Crystal Structure and Properties of N-Phenyl-N'-(2-nitrobenzoyl)thiourea

J.H. HU^{1,*}, Y. LI² and N.P. YAN¹

¹College of Chemical and Biological Engineering, Lanzhou Jiaotong University, Lanzhou, Gansu Province, P.R. China

²College of Chemistry and Chemical Engineering, Northwest Normal University, Lanzhou, Gansu Province, P.R. China

*Corresponding author: Tel: +86 13359460506; E-mail: hujinghan62@163.com

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N-phenyl-N'-(2-nitrobenzoyl)thiourea has been synthesized in high yield under PEG-400 as the phase-transfer catalyst. The structure of the synthesized substituted thiourea was determined by single crystal X-ray diffraction. In DMSO solutions, **L1** exhibited selective recognition for F⁻, OAc⁻ and H₂PO₄⁻. In addition, the compound is also a considerable plant-growth regulator.

Key Words: Aroylthiourea, Anion recognition, Crystal structure, Biological activity.

INTRODUCTION

The development of selective optical signaling systems for anions has received considerable attention in recent decades due to their important roles in biological and environmental processes¹. Since the first synthetic sensor for inorganic anions was reported in 1968², great progress has been made³⁻⁵. Recently, thiourea and its derivatives as neutral receptors have been growing in the field of host-guest chemistry^{6,7}. Among a variety of possible H-bond donor groups, systems employing thiourea moieties have been proven to be very efficient in the design of neutral anion binding receptors⁸. We synthesized a anion sensor bearing NH groups, in addition, in order to achieve colorimetric recognition, we induced nitrophenyl group as signal report group. The results show that it can be used as anion sensor in DMSO, at the same time, by investigating the biological activity of the title compound, we found that the compound has high plant growth regulating activity at low concentration.

EXPERIMENTAL

All reagents for synthesis were analytical grade, commercially and were used without further purification. All the anions were added in the form of tetrabutylammonium (TBA) salts, which were purchased from Sigma-Aldrich Chemical, stored in a vacuum desiccator.

Melting points were measured on an X-4 digital melting-point apparatus (uncorrected). The infrared spectra were performed on a Digilab FTS-3000 FT-IR spectrophotometer. UV-VIS spectra were recorded on a Shimadzu UV-2550 spectrometer. ¹H NMR spectra were recorded on a Varian Mercury

Plus-400 MHz spectrometer with DMSO-*d*₆ as solvent and TMS as an internal reference. Electrospray ionization mass spectra (ESI-MS) were measured on an Agilent 1100 LC-MSD-Trap-VL system. Elemental analyses were performed by Thermo Scientific Flash 2000 organic elemental analyzer.

All the UV-visible experiments were carried out in DMSO on a Shimadzu UV-2550 spectrometer. Any changes in the UV-visible spectra of sensor **L1** were recorded upon the addition of tetrabutylammonium salts while keeping the concentration of sensor **L1** constant in all experiments. The solution of the sensor **L1** (2.0 × 10⁻⁴ M) in DMSO was prepared and stored in dry atmosphere. The solution was used for all spectroscopic studies after appropriate dilution. Solutions of 1.0 × 10⁻² mol L⁻¹ tetrabutylammonium salts of the respective anions were prepared.

Synthesis: N-Phenyl-N'-(2-nitrobenzoyl)thiourea was synthesized in high yield as shown in **Scheme-I**: powdered potassium thiocyanate (7.5 mmol) 2-nitrobenzoyl chloride (5 mmol), PEG-400 (3 % with respect to ammonium thiocyanate) and dichloromethane 20 mL were placed in a dried round-bottomed flask containing a magnetic stirrer bar and stirred at room temperature for 1 h, then aniline (5 mmol) was added and the mixture were stirred for additional 0.5 h. The corresponding thiourea precipitates immediately. The product was filtered, washed with water to remove inorganic salts, dried and recrystallized from DMF-EtOH-H₂O to give the title compound. Yield, 78 %; m.p. 179-181 °C; ¹H NMR (CDCl₃, 400 MHz) δ: 12.28 (s, 1H, NH), 11.62 (s, H, NH), 7.24-8.14 (m, 9H, Ar-H); ¹³C NMR (CDCl₃, 400 MHz) δ: 124.25, 124.94, 127.09, 128.27, 128.86, 130.03, 131.93, 134.27, 137.18, 145.80, 166.68, 177.88; ESI-MS: 301.7 (*m/z* + 1). IR (KBr,

Scheme-I: Synthetic route of **L1**

$\nu_{\max}$, cm^{-1}): 3169, 3028 (NH), 1691 (C=O), 1258 (C=S); anal. calcd. (%) for $\text{C}_{14}\text{H}_{11}\text{N}_3\text{O}_3\text{S}$: C: 55.80, H: 3.68, N: 13.95; Found C: 55.77, H: 3.70, N: 13.97.

