


 Cite this: *RSC Adv.*, 2024, 14, 16546

Design of potent tyrosinase inhibiting *N*-arylated-4-yl-benzamides bearing 2-aminothiazole-triazole bi-heterocycles: mechanistic insight through enzyme inhibition, kinetics and computational studies†

 Farhan Mahmood Khan,^a Muhammad Athar Abbasi,^{ID} *^a Aziz-ur Rehman,^a Sabahat Zahra Siddiqui,^a Abdul Rehman Sadiq Butt,^a Hussain Raza,^{ID} ^b Mubashir Hassan,^c Syed Adnan Ali Shah,^{d,e} Muhammad Shahid^f and Song Ja Kim^{ID} ^b

By using a convergent methodology, a unique series of *N*-arylated 4-yl-benzamides containing a bi-heterocyclic thiazole-triazole core was synthesized and the structures of these hybrid molecules, **9a–k**, were corroborated through spectral analyses. The *in vitro* studies of these multi-functional molecules demonstrated their potent mushroom tyrosinase inhibition relative to the standard used. The kinetics mechanism was exposed by lineweaver–burk plots which revealed that, **9c**, inhibited mushroom tyrosinase non-competitively by forming an enzyme–inhibitor complex. The inhibition constant K_i calculated from Dixon plots for this compound was 0.016 μM . The computational study was also consistent with the experimental results and these molecules disclosed good results of all scoring functions and interactions, which suggested a good binding to mushroom tyrosinase. So, it was predicted from the inferred results that these molecules might be considered as promising medicinal scaffolds for the diseases associated with the over-expression of this enzyme.

 Received 11th February 2024
 Accepted 6th May 2024

DOI: 10.1039/d4ra01063a

rsc.li/rsc-advances

1. Introduction

Heterocyclic compounds are very important biologically and are found in nature as components of vitamins, alkaloids, hormones, and many other natural products. Several heterocycles are used in combinatorial and supramolecular chemistry, as well as in medicine, agriculture, and polymer industries.^{1–3}

Thiazoles are a part of numerous natural products *e.g.*, epothilone, thiostrepton, thiamine pyrophosphate (TPP), carboxylase vitamin B₁, and penicillin.^{4,5} Thiazoles are claimed to play a role in drug development for the treatment of allergies,

HIV infections, hypertension, bacterial infections, schizophrenia and pain,⁶ as novel inhibitors of bacterial DNA gyrase B,⁷ and as fibrinogen receptor antagonists with antithrombotic activity.⁸ They displayed magnificent pharmaceutical activities, for instance, antifungal,⁹ antimicrobial,¹⁰ anti-inflammatory,¹¹ analgesic,¹² anticancer,¹³ and anticonvulsant activities.¹⁴

One of the isomeric forms of triazole is 1,2,4-triazole.¹⁵ These triazole compounds have various biological activities, including anticancer, antimicrobial, anti-inflammatory, antioxidant, anti-HIV and antimalarial.^{16,17} Some of the marketed drugs containing 1,2,4-triazole moiety, like anastrozole, letrozole, and vorozole are used for the treatment of breast cancer. Several compounds bearing 1,2,4-triazole moiety are stated as potent inhibitors of tyrosinase.¹⁸

Tyrosinase is a copper containing metalloenzyme with two Cu⁺² ions in its active site. It is commonly found in plants, animals, and microorganisms.¹⁹ Tyrosinase is essential in facilitating the enzymatic browning of fruits and vegetables, which results in a significant loss of flavour and nutritional content.²⁰ In humans, dopaquinone is converted to melanin by a series of reactions and tyrosinase plays an important role for its creation, which can result in hyperpigmentation disorders including melisma and seborrheic.²¹ Tyrosinase is an enzyme that is only present in melanoma cells, therefore, its blocking is helpful to develop a highly targeted treatment for skin cancer.²²

^aDepartment of Chemistry, Government College University, Lahore, 54000, Pakistan. E-mail: abbasi@gcu.edu.pk; Tel: (+92)-42-111000010 Ext. 266

^bDepartment of Biological Sciences, College of Natural Sciences, Kongju National University, Gongju, 32588, South Korea

^cThe Steve and Cindy Rasmussen Institute for Genomic Medicine, Nationwide Children Hospital, Columbus, Ohio, 43205, USA

