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Article Synthesis, Characterization and Single Crystal X-ray Diffraction Analysis of Fused Triazolo/Thiadiazole Clubbed with Indole Scaffold

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Abstract: The present synthetic strategy involves the synthesis of indolyl-triazolo-thiadiazole heterocyclic ring systems **8–13** from the condensation of 4-amino-5-(1*H*-indol-2-yl)-3*H*-1,2,4-triazole-3-thione **1** with the aromatic carboxylic acid derivatives **2–7** in presence of POCl₃ for 1 h. All compounds were obtained in very good yields and have been well-characterized using spectroscopic techniques. Exclusively, good quality crystals from the target organic hybrid 8-(1*H*-indol-2-yl)-5-(*p*-tolyl)-[1,2,4]triazolo [3,4-b][1,3,4]thiadiazole **9** were obtained and found suitable for X-ray single crystal diffraction measurement, which is used to confirm and analyze the molecular and supramolecular structure aspects of **9**. The solid-state structure of the synthesized molecule **9** agrees very well with other characterizations. The packing of **9** is dominated by the N … H, S … H, C … C and S … C non-covalent interactions, which agree with the Hirshfeld surface analysis. The percentages of these contacts are calculated to be 20.3%, 5.4%, 9.4% and 4.3%, respectively.

Keywords: fused heterocycle; triazole; thiadiazole; computational study

1. Introduction

Indole chemistry is a wonderful area of research and has gained a lot of attention from researchers for centuries. This fused heterocycle pharmacophore ring exists in many biological hits, which makes this unit a privileged molecule [1]. The indole core structure is found in alkaloid polycyclic compounds and has shown broad-spectrum actions in many pharmacological applications. Among their biological activities, these compounds have been applied as targeting enzyme inhibitors in cancer research, specifically PARP-1 and EGFR inhibitors [2–4], α -glucosidase inhibitor [5] and anti-HIV (anti-human immunod-eficiency virus) [6], and have also exhibited other activities, including anti-microbial [7], anti-inflammation [8], anti-vascular [9], ischemia/reperfusion injury [10], and anti-malarial potential [11].

On the other hand, the 4-amino-1,2,4-triazole-3-thione motif is an interesting scaffold that can be utilized for the synthesis of heterocyclic compounds via annulation or Schiff base formation because it has good functionality. This pharmacophore showed interesting biological patterns, including anti-tumor [12], anti-malarial [13], anti-microbial [14], anti-convulsant [15], and anti-proliferative behaviors [16]. Additionally, it has been used as an



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Copyright: © 2023 by the authors. Licensee MDPI, Basel, Switzerland. This article is an open access article distributed under the terms and conditions of the Creative Commons Attribution (CC BY) license (https:// creativecommons.org/licenses/by/ 4.0/). enzyme inhibitor, such as for protein tyrosine phosphatase 1B [17], AChE enzyme [18], urease enzymes [19] and dizinc metallo- β -lactamase [20]. L. Gavara et al. have designed and synthesized a new Schiff base derived from the 4-amino-1,2,4-triazole-3-thione core structure, which showed high efficacy and more selectivity against metallo- β -lactamases (MBLs) [21]. Recently, Boraei et al. [22] utilized this synthon combined with indole moiety for annulation, and yielded a new triazolo/thiadiazole heterocycle with a confirmed molecular structure, which makes this scaffold interesting and will lead to further research. Triazole and fused rings have shown great biological importance in the medical chemistry and drug discovery research fields [23–27].

The chemical insights yielded by the newly synthesized molecules have attracted the attention of many researchers. One of the most powerful tools that can provide insights into the intermolecular interactions in molecular crystals is Hirshfeld surface analysis. The shape- and size-based Hirshfeld surface analysis approach allows the quantitative and qualitative exploration and visualization of the intermolecular contacts in crystalline molecules. In this context, based on the findings mentioned above and in continuation of our research program focused on heterocyclic chemistry [28–30], we have synthesized new fused triazolo/thiadiazole heterocycles combined with an indole scaffold. The molecular structures of the desired molecules are fully characterized based on spectrophotometric tools, in addition to single-crystal X-ray diffraction (SCXRD) analysis. Hirshfeld surface analysis has also been performed.