Structure determination: The crystal structure of the title compound atom scheme-labeling was shown in Fig. 1. The crystal data and structure refinement details for $\text{C}_{14}\text{H}_{11}\text{N}_3\text{O}_3\text{S}$ are summarized in Table-1, selected bond distances and angles are listed in Tables 2 and 3.

Fig. 1. Crystal structure of the title compound

TABLE-1
CRYSTAL DATA OF THE TITLE COMPOUND

Formula	$\text{C}_{14}\text{H}_{11}\text{N}_3\text{O}_3\text{S}$
f.w.	301.32
Crystal system,	Triclinic
Space group	P_1
a (Å)	6.2013(4)
b (Å)	9.9667(6)
c (Å)	11.9444(8)
α (°)	74.188(3)
β (°)	85.421(3)
γ (°)	83.191(3)
V (Å ³), Z	704.42(8), 2
$F_{(000)}$	312
Reflections collected/unique (R_{int})	4293/2993 [$R_{\text{int}} = 0.0096$]
Obsd reflns [$I \geq 2\sigma(I)$]	2993
data/restraints/parameter	2993/0/198
D_c (Mg m^{-3})	1.421
μ (mm^{-1})	0.243
Goodness-of-fit on F^2	1.051
R_1, wR_2 [$I \geq 2\sigma(I)$]	0.0471, 0.1381

Crystal structure: The structure of the N-phenyl-N'-(2-nitrobenzoyl)thiourea is shown in Fig. 1, the intramolecular N-H...O (2.135 Å) hydrogen bonding between the amide-N and benzoyl-O atoms completes a nearly planar six-membered ring in the central part of the molecule. This group makes dihedral angles of 34.98 (1) and 76.39(1)° with the *o*-nitrobenzoyl and other phenyl rings, respectively.

TABLE-2
SELECTED BOND DISTANCES (nm) AND
BOND ANGLES (°) FOR THE COMPOUND

Bond lengths			
S(1)-C(8)	1.650(2)	C(1)-C(7)	1.508(3)
C(2)-N(3)	1.458(3)	C(7)-O(1)	1.206(3)
C(7)-N(1)	1.367(3)	C(8)-N(2)	1.316(3)
C(8)-N(1)	1.394(2)	C(9)-N(2)	1.422(2)
Bond angles			
C(3)-C(2)-N(3)	117.53(19)	C(1)-C(2)-N(3)	120.15(17)
C(7)-N(1)-C(8)	128.70(19)	C(8)-N(2)-C(9)	129.25(19)

TABLE-3
DATA OF HYDROGEN BOND (nm)
AND ANGLE (°) OF TITLE COMPOUND

D-H...A	D-H (nm)	H...A (nm)	D...A (nm)	D-H...A (°)	Symmetry code
N2-H2N...O1	—	—	2.698(6)	144.8	-x+1, -y+1, -z+2

Fig. 2 showed that the crystals were linked to one-dimensional chain-like supermolecular through the intermolecular hydrogen bonds, which form a eight-membered ring with C8A, N1A, H1NA, S1C, C8C, N1C, H1NC, S1A. Moreover, it is found that the supramolecular are connected by intermolecular hydrogen bond of N(1C)-H(1NC)...S(1A) along the *oc*-axis direction and parallel packing along the *oa*-axis direction. As shown in Fig. 2, we can find the parallel packing of the phenyl rings in the crystal of the compound; however, the distance between the centers of two parallel phenyl rings is 0.6201 nm, which is out of the accepted range of the intermolecular π - π interaction⁹.

Fig. 2. Packing diagram of the title compound

RESULTS AND DISCUSSION

Anion recognition properties: Compound **L1** (2×10^{-5} M) exhibited strong absorption at 303 nm in DMSO solution. When adding 50 equiv of F^- , OAc^- , H_2PO_4^- to the DMSO solution of **L1** (Fig. 3), it indicated that the receptor **L1** exhibited selective recognition for F^- , OAc^- and H_2PO_4^- in DMSO. The same test were applied to Cl^- , Br^- , I^- , HSO_4^- , ClO_4^- , no obvious changes were observed.

