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N-(3-Chloropropionyl)-N'-phenylthio-urea

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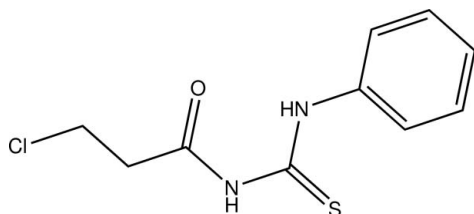
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Key indicators: single-crystal X-ray study; $T = 298$ K; mean $\sigma(\text{C}-\text{C}) = 0.004$ Å; R factor = 0.041; wR factor = 0.113; data-to-parameter ratio = 15.5.

The title compound, $\text{C}_{10}\text{H}_{11}\text{ClN}_2\text{OS}$, adopts a *cis-trans* configuration with respect to the position of the phenyl and 3-chloropropionyl groups relative to the thiono group across the C–N bonds. The benzene ring is perpendicular to the propionyl thiourea fragment with a dihedral angle of 82.62 (10)°. An intramolecular N–H···O interaction occurs. The crystal structure is stabilized by intermolecular N–H···S hydrogen bonds, which link pairs of molecules, building up $R_2^2(8)$ ring motifs, and C–H··· π interactions.

Related literature

For related structures, see: Ismail *et al.* (2007); Ismail & Yamin (2009). For hydrogen-bond motifs, see: Etter *et al.* (1990); Bernstein *et al.* (1995). For reference bond lengths, see: Allen *et al.* (1987).



Experimental

Crystal data

$\text{C}_{10}\text{H}_{11}\text{ClN}_2\text{OS}$
 $M_r = 242.72$
 Triclinic, $P\bar{1}$
 $a = 5.8088$ (12) Å

$b = 10.467$ (2) Å
 $c = 10.660$ (2) Å
 $\alpha = 112.811$ (3)°
 $\beta = 101.855$ (3)°

$\gamma = 95.483$ (3)°
 $V = 573.6$ (2) Å³
 $Z = 2$
 Mo $K\alpha$ radiation

$\mu = 0.49$ mm⁻¹
 $T = 298$ K
 $0.49 \times 0.45 \times 0.27$ mm

Data collection

Bruker SMART APEX CCD area-detector diffractometer
 Absorption correction: multi-scan (SADABS; Bruker, 2000)
 $T_{\min} = 0.795$, $T_{\max} = 0.879$

5617 measured reflections
 2103 independent reflections
 1798 reflections with $I > 2\sigma(I)$
 $R_{\text{int}} = 0.016$

Refinement

$R[F^2 > 2\sigma(F^2)] = 0.041$
 $wR(F^2) = 0.113$
 $S = 1.04$
 2103 reflections

136 parameters
 H-atom parameters constrained
 $\Delta\rho_{\text{max}} = 0.54$ e Å⁻³
 $\Delta\rho_{\text{min}} = -0.43$ e Å⁻³

Table 1

Hydrogen-bond geometry (Å, °).

Cg1 is the centroid of the C5–C10 ring.

$D-H\cdots A$	$D-H$	$H\cdots A$	$D\cdots A$	$D-H\cdots A$
N2–H2A···O1	0.86	2.01	2.677 (3)	134
N1–H1A···S1 ⁱ	0.86	2.53	3.3709 (19)	165
C1–H1C···Cg1 ⁱⁱ	0.97	2.84	3.466	123

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

Data collection: SMART (Bruker, 2000); cell refinement: SAINT (Bruker, 2000); data reduction: SAINT; 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, PARST (Nardelli, 1995) and PLATON (Spek, 2009).

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Supplementary data and figures for this paper are available from the IUCr electronic archives (Reference: DN2535).