^dFaculty of Pharmacy, Universiti Teknologi MARA Cawangan Selangor Kampus Puncak Alam, Bandar Puncak Alam, Selangor, 42300, Malaysia

^eAtta-ur-Rahman Institute for Natural Product Discovery (AuRIns), Universiti Teknologi MARA Cawangan Selangor Kampus Puncak Alam, Bandar Puncak Alam, Selangor, 42300, Malaysia

^fDepartment of Biochemistry, University of Agriculture, Faisalabad, 38040, Pakistan

† Electronic supplementary information (ESI) available. See DOI: <https://doi.org/10.1039/d4ra01063a>



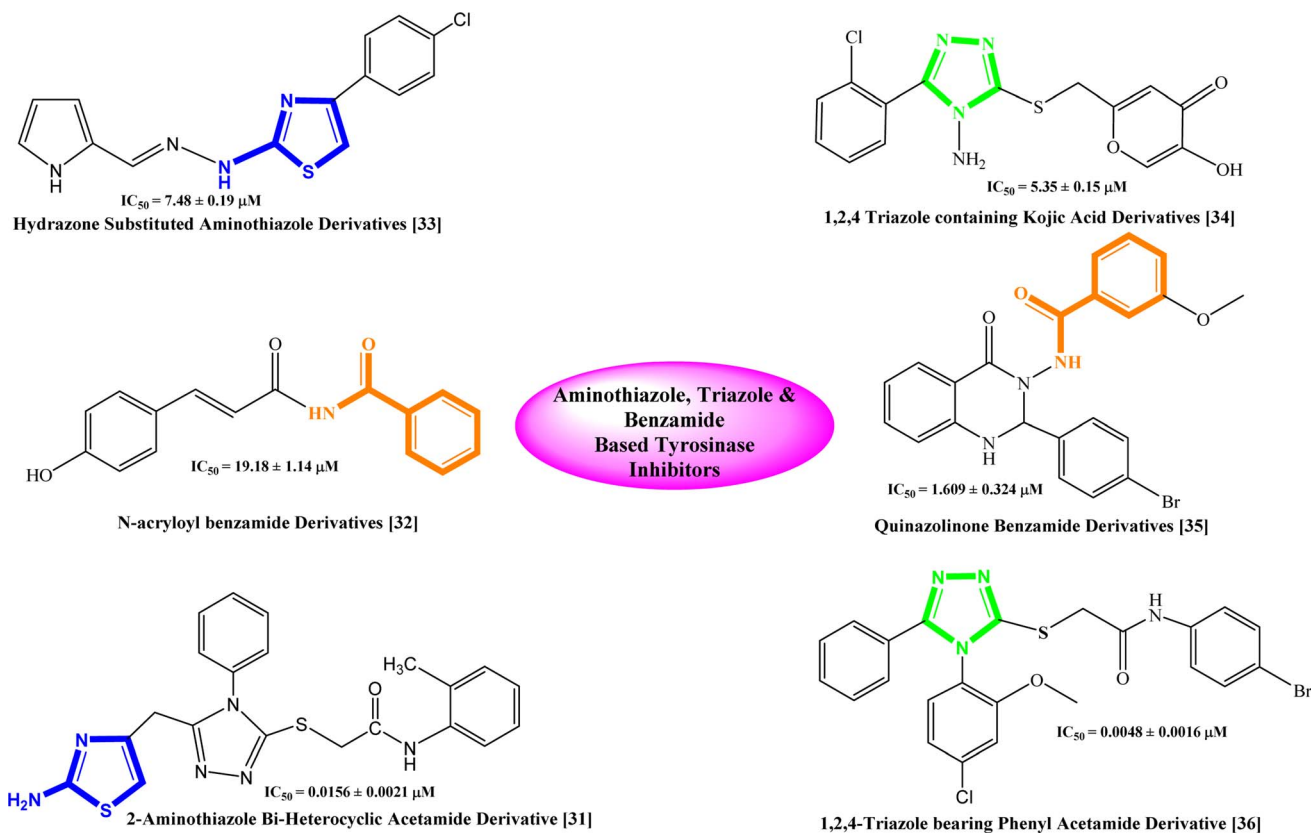


Fig. 1 Rationale of the current study.