2. Materials and Methods

2.1. General

Melting points were measured via a melting point apparatus (SMP10) with open capillaries and are inclded uncorrected. Nuclear magnetic resonance (¹H-NMR and ¹³C-NMR) spectra were determined using DMSO- d_6 on Bruker AC 300 and 400 MHz spectrometers, respectively, in the presence of tetramethylsilane as an internal standard. Chemical shifts are described in δ (ppm) and coupling constants are given in Hz. Elemental analysis was performed on a Flash EA-1112 instrument. A Finnigan MAT 95XP was used to record the mass spectra of HREI experiments. A Jeol JMS HX110 mass spectrometer was used to capture the FAB-MS data.

2.2. General Procedures

A mixture of 3.0 mmol of appropriate aromatic carboxylic acid derivative **2**–7 (3.0 mmol) was refluxed in 10 mL phosphorus oxychloride for 1 h, then left to cool to room temperature, before being poured over ice water. The formed precipitates were collected by filtration, dried and recrystallized from DMF to produce the pure compounds **8–13**. Only 8-(1*H*-indol-2-yl)-5-(*p*-tolyl)-[1,2,4]triazolo [3,4-b][1,3,4]thiadiazole 9 was obtained in a sufficient crystal form for SCXRD measurement.

8-(1H-Indol-2-yl)-5-phenyl-[1,2,4]triazolo [3,4-b][1,3,4]thiadiazole 8

Yield: 87%, m.p. > 300 °C; ¹H NMR (DMSO- d_6 , 300 MHz) δ 7.09 (dd, J = 7.8, J = 7.2 Hz, 1H), 7.22 (dd, J = 7.2, J = 7.8 Hz, 1H), 7.47–7.72 (m, 6H), 8.13 (d, J = 6.6 Hz, 2H), 12.13 (br.s, 1H); ¹³C NMR (DMSO- d_6 , 75 MHz) δ 102.67, 111.97, 119.88, 121.00, 122.27, 123.27, 127.30, 127.58, 128.95, 129.63, 132.95, 137.15, 141.13, 153.45, 167.01; HRMS (EI) calcd for C₁₇H₁₁N₅S (M+): 317.0715. Found: 317.0726.

8-(1*H*-Indol-2-yl)-5-(*p*-tolyl)-[1,2,4]triazolo [3,4-*b*][1,3,4]thiadiazole 9

Yield: 83%, m.p. > 300 °C; ¹H NMR (DMSO- d_6 , 400 MHz) δ 2.42 (s, 3H), 7.08 (dd, J = 8.0, J = 7.5 Hz, 1H), 7.22 (dd, J = 7.5, J = 8.4 Hz, 1H), 7.45–7.51 (m, 4H), 7.71 (d, J = 8.0 Hz, 1H), 7.99 (d, J = 8.0 Hz, 2H), 12.08 (br.s, 1H); ¹³C NMR (DMSO- d_6 , 100 MHz) δ 21.19, 102.69, 112.04, 119.95, 121.09, 122.94, 123.33, 126.31, 127.29, 127.66, 130.24, 137.21, 143.52, 167.13; HRMS (FAB +ve) calcd for C₁₈H₁₄N₅S (M + 1): 332.0970. Found: 332.0955.