References

- Allen, F. H., Kennard, O., Watson, D. G., Brammer, L., Orpen, A. G. & Taylor, R. (1987). *J. Chem. Soc. Perkin Trans. 2*, pp. S1–19.
 Bernstein, J., Davis, R. E., Shimoni, L. & Chang, N.-L. (1995). *Angew. Chem. Int. Ed. Engl.* **34**, 1555–1573.
 Bruker (2000). *SADABS, SMART and SAINT*. Bruker AXS Inc., Madison, Wisconsin, USA.
 Etter, M. C., MacDonald, J. C. & Bernstein, J. (1990). *Acta Cryst.* **B46**, 256–262.
 Ismail, N. L., Othman, E. & Yamin, B. M. (2007). *Acta Cryst.* **E63**, o2442–o2443.
 Ismail, N. & Yamin, B. M. (2009). X-ray Struct. Anal. Online, **25**, 39–40.
 Nardelli, M. (1995). *J. Appl. Cryst.* **28**, 659.
 Sheldrick, G. M. (2008). *Acta Cryst.* **A64**, 112–122.
 Spek, A. L. (2009). *Acta Cryst.* **D65**, 148–155.

supporting information

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***N*-(3-Chloropropionyl)-*N'*-phenylthiourea**

Eliyanti A. Othman, Siti K. C. Soh and Bohari M. Yamin

S1. Comment

The presence of both alpha and ipso chlorine atoms in 2-chloropropionyl chloride could lead to a complicated reaction when reacted with a nucleophile such as thiocyanate depending on the solvent used. For example, the reaction of 2-chloropropionyl chloride with ammonium thiocyanate and succeedingly with aniline in acetone was found to give 4,5,6-trimethyl-1-phenyl-3,4-dihydropyrimidine-2(1*H*)-thione (Ismail *et al.*, 2007) instead of the expected *N*-phenyl-*N'*-(2-chloropropionyl) thiourea. However, in the present study, the same reaction but with 3-chloropropionyl chloride, the expected *N*-phenyl-*N'*-(3-chloropropionyl)thiourea (I) was indeed obtained.

The whole molecule is not planar (Fig.1). The benzene (C5—C10) ring and propionylthiourea fragment, S1/N1/N2/(C1—C4) are each planar with maximum deviation of 0.062 (2)Å for C3 atom from the least square plane. The benzene ring is roughly perpendicular to the propionylthiourea fragment with dihedral angle between the two planes of 82.62 (10)°. A smaller dihedral angle of 12.68 (7)° was observed in an analogous compound of *N*-(3-chloropropionyl)-*N'*-(4-fluorophenyl) thiourea (II) (Ismail & Yamin, 2009). The trans-cis configuration of the propionyl and phenyl groups relative to the thiono group respectively, across their C—N bonds, is maintained. The bond lengths and angles are in normal range (Allen *et al.*, 1987) and comparable to those in (II).

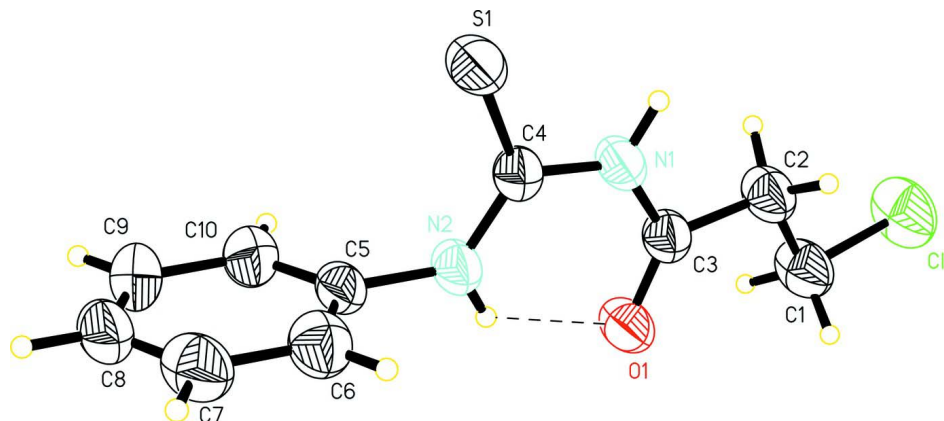
There is one intrahydrogen bond, N2—H2A···O1 forming the pseudo-six membered ring, N2/C4/N1/C3/O1···H2A. In the crystal structure, the molecules are linked by N1—H1A···S1 intermolecular hydrogen bond forming dimers through a R₂²(8) ring (Etter *et al.*, 1990 ; Bernstein *et al.*, 1995) extending along the c-axis (Table 1, Fig.2). In addition, there is also a C1—H1C···π bond with the benzene (C5—C10) ring (Table 1).