However, it is probable that these inhibitors only target the cancer cells, leaving the DNA of healthy cells unaffected.^{23–25}

Sulpiride and amisulpiride are two active benzamides which have been broadly applied in psychiatry and in the treatment of neurodegenerative disorders, schizophrenia, and Alzheimer's disease.^{26,27} Some benzamide derivatives have been reported to exhibit good enzyme inhibitory, antimicrobial, anti-inflammatory, serotonin (5-HT) and antitumor activities.^{28,29}

Based on the recent studies, phenolic compounds and their derivatives and several compounds including terpenoid, phenyl, pyridine, piperidine, pyridinone, hydroxypyridinone, thiosemicarbazone, thiosemicarbazide, azole, thiazolidine, kojic acid, benzaldehyde and xanthate derivatives were characterized as potent tyrosinase inhibitors.³⁰ Similarly, some bi-heterocyclic compounds have been reported previously as tyrosinase inhibitors and some of them are shown with their IC_{50} value in Fig. 1,^{31–36} However, to completely cope with the issues of skin disorders, *i.e.* ephelides, melasma, and senile lentigenes, it is imperative to find some more unique and secure inhibitors of this enzyme.

2. Results and discussion

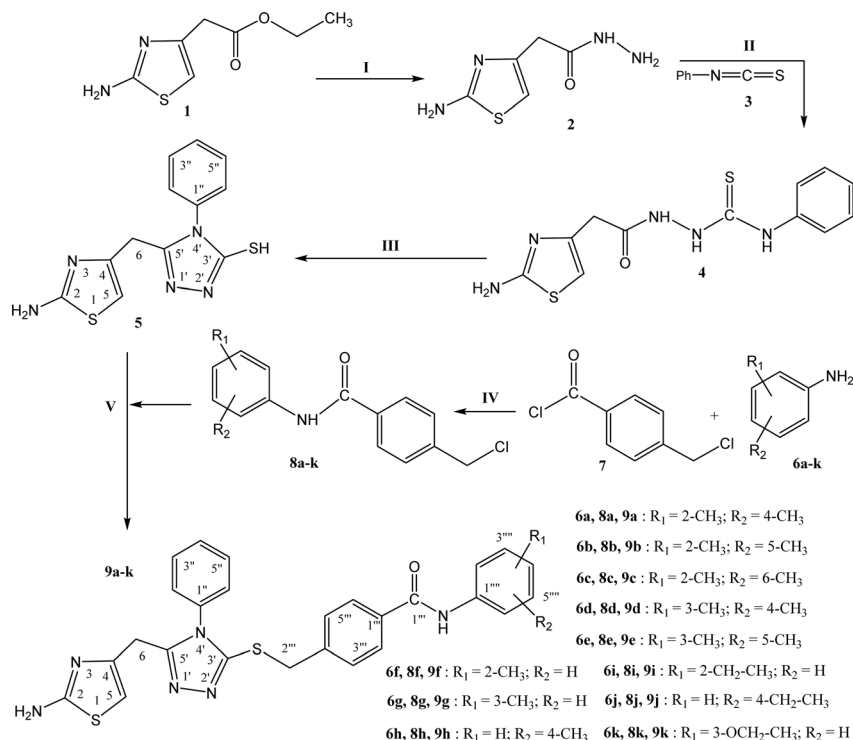
2.1 Chemistry

The designed bi-heterocyclic *N*-arylated benzamides were synthesized by a convergent protocol as described in Scheme 1. The synthesis was initiated by refluxing ethyl 2-(2-amino-1,3-thiazol-4-yl)acetate (1) in methanol and hydrazine hydrate to

transform this starter into 2-(2-amino-1,3-thiazol-4-yl)aceto-hydrazide (2). The hydrazide, 2, was added in methanol and refluxed with phenyl isothiocyanate (3) to convert it into a solid intermediary compound 2-[2-(2-amino-1,3-thiazol-4-yl)acetyl]-*N*-phenyl-1-hydrazinecarbothioamide (4) which was cyclized to a nucleophilic molecule 5-[2-(2-amino-1,3-thiazol-4-yl)methyl]-4-phenyl-4*H*-1,2,4-triazole-3-thiol (5). In a complementary set of reactions, the substituted anilines (6a–k) were reacted with 4-chloromethyl benzoyl chloride (7) in an aqueous basic medium, to acquire the required electrophiles, 4-chloromethyl-*N*-(aryl)benzamides (8a–k). Finally, the nucleophile 5 was dissolved in DMF and a nip of LiH was added into it. To activate its mercapto position, the mixture was stirred for about 30 minutes, and in this last step, the electrophiles, 8a–k, were coupled in equimolar amounts with the activated 5 to yield the targeted bi-heterocyclic benzamides (9a–k).