3-(1*H*-Indol-2-yl)-6-(4-methoxyphenyl)-[1,2,4]triazolo [3,4-*b*][1,3,4]thiadiazole 10

Yield: 82%, m.p. > 300 °C; ¹H NMR (DMSO- d_6 , 400 MHz) δ 3.91 (s, 3H), 7.11 (t, J = 7.5 Hz, 1H), 7.18–7.31 (m, 3H), 7.40–7.58 (m, 2H), 7.73 (d, J = 7.9 Hz, 1H), 8.10 (d,

 $J = 8.8 \text{ Hz}, 2\text{H}, 12.14 \text{ (s, 1H); } {}^{13}\text{C NMR} \text{ (DMSO-}d_6, 100 \text{ MHz}) \delta 56.19, 103.15, 112.51, 115.57, 120.45, 121.56, 123.40, 123.53, 128.10, 129.70, 137.62, 141.51, 162.90, 163.40; elemental analysis calculated for [C₁₈H₁₃N₅OS]: C, 62.23; H, 3.77; N, 20.16; S, 9.23. Found: C, 62.34; H, 3.86; N, 20.08; S, 9.35.$

8-(1*H*-Indol-2-yl)-5-(4-chlorophenyl)-[1,2,4]triazolo [3,4-*b*][1,3,4]thiadiazole 11

Yield: 92%, m.p. >300 °C; ¹H NMR (DMSO- d_6 , 300 MHz) δ 7.09 (dd, J = 8.1, J = 7.2 Hz, 1H), 7.22 (dd, J = 7.2, J = 8.4 Hz, 1H), 7.46 (s, 1H), 7.49 (d, J = 8.4 Hz, 1H), 7.69–7.76 (m, 4H), 8.16 (d, J = 8.4 Hz, 2H), 12.13 (br.s, 1H); ¹³C NMR (DMSO- d_6 , 75 MHz) δ 102.74, 111.96, 119.96, 121.06, 122.81, 123.36, 127.61, 127.88, 129.12, 129.77, 137.20, 137.66, 141.18, 153.60, 165.97; HRMS (FAB +ve) calcd for C₁₇H₁₁N₅SCl (M+1): 352.0424. Found: 352.0441.

6-(2-Bromophenyl)-3-(1H-indol-2-yl)-[1,2,4]triazolo [3,4-b][1,3,4]thiadiazole 12

Yield: 73%, m.p. 296–297 °C; ¹H NMR (400 MHz, DMSO) δ 7.11 (d, *J* = 7.2 Hz, 1H), 7.24 (d, *J* = 8.1 Hz, 1H), 7.43 (d, *J* = 1.4 Hz, 1H), 7.52 (d, *J* = 8.3 Hz, 1H), 7.84–7.62 (m, 4H), 8.25 (dd, *J* = 7.7, 1.7 Hz, 1H), 12.18 (s, 1H); ¹³C NMR (DMSO-*d*₆, 100 MHz) δ 103.30, 112.53, 120.47, 121.61, 121.88, 123.20, 123.93, 128.08, 129.09, 129.88, 132.87, 134.27, 134.71, 137.70, 141.60, 150.53, 165.88; elemental analysis calculated for [C₁₇H₁₀BrN₅S]: C, 51.53; H, 2.54; Br, 20.16; N, 17.67; S, 8.09. Found: C, 51.70; H, 2.55; Br, 20.18; N, 17.61; S, 8.13.

3-(1H-indol-2-yl)-6-(pyridin-3-yl)-[1,2,4]triazolo [3,4-b][1,3,4]thiadiazole 13

Yield: 77%, m.p. >300 °C; ¹H NMR (DMSO- d_6 , 400 MHz) δ 7.19 (m, 2H), 7.52 (s, 2H), 7.73 (d. *J* = 7.5 Hz, 2H), 8.55 (s, 1H), 8.90 (s, 1H), 9.35 (s, 1H), 12.16 (s, 1H); ¹³C NMR (DMSO- d_6 , 100 MHz) δ 103.50, 112.56, 120.48, 121.58, 123.24, 123.89, 124.95, 128.17, 135.59, 137.78, 148.31, 153.76, 162.82, 165.17; elemental analysis calculated for [C₁₆H₁₀N₆S]: C, 60.36; H, 3.17; N, 26.40; S, 10.07. Found: C, 60.41; H, 3.13; N, 26.32; S, 10.11.