S2. Experimental

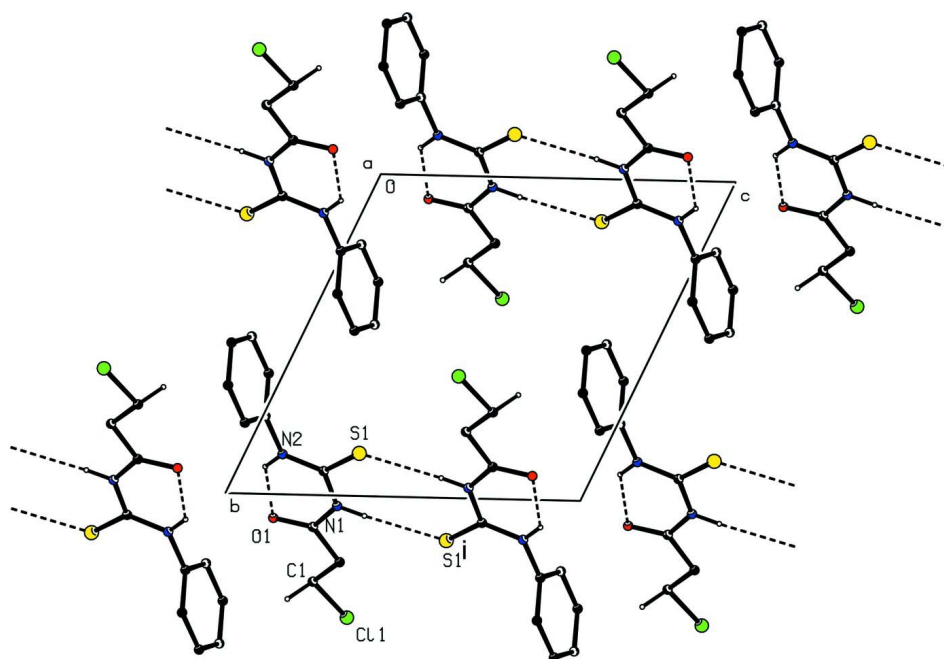
30 ml acetone solution of aniline was added into 30 ml acetone containing 3-chloropropionyl isothiocyanate (1.49, 0.01 mol). The mixture was refluxed for 2 hours. The solution was filtered and left to evaporate at room temperature. The white precipitate obtained after a few days, was washed with water and cold ethanol. The colorless crystals were obtained by recrystallization from ethanol. Yield 90%; m.p 389.7-391.2 K; ¹H NMR (400 MHz, CDCl₃-d₆):δ (ppm)= 12.32 (s,1H), 9.99 (s,1H), 7.63 (m,4H, H_{Ar}), 7.41 (s,1H, H_{Ar}), 3.8 (d,2H,Cl—CH₂), 2.9 (d,2H,CH₂). ¹³C NMR (400 MHz, CDCl₃-d₆):δ (ppm)= 38.32, 39.76, 124.70, 127.35, 127.35, 129.08, 129.08, 137.30, 171.22, 178.49.