The detailed structural analysis of a compound, 9c, is discussed hereby. It was synthesized as a light brown amorphous solid, having melting point of 110–111 °C. The molecular formula, $C_{28}H_{26}N_6OS_2$, was affirmed by its CHN analysis data, supported by the count of number of protons in its $^1\text{H-NMR}$ (Fig. 2 and 3) and carbons in its $^{13}\text{C-NMR}$ spectrum (Fig. 4). Different functionalities in the molecule were ascertained by its IR spectrum. The characteristic peaks appeared at ν 3346 (N–H stretch), 3056 (C–H stretch), 2927 (–CH₂– stretch), 1679 (C=O stretch), 1560 (C=C stretch), 1520 (C=N stretch), 1166 (C–N–C bond stretch), 626 (C–S stretch) cm^{-1} . In its $^1\text{H-NMR}$ spectrum, the phenyl ring linked with nitrogen (4') of 1,2,4-triazole





Scheme 1 Outline for the synthesis of 4-(((5-((2-amino-1,3-thiazol-4-yl)methyl)-4-phenyl-4H-1,2,4-triazol-3-yl)sulfanyl)methyl)-N-(aryl)benzamides. Reagents & conditions: (I) MeOH/N₂H₄·H₂O/refluxing for 2 h (yield = 91%). (II) MeOH/**3**/refluxing for 1 h (yield = 88%). (III) The ppt. of **4** dissolved in 10% NaOH/filtration/acidification of filtrate in cold state to get ppt. of **5** (yield = 87%). (IV) Aqueous Na₂CO₃ solution/pH 9–10/vigorous manual shaking at RT for 2–3 h (yield = 89%). (V) DMF/LiH/stirring for 12–24 h (yield = 92–98%).