2.3. X-ray Structure Determinations

The technical procedure for the chemical structural elucidation of compound **9** via X-ray single-crystal diffraction analysis is given in the Supplementary Materials (The protocol and refinement crystal data Tables S1–S6). Crystal data were refined using the software from [31–34]. Table 1 lists the data regarding the synthesized crystals of compound **9**.

	9
CCDC	2239222
empirical formula	$C_{18}H_{13}N_5S$
fw	331.39
temp. (K)	120(2) K
$\lambda(\text{\AA})$	0.71073 Å
cryst syst	Monoclinic
space group	$P2_{1}/n$
a (Å)	a = 7.8707(2) Å
b (Å)	b = 15.9681(4) Å
<i>c</i> (Å)	c = 11.9798(4) Å
β (deg)	100.283(3)°
V (Å ³)	1481.44(7) Å ³
Z	4
$ ho_{\rm calc} ({\rm Mg/m^3})$	1.486 Mg/m ³
μ (Mo K α) (mm ⁻¹)	0.228 mm^{-1}
No. reflns.	7305
Unique reflns.	3997
Completeness to $\theta = 25.242^{\circ}$	100%
$GOOF(F^2)$	1.031
R _{int}	0.0213
$R_1 \stackrel{a}{} (I \ge 2\sigma)$	0.0454
$wR_2^{b} (I \ge 2\sigma)$	0.1095

Table 1. Crystal data for 9.

^a $R_1 = \Sigma ||F_0| - |F_c|| / \Sigma |F_0|$. ^b $wR_2 = \{\Sigma [w(F_0^2 - F_c^2)^2] / \Sigma [w(F_0^2)^2] \}^{1/2}$.

2.4. Hirshfeld Surface Analysis

The Crystal Explorer 17.5 program was used to perform the topology analysis of studied compound 9 [35].

3. Results and Discussion

3.1. Chemistry and Characterizations

4-Amino-5-(1*H*-indol-2-yl)-3*H*-1,2,4-triazole-3-thione **1** was mixed separately with benzoic acid, *p*-toluic acid, anisic acid, *p*-chlorobenzoic acid, o-bromobenzoic acid and nicotinic acid **2**–7 in POCl₃ for 1 h; this afforded the indolyl-triazolo-thiadiazoles **8–13**, respectively, in excellent yields (Scheme 1). ¹H- and ¹³C-NMR exhibited the characteristic signals of aromatic protons, between 7.08 ppm and 8.28 ppm, whereas the aromatic carbons were found at 102.0–167.0 ppm. In addition, the NMR study of compound **9** showed the proton for the methyl group at 2.42 ppm, and the carbon of the methyl group at 21.19 ppm. Moreover, the NMR of compound **10** revealed methoxy protons at 3.91 ppm and methoxy carbon at 56.19 ppm. Crystals suitable for the X-ray single-crystal analysis of compound **9** were obtained via recrystallization from DMF.



Scheme 1. Synthesis of the target fused heterocycles.

3.2. Crystal Structure Description

Figure 1 shows the molecular structure of compound 9, obtained by single-crystal X-ray diffraction analysis. Table 1 lists some of the selected bon angles and bond distances of the desired compound. The monoclinic system was observed in the crystallized compound and centrosymmetric $P2_1/n$ space group, with lattice parameters a = 7.8707(2) Å, b = 15.9681(4) Å, c = 11.9798(4) and $\beta = 100.283(3)^{\circ}$. The asymmetric unit contained one molecule, while the unit cell comprised four molecules. The crystal density was 1.486 Mg/m³ and the unit cell volume was 1481.44(7) Å³. The molecule comprised a number of aromatic ring systems, which were perfectly planar but showed some twists. The phenyl group of the aryl moiety was twisted from the mean plane of the fused aromatic ring system (**A**) by only 2.85°. The two fused ring systems **A** and **C** were nearly coplanar with one another. The twist angle between the mean plane of the two fused ring systems was only 0.60°. Selected bond lengths and angles are depicted in Table 2.



Figure 1. ORETP of crystalized compound 9.

