)
O2—C7—C8	120.86 (17)	N2—N3—H2N	105.3 (19)
N1—C7—C8	117.53 (15)	N2—N3—H3N	105.6 (16)
C7—C8—S1	110.54 (14)	H2N—N3—H3N	112 (2)
C7—C8—H8A	109.5	C1—S1—C8	95.78 (9)
C6—C1—C2—C3	1.6 (3)	O2—C7—N1—C6	-178.88 (18)
S1—C1—C2—C3	-177.5 (2)	C8—C7—N1—C6	-0.2 (2)
C1—C2—C3—C4	-0.8 (4)	O2—C7—N1—C9	-3.3 (2)
C2—C3—C4—C5	-0.7 (4)	C8—C7—N1—C9	175.45 (16)
C3—C4—C5—C6	1.2 (3)	C5—C6—N1—C7	155.73 (17)
C4—C5—C6—C1	-0.3 (3)	C1—C6—N1—C7	-25.7 (2)
C4—C5—C6—N1	178.29 (17)	C5—C6—N1—C9	-19.7 (2)
C2—C1—C6—C5	-1.1 (3)	C1—C6—N1—C9	158.89 (15)
S1—C1—C6—C5	178.02 (14)	C10—C9—N1—C7	-76.87 (18)
C2—C1—C6—N1	-179.71 (19)	C10—C9—N1—C6	98.95 (17)
S1—C1—C6—N1	-0.6 (2)	O1—C10—N2—N3	4.2 (3)
O2—C7—C8—S1	-135.26 (18)	C9—C10—N2—N3	-178.15 (17)
N1—C7—C8—S1	46.0 (2)	C2—C1—S1—C8	-143.10 (19)
N1—C9—C10—O1	-26.9 (2)	C6—C1—S1—C8	37.80 (16)
N1—C9—C10—N2	155.47 (14)	C7—C8—S1—C1	-58.15 (15)

*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>
N3—H3N $\cdots$ O2 <sup>i</sup>	0.96 (3)	2.35 (3)	3.299 (3)	171 (2)
N2—H1N $\cdots$ O1 <sup>ii</sup>	0.87 (2)	2.07 (2)	2.935 (2)	175.5 (18)
C3—H3 $\cdots$ O2 <sup>iii</sup>	0.93	2.48	3.406 (3)	174
C8—H8B $\cdots$ O1 <sup>iv</sup>	0.97	2.57	3.442 (2)	150

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



Fig. 1

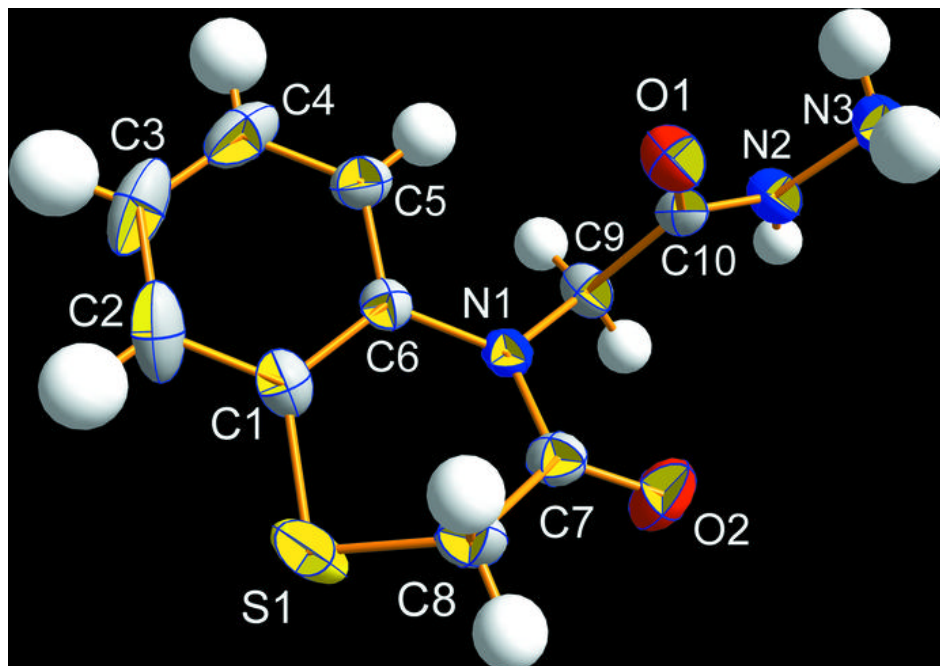


Fig. 2