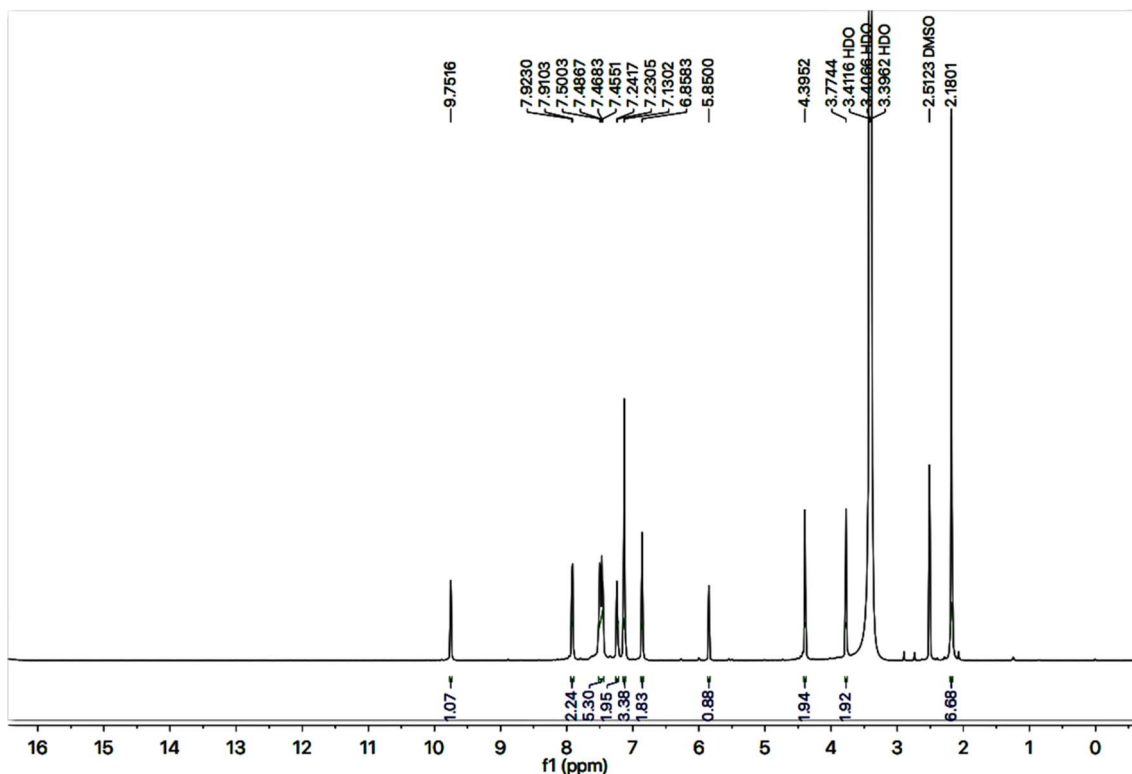
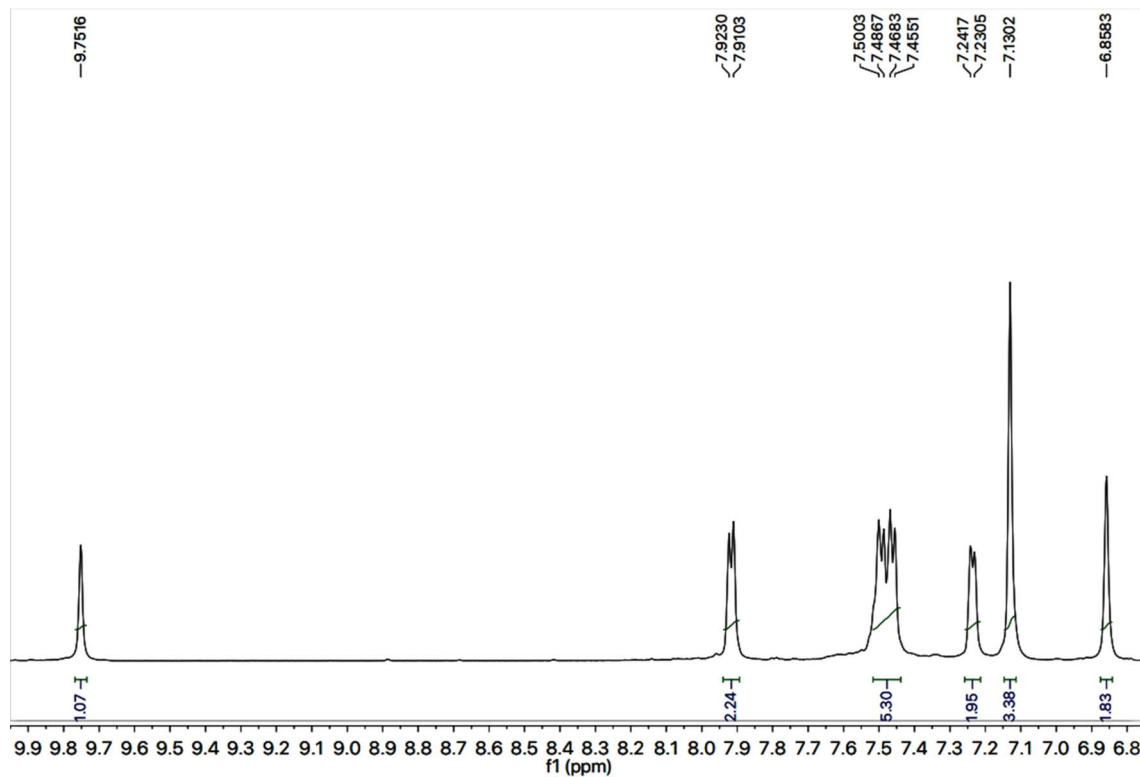
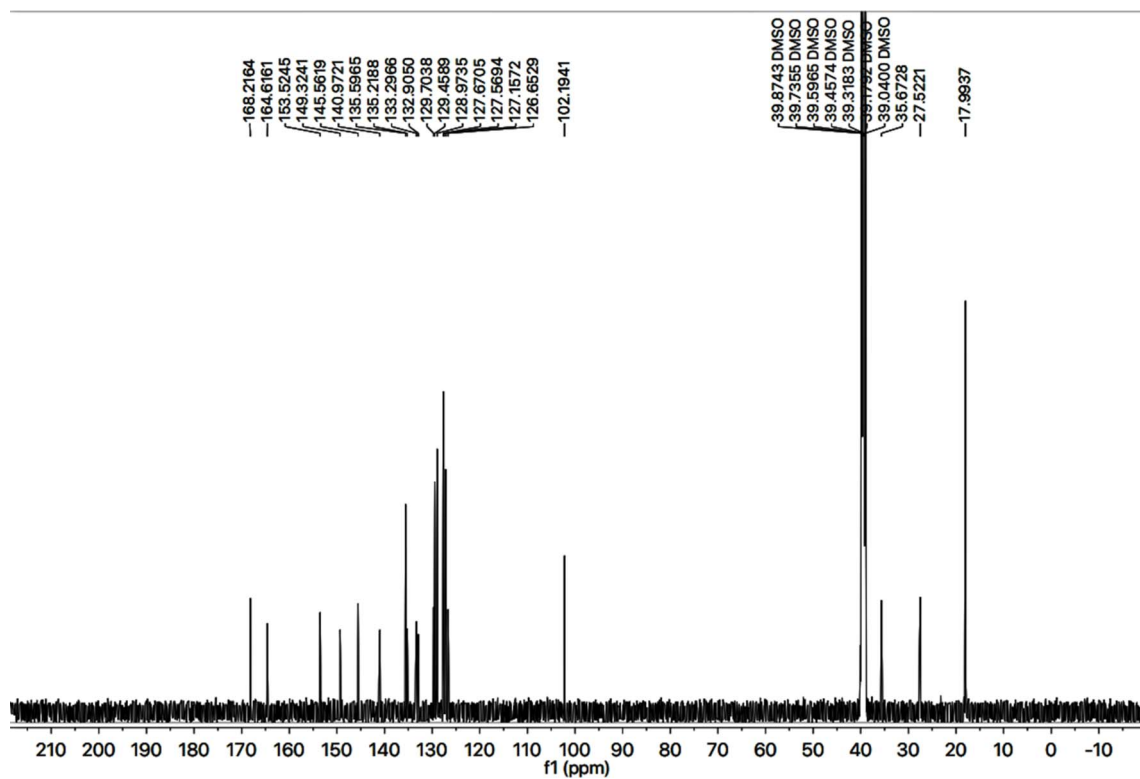