Table 2.	Bond lengths	(Å)) and angles	(°)	of 9
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Bond	Length/Å	Bond	Length/Å
S(1)-C(10)	1.7275(18)	N(3)-C(10)	1.306(2)
S(1)-C(11)	1.7734(17)	N(4)-C(10)	1.360(2)
N(1)-C(7)	1.376(2)	N(4)-C(9)	1.370(2)
N(1)-C(8)	1.381(2)	N(4)-N(5)	1.3747(19)
N(2)-C(9)	1.324(2)	N(5)-C(11)	1.304(2)
N(2)-N(3)	1.402(2)		
Bonds	Angle/°	Bonds	Angle/°
C(10)-S(1)-C(11)	87.61(8)	N(1)-C(7)-C(6)	130.16(16)
C(7)-N(1)-C(8)	108.31(14)	N(1)-C(7)-C(2)	107.94(15)
C(9)-N(2)-N(3)	109.54(14)	N(1)-C(8)-C(9)	120.90(15)
C(10)-N(3)-N(2)	104.79(14)	N(2)-C(9)-N(4)	107.75(14)
C(10)-N(4)-C(9)	105.87(14)	N(2)-C(9)-C(8)	127.46(16)
C(10)-N(4)-N(5)	118.88(14)	N(4)-C(9)-C(8)	124.79(15)
C(9)-N(4)-N(5)	135.17(14)	N(3)-C(10)-N(4)	112.04(15)
C(11)-N(5)-N(4)	107.48(14)	N(3)-C(10)-S(1)	138.55(14)
N(5)-C(11)-S(1)	116.67(13)	N(4)-C(10)-S(1)	109.37(12)
C(1)-C(8)-N(1)	109.98(15)	N(5)-C(11)-C(12)	123.27(15)

The molecular units of this compound are controlled by a range of weak non-covalent interactions, including the N ... H, S ... H and C ... C interactions. The molecules are connected to each other by N1-H1 ... N2, N1-H1 ... N3 and C14-H14 ... S1 interactions (Table 3). The different N ... H and S ... H contacts are shown in Figure 2A, while the resulting packing scheme is presented in Figure 2B. Additionally, the packing of **9** is controlled by different levels of C ... C interactions, which are kinds of π - π stacking interactions (Figure 2C). The shortest C ... C interactions are depicted in Table 4.

Table 3. Hydrogen bonds for **9** (Å and °).

D-H A	d(D-H)	d(H A)	d(D A)	<(DHA)
N1-H1 N2#1	0.85(2)	2.12(2)	2.959(2)	169(2)
N1-H1 N3#1	0.85(2)	2.60(2)	3.292(2)	139(2)
C14-H14 S1#2	0.95	2.905	2.905	179.35

Symm. codes: #1 - x + 1, -y + 1, -z and #2 - 1/2 + x, 1.5 - y, -1/2 + z.



Figure 2. The N . . . H/S . . . H contacts (**A**), molecular packing view via N . . . H/S . . . H contacts (**B**) and π - π stacking interactions (**C**) for 9.

Table 4. The shortest C . . . C for π - π stacking interactions.

Contact	Distance	Symmetry Code
C1 C18	3.371	1 - x, 1 - y, 1 - z
C3 C13	3.342	1 - x, 1 - y, 1 - z
C9 C15	3.288	2 - x, 1 - y, 1 - z
C10 C17	3.349	2 - x, 1 - y, 1 - z

3.3. Hirshfeld Surface Analysis

The stability of crystalline materials is controlled by different intermolecular interactions that occur among the adjacent molecules [36,37]. The types, strengths and weights of the intermolecular interactions in the crystal structure could be simply analyzed using Hirshfeld calculations. There are three important maps, which are the d_{norm} , shape index and curvedness maps. A view of these surfaces is shown in Figure 3. In the d_{norm} map, the presence of red regions indicates short contacts, which are considered of great importance to crystal stability.