S3. Refinement

After locating the hydrogen atoms from a different-Fourier map, they were positioned geometrically with C—H=0.93-0.97Å and N—H=0.86Å respectively, and constrained to all their parent atoms with U_{iso}(H)=1.2U_{eq}(parent atom). There is a highest peak and deepest hole of 0.97 and 0.81 Å respectively from S1 atom.

**Figure 1**

The molecular structure of (I), with displacement ellipsoids drawn at the 50% probability level. H atoms are represented as small spheres of arbitrary radii.

**Figure 2**

A packing diagram of (I) viewed down the *a* axis. Hydrogen bonds are shown by dashed lines. H atoms not involved in hydrogen bondings have been omitted for clarity. [Symmetry code: (i) $-x+2, -y+2, -z+1$]

N-(3-chloropropionyl)-*N'*-phenylthiourea

Crystal data

$C_{10}H_{11}ClN_2OS$

$M_r = 242.72$

Triclinic, *P*1

Hall symbol: $-P\ 1$

$a = 5.8088$ (12) Å

$b = 10.467$ (2) Å

$c = 10.660$ (2) Å

$\alpha = 112.811$ (3)°

$\beta = 101.855$ (3)°

$\gamma = 95.483$ (3)°

$V = 573.6$ (2) Å³

$Z = 2$

$F(000) = 252$

$D_x = 1.405$ Mg m⁻³

Mo $K\alpha$ radiation, $\lambda = 0.71073 \text{ \AA}$
 Cell parameters from 3405 reflections
 $\theta = 2.1\text{--}25.5^\circ$
 $\mu = 0.49 \text{ mm}^{-1}$

$T = 298 \text{ K}$
 Block, colourless
 $0.49 \times 0.45 \times 0.27 \text{ mm}$

Data collection

Bruker SMART APEX CCD area-detector
 diffractometer
 Radiation source: fine-focus sealed tube
 Detector resolution: $83.66 \text{ pixels mm}^{-1}$
 ω scan
 Absorption correction: multi-scan
 (SADABS; Bruker, 2000)
 $T_{\min} = 0.795$, $T_{\max} = 0.879$

5617 measured reflections
 2103 independent reflections
 1798 reflections with $I > 2\sigma(I)$
 $R_{\text{int}} = 0.016$
 $\theta_{\text{max}} = 25.5^\circ$, $\theta_{\text{min}} = 2.1^\circ$
 $h = -7 \rightarrow 7$
 $k = -12 \rightarrow 12$
 $l = -12 \rightarrow 12$

Refinement

Refinement on F^2
 Least-squares matrix: full
 $R[F^2 > 2\sigma(F^2)] = 0.041$
 $wR(F^2) = 0.113$
 $S = 1.04$
 2103 reflections
 136 parameters
 0 restraints
 Primary atom site location: structure-invariant
 direct methods

Secondary atom site location: difference Fourier
 map
 Hydrogen site location: inferred from
 neighbouring sites
 H-atom parameters constrained
 $w = 1/[\sigma^2(F_o^2) + (0.0559P)^2 + 0.2663P]$
 where $P = (F_o^2 + 2F_c^2)/3$
 $(\Delta/\sigma)_{\text{max}} < 0.001$
 $\Delta\rho_{\text{max}} = 0.54 \text{ e \AA}^{-3}$
 $\Delta\rho_{\text{min}} = -0.43 \text{ e \AA}^{-3}$

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	$U_{\text{iso}}^*/U_{\text{eq}}$
Cl1	0.23811 (13)	1.37819 (8)	0.50799 (7)	0.0777 (3)
S1	1.10045 (13)	0.86771 (7)	0.31949 (6)	0.0653 (2)
O1	0.5152 (3)	1.07380 (16)	0.16725 (14)	0.0539 (4)
N1	0.7823 (3)	1.03263 (18)	0.32922 (17)	0.0497 (4)
H1A	0.8413	1.0614	0.4186	0.060*
N2	0.7847 (3)	0.87565 (18)	0.10659 (17)	0.0484 (4)
H2A	0.6781	0.9154	0.0753	0.058*
C1	0.3405 (4)	1.2689 (3)	0.3640 (2)	0.0594 (6)
H1B	0.2091	1.1935	0.2967	0.071*
H1C	0.3931	1.3244	0.3167	0.071*
C2	0.5422 (4)	1.2067 (2)	0.4124 (2)	0.0539 (5)
H2B	0.4968	1.1623	0.4707	0.065*