Fig. 2 ¹H-NMR spectrum of compound **9c**.



Fig. 3 Down field region of ^1H -NMR spectrum of 9c.Fig. 4 ^{13}C -NMR spectrum of compound 9c.

heterocyclic ring was identified by a broad doublet at 7.23 (br. d, $J = 6.7$, 2H, H-2'' & H-6'') and a multiplet in the aromatic region at δ 7.50–7.48 (m, 3H, H-3'', H-4'' & H-5''). The 4-substituted benzamide unit was predictable by an amidic signal at δ 9.75 (s, 1H, -CO-NH-1''') along with two prominent *ortho*-coupled broad doublets in the aromatic region at δ 7.91 (br. d, $J = 7.6$, 2H, H-2''' & H-6''') and 7.45 (br. d, $J = 7.9$, 2H, H-3''' & H-5'''). The 2,6-dimethylphenyl moiety connected with nitrogen of benzamide unit was signified by a broad singlet at δ 7.13 (br. s, 3H, H-3''', H-4'''' & H-5''') in the aromatic region, along with a typical signal of two symmetrical methyl groups resonating at 2.18 (s, 6H, CH₃-2'''' & CH₃-6'''). The 2-amino-1,3-thiazol-4-yl unit was characterized by two singlets at δ 6.85 (br. s, 2H, H₂N-2) and 5.85 (s, 1H, H-5), while a singlet at δ 3.77 (s, 2H, CH₂-6) was assignable to a methylene group joining the two heterocycles in the molecule. Another singlet at δ 4.39 (s, 2H, CH₂-8''') was rational for the methylene group linking the benzamide unit with the mercapto position of the triazole moiety.

The 2-amino-1,3-thiazol-4-yl moiety in its ¹³C-NMR spectrum (Fig. 4) was clearly indicated by two quaternary signals at δ 168.21 (C-2), and 145.56 (C-4), along with a methine signal at δ 102.19 (C-5). Similarly, the other heterocycle *i.e.* (1,2,4-triazolo-2-yl)sulfanyl was also identified by two quaternary signals at δ 153.52 (C-5') and 149.32 (C-3') while a methylene linking the two heterocycles (4-position of the former heterocycle with 5'-position of the latter heterocycle) was apparent at δ 27.52 (C-6). The 2,6-dimethylphenyl moiety was also corroborated with two quaternary signals at δ 135.59 (C-2'''' & C-6''') and 135.21 (C-1'''), along with symmetrical duplet methines at δ 127.56 (C-3'''' & C-