The decomposition of the fingerprint plot gives the weight of each contact in the crystal structure, and also sheds light on the strength of each contact. The percentages of all contacts contributing to molecular packing are presented in Figure 4. The most important short contacts are the N ... H, S ... H, C ... C and S ... C interactions. Their percentages are calculated to be 20.3%, 5.4%, 9.4% and 4.3%, respectively. Decomposed fingerprint plots of these short contacts are shown in Figure 5. The majority of these interactions appeared in the fingerprint plots as sharp spikes, which confirms that that these interactions are important. A list of the short N ... H, S ... H, C ... C and S ... C contacts contributed to a large extent in the molecular packing, including H ... H and H ... C interactions. The percentages of these interactions are 38.3% and 18.3%, respectively.



Figure 3. Hirshfeld surfaces of 9 showing the most important interactions: $N \dots H(A)$, $S \dots H(B)$, $C \dots C(C)$ and $C \dots S(D)$.



Figure 4. Intermolecular contacts and their percentages in 9.



Figure 5. Decomposed fingerprint plots for the short contacts in 6.

Contact	Distance	Contact	Distance
N2 H1	1.963	C9 C15	3.288
N3 H1	2.483	C10 C17	3.349
N3 H6	2.512	C6 S1	3.469
S1 H14	2.772	C7 S1	3.42

Table 5. Short contacts and the corresponding interaction distances.

Another important feature of molecular packing is the π - π stacking interactions. The presence of these types of intermolecular contact is indicated by the presence of short C9... C15 (3.288 Å) and C10... C17 (3.349 Å) interactions. Additionally, the presence of π - π stacking interactions is clearly implied by the presence of red/blue triangles and a flat green area in the shape index and curvedness maps, respectively (Figure 3).

4. Conclusions

New heterocyclic systems including indole, triazole amd thiadiazole rings **8–13** were synthesized from the reaction of 4-amino-5-(1*H*-indol-2-yl)-3*H*-1,2,4-triazole-3-thione and benzoic acid derivatives in POCl₃. The success of our strategy to design target compounds is proven by our obtaining of the hybrid heterocycle **9** in a good crystalline form. Its structure was revealed by measuring its X-ray single-crystal structure. The resulting X-ray structure was used to analyze the molecular packing of the newly synthesized compound **9**. Additionally, the type, strength and weight of the intermolecular interactions in the crystal structure were analyzed using Hirshfeld calculations. It was found that the N ... H (20.3%), S ... H (5.4%), C ... C (9.4%) and S ... C (4.3%) non-covalent interactions were the most important.

Supplementary Materials: The following supporting information can be downloaded at: https://www. mdpi.com/article/10.3390/cryst13030423/s1, X-ray determination protocol for compound **9** [31,33,34]. Table S1: Atomic coordinates (×10⁴) and equivalent isotropic displacement parameters (Å2 × 10³) for **9**. U(eq) is defined as one third of the trace of the orthogonalized Uij tensor; Table S2: Selected bond lengths [Å] and angles [°] for **9**; Table S3: Bond lengths [Å] and angles [°] for **9**; Table S4: Anisotropic displacement parameters (Å2 × 10³) for **9**. The anisotropic displacement factor exponent takes the form: -2p2[h2 a*2U11 + ... + 2 h k a* b* U12; Table S5: Hydrogen coordinates (×10⁴) and isotropic displacement parameters (Å2 × 10³) for **9**; Table S6: Torsion angles [°] for **9**.

Author Contributions: Conceptualization, A.B., A.T.A.B. and M.F.Y.; methodology, M.S.A., M.F.Y. and E.M.G.; software, M.H. and S.M.S.; formal analysis, M.S.A., A.T.A.B., M.F.Y., E.M.G. and S.O.A.; X-ray crystal structure: M.H.; investigation, M.S.A. and S.M.S.; resources, A.T.A.B., A.B. and M.S.A.; writing—original draft preparation, A.T.A.B., M.F.Y., E.M.G., S.M.S. and A.B.; data curation, S.O.A.; writing—review and editing: All authors have prepare. All authors have read and agreed to the published version of the manuscript.

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