H2C	0.6809	1.2815	0.4696	0.065*
C3	0.6079 (4)	1.0991 (2)	0.2902 (2)	0.0434 (5)
C4	0.8769 (4)	0.9252 (2)	0.2443 (2)	0.0452 (5)
C5	0.8541 (4)	0.7586 (2)	0.00691 (19)	0.0435 (5)
C6	1.0500 (4)	0.7804 (2)	-0.0412 (2)	0.0543 (5)
H6A	1.1404	0.8707	-0.0087	0.065*
C7	1.1101 (5)	0.6653 (3)	-0.1391 (3)	0.0618 (6)
H7A	1.2421	0.6787	-0.1725	0.074*
C8	0.9790 (5)	0.5329 (3)	-0.1872 (2)	0.0635 (7)
H8A	1.0221	0.4565	-0.2524	0.076*
C9	0.7837 (5)	0.5125 (2)	-0.1393 (3)	0.0646 (7)
H9A	0.6934	0.4221	-0.1724	0.078*
C10	0.7203 (4)	0.6258 (2)	-0.0418 (2)	0.0541 (5)
H10A	0.5872	0.6120	-0.0094	0.065*

Atomic displacement parameters (Å²)

	U^{11}	U^{22}	U^{33}	U^{12}	U^{13}	U^{23}
Cl1	0.0802 (5)	0.0871 (5)	0.0664 (4)	0.0432 (4)	0.0358 (4)	0.0181 (4)
S1	0.0877 (5)	0.0664 (4)	0.0387 (3)	0.0470 (3)	0.0082 (3)	0.0157 (3)
O1	0.0636 (9)	0.0576 (9)	0.0367 (8)	0.0276 (7)	0.0084 (7)	0.0145 (7)
N1	0.0632 (11)	0.0471 (10)	0.0316 (8)	0.0248 (8)	0.0058 (7)	0.0090 (7)
N2	0.0596 (11)	0.0462 (10)	0.0345 (9)	0.0255 (8)	0.0069 (7)	0.0110 (7)
C1	0.0641 (14)	0.0675 (15)	0.0476 (12)	0.0343 (12)	0.0188 (11)	0.0180 (11)
C2	0.0604 (13)	0.0556 (13)	0.0400 (11)	0.0260 (11)	0.0097 (10)	0.0126 (10)
C3	0.0480 (11)	0.0397 (10)	0.0384 (11)	0.0137 (9)	0.0082 (8)	0.0125 (8)
C4	0.0560 (12)	0.0387 (10)	0.0384 (10)	0.0162 (9)	0.0102 (9)	0.0130 (8)
C5	0.0524 (11)	0.0440 (11)	0.0313 (9)	0.0205 (9)	0.0066 (8)	0.0126 (8)
C6	0.0513 (12)	0.0552 (13)	0.0508 (12)	0.0123 (10)	0.0090 (10)	0.0182 (10)
C7	0.0600 (14)	0.0798 (18)	0.0559 (14)	0.0321 (13)	0.0245 (11)	0.0301 (13)
C8	0.0891 (18)	0.0620 (15)	0.0459 (13)	0.0419 (14)	0.0251 (12)	0.0194 (11)
C9	0.0920 (19)	0.0416 (12)	0.0555 (14)	0.0182 (12)	0.0218 (13)	0.0130 (10)
C10	0.0650 (14)	0.0484 (12)	0.0517 (12)	0.0169 (10)	0.0219 (11)	0.0192 (10)