Fig. 3. UV-visible absorption of sensor **L1** ($c = 2.0 \times 10^{-5}$ M) in the presence of 50 equiv. various anions in DMSO

UV-visible titrations were carried out in DMSO at a concentration of 2.0×10^{-5} mol L⁻¹ with the addition of tetrabutylammonium OAc⁻, the spectra are shown in (Fig. 4). The plot gives a 1:1 stoichiometric ratio between OAc⁻ and **L1** in (Fig. 5), by the association constants K_a of the sensor¹⁰ **L1** toward OAc⁻ was obtained as 1.5×10^5 M⁻¹.

Fig. 4. UV-visible spectral titration of sensor **L1** with OAc⁻

Fig. 5. Plot of the absorbance at 303 nm vs. equivalents of OAc⁻ in DMSO. $C(\mathbf{L1}) = 2 \times 10^{-5}$ M, the correlation coefficient is 0.998

The whole reaction process was shown in **Scheme-II**. This phenomenon may be attributed to its basicity and geometry configuration to converge the two -NH groups at a point *via* -NH- anions hydrogen binding¹¹.

Scheme-II: Proposed binding mode of **L1** with OAc⁻ in DMSO

Biological activity: We also investigated plant growth regulation activity. Here, we adopted the method of plate culture. The compound was collected to the solutions in concentration of 100, 10, 1, 0.1, 0.01 and 0.001 ppm and rape seeds were cultured in a 10 cm petri-dish with 10 mL different solutions and a circular filter paper. After growing at room temperature for 4 days, the root length was gained and the percentage plant growth activity was calculated according to the following equation:

$$\text{Percentage plant growth activity} = \frac{(N - N_1)}{N_1} \times 100 \%$$

In the equation, N is the root length cultured in compound solutions and N_1 is the root length cultured in the distilled water under the same condition. The data are shown in Table-4.

TABLE-4 PLANT GROWTH REGULATING ACTIVITY DATA (RAPE)						
Compound	Plant growth activity (%) ^a					
	100 ppm	10 ppm	1 ppm	0.1 ppm	0.01 ppm	0.001 ppm
L1	-100	-22.1	67.4	68.2	49.6	16.3
Heteroauxing	-100	-92.5	-91.2	-80.2	-28.9	-10.7

^aSolution was prepared in the proportion of H₂O:DMF = 99.5:0.5 and 0.1 g Tween-80 was added to promote the compound to dissolved.

From the results summarized in Table-4, compared with heteroauxing, the compound exhibited root elongation activity and with the concentration was decreased from 100-0.001 ppm, the root elongation activity becomes strengthener gradually. It was obvious that the compound shown excellent plant-growth regulation activity in the concentration of 0.1 ppm, which the growth activity was reached to 68.2 %.

Conclusion

In summary, we have synthesized N-phenyl-N'-(2-nitrobenzoyl)thiourea in high yield. The crystal structure show that the intramolecular N-H...O hydrogen bonding between the amide-N and benzoyl-O atoms completes a nearly planar six-membered ring in the central part of the molecule, moreover, in the crystals, the moleculars of **L1** were linked to one-dimensional chain-like superamolecular through the intermolecular hydrogen bonds. The anion-binding properties of **L1** has been studied by UV-visible, which is shown to be capable of strongly binding for OAc⁻ through hydrogen bonding interaction in DMSO and formed a 1:1 stoichiometry complex. In addition, investigation of biological activity shows the compound show good plant-growth regulation activity.

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REFERENCES

1. S.Y. Chung, S. W. Nam, J. Lim, S. Park and J. Yoon, *Chem. Commun.*, 2866 (2009).
2. C.H. Park and H.E. Simmons, *J. Am. Chem. Soc.*, **90**, 2431 (1968).
3. F.P. Schmidtchen and M. Berger, *Chem. Rev.*, **97**, 1609 (1997).
4. M.M.G. Antonisse and D.N. Reinhoudt, *Chem. Commun.*, 443 (1998).
5. T.S. Snowden and E.V. Anslyn, *Curr. Opin. Chem. Biol.*, **3**, 740 (1999).
6. D.A. Jose, D.K. Kumar, B. Ganguly and A. Das, *Org. Lett.*, **6**, 3445 (2004).
7. L.S. Evans, P.A. Gale, M.E. Light and R. Quesada, *Chem. Commun.*, 965 (2006).
8. J.H. Hu, W.X. Xu and T.B. Wei, *Spectrosc. Spectr. Anal.*, **6**, 27 (2007).
9. S. Tsuzuki, K. Honda and R. Azumi, *J. Am. Chem. Soc.*, **124**, 12200 (2002).
10. X.D. Yu, H. Lin, Z.S. Cai and H.K. Lin, *Tetrahedron Lett.*, **48**, 8615 (2007).
11. J. Shao, H. Lin and H.K. Lin, *Dyes Pigments*, **80**, 259 (2009).