5''') and another methine at δ 126.65 (C-4'''). The symmetrical 2,6-dimethyl groups on this aromatic ring were verified by a signal at δ 17.99 (CH₃-2'''' & CH₃-6'''). The 4-substituted benzamide unit was established by a distinct signal of carbonyl carbon at δ 164.61 (C-7''') in addition to other two quaternary signals at δ 140.97 (C-4'''), and 133.29 (C-1''') along with two methine duplet resonances at δ 128.97 (C-3''' & C-5''') and 127.67 (C-2''' & C-6'''). The signal at δ 35.67 (C-8''') was assignable to a methylene joining the benzamide from its 4-position to the sulfur atom joined with 1,2,4-triazole heterocycle. The phenyl group attached to the nitrogen (4') atom of 1,2,4-triazole ring was distinctive by a quaternary signal at δ 132.90 (C-1''), a methine signal at 129.70 (C-4'') along with two symmetrical methine duplets at δ 129.45 (C-3'' & C-5'') and δ 127.15 (C-2'' & C-6''). The C-H connectivities in the carbon skeleton were comprehensively certified by its HMBC spectrum and the important correlations are demonstrated on this spectrum in (Fig. 5).

So, based upon aforementioned evidences and the EIMS spectra of compound **9c** (Fig. 6) the structure of **9c** was confirmed and it was named as 4-[(5-[(2-amino-1,3-thiazol-4-yl)methyl]-4-phenyl-4*H*-1,2,4-triazol-3-yl)sulfanyl)methyl]-*N*-(2,6-dimethylphenyl)benzamide. Similarly, the structures of all other synthesized derivatives were verified through rigorous spectral analyses (Fig. S1–S20†).

2.2 Mushroom tyrosinase inhibition

These synthesized bi-heterocyclic benzamides (**9a–k**) were assessed for their inhibiting potentials against tyrosinase

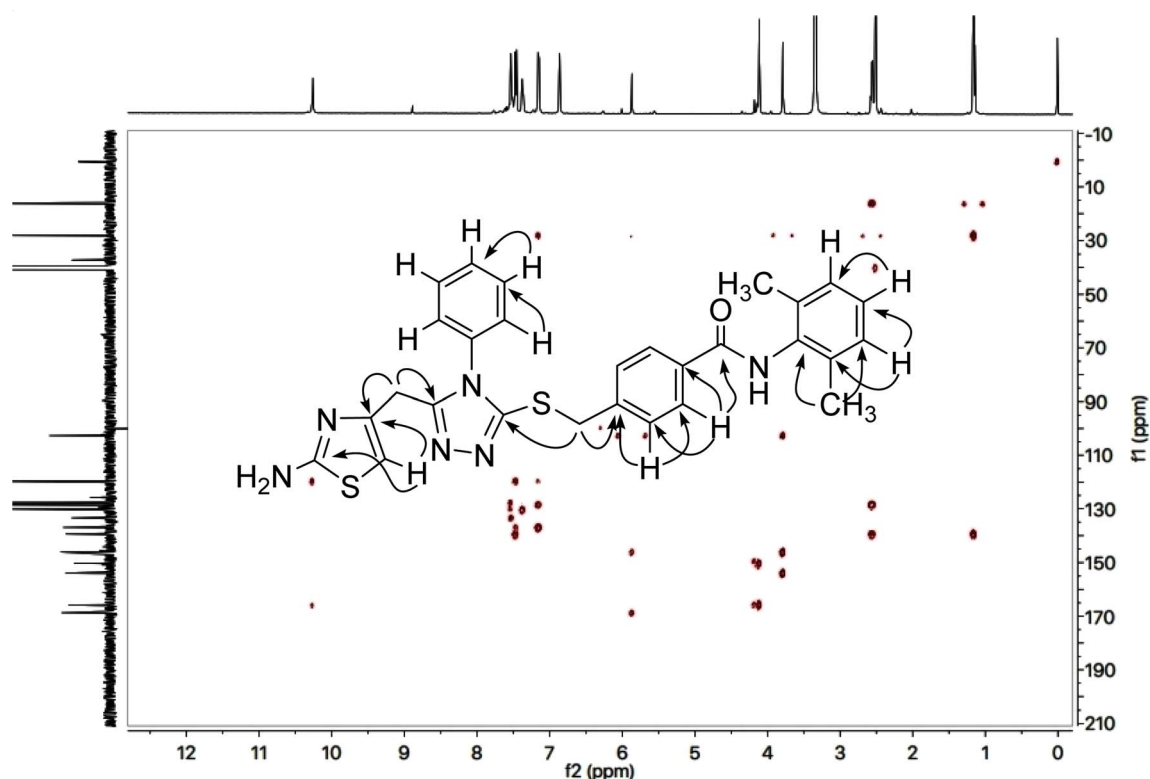
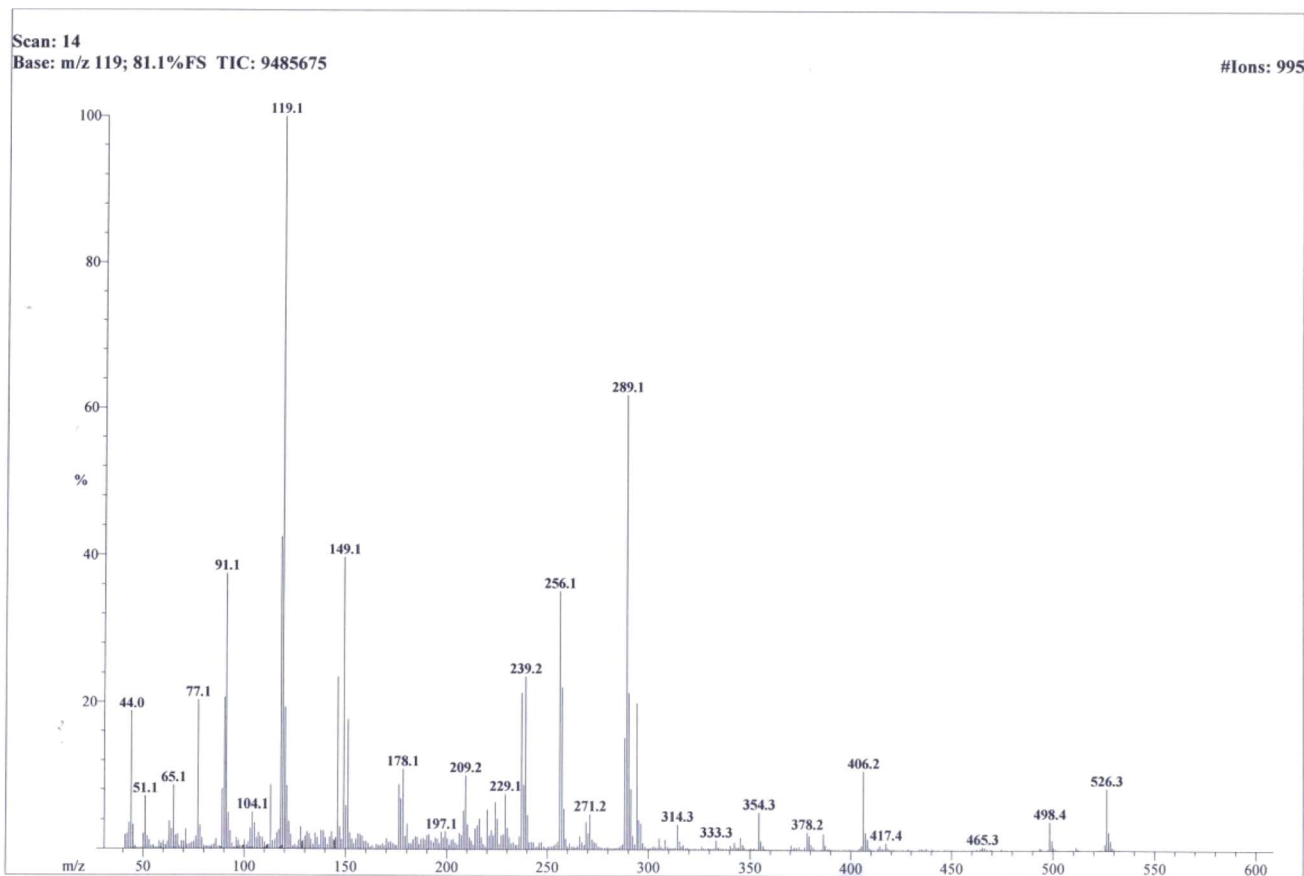
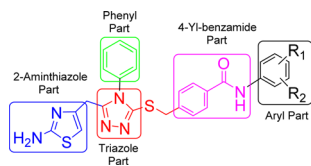