Geometric parameters (Å, °)

Cl1—C1	1.778 (2)	C2—H2B	0.9700
S1—C4	1.669 (2)	C2—H2C	0.9700
O1—C3	1.221 (2)	C5—C10	1.369 (3)
N1—C3	1.367 (3)	C5—C6	1.375 (3)
N1—C4	1.386 (3)	C6—C7	1.385 (3)
N1—H1A	0.8600	C6—H6A	0.9300
N2—C4	1.320 (3)	C7—C8	1.361 (4)
N2—C5	1.433 (2)	C7—H7A	0.9300
N2—H2A	0.8600	C8—C9	1.367 (4)
C1—C2	1.490 (3)	C8—H8A	0.9300
C1—H1B	0.9700	C9—C10	1.381 (3)
C1—H1C	0.9700	C9—H9A	0.9300

C2—C3	1.503 (3)	C10—H10A	0.9300
C3—N1—C4	128.77 (17)	N2—C4—N1	117.03 (18)
C3—N1—H1A	115.6	N2—C4—S1	123.88 (16)
C4—N1—H1A	115.6	N1—C4—S1	119.09 (15)
C4—N2—C5	122.97 (17)	C10—C5—C6	120.60 (19)
C4—N2—H2A	118.5	C10—C5—N2	119.22 (19)
C5—N2—H2A	118.5	C6—C5—N2	120.17 (19)
C2—C1—C11	111.32 (16)	C5—C6—C7	118.7 (2)
C2—C1—H1B	109.4	C5—C6—H6A	120.7
C11—C1—H1B	109.4	C7—C6—H6A	120.7
C2—C1—H1C	109.4	C8—C7—C6	121.0 (2)
C11—C1—H1C	109.4	C8—C7—H7A	119.5
H1B—C1—H1C	108.0	C6—C7—H7A	119.5
C1—C2—C3	111.71 (17)	C7—C8—C9	119.9 (2)
C1—C2—H2B	109.3	C7—C8—H8A	120.1
C3—C2—H2B	109.3	C9—C8—H8A	120.1
C1—C2—H2C	109.3	C8—C9—C10	120.1 (2)
C3—C2—H2C	109.3	C8—C9—H9A	119.9
H2B—C2—H2C	107.9	C10—C9—H9A	119.9
O1—C3—N1	122.97 (18)	C5—C10—C9	119.7 (2)
O1—C3—C2	123.17 (18)	C5—C10—H10A	120.1
N1—C3—C2	113.86 (17)	C9—C10—H10A	120.1
C11—C1—C2—C3	172.29 (17)	C4—N2—C5—C6	-86.1 (3)
C4—N1—C3—O1	-3.2 (4)	C10—C5—C6—C7	-0.5 (3)
C4—N1—C3—C2	177.1 (2)	N2—C5—C6—C7	-179.04 (19)
C1—C2—C3—O1	4.3 (3)	C5—C6—C7—C8	-0.1 (3)
C1—C2—C3—N1	-176.1 (2)	C6—C7—C8—C9	0.4 (4)
C5—N2—C4—N1	-175.79 (19)	C7—C8—C9—C10	-0.3 (4)
C5—N2—C4—S1	5.2 (3)	C6—C5—C10—C9	0.6 (3)
C3—N1—C4—N2	-2.1 (3)	N2—C5—C10—C9	179.2 (2)
C3—N1—C4—S1	176.94 (18)	C8—C9—C10—C5	-0.2 (4)
C4—N2—C5—C10	95.3 (3)		

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

Cg1 is the centroid of the C5—C10 ring.

$D-H\cdots A$	$D-H$	$H\cdots A$	$D\cdots A$	$D-H\cdots A$
N2—H2A \cdots O1	0.86	2.01	2.677 (3)	134
N1—H1A \cdots S1 ⁱ	0.86	2.53	3.3709 (19)	165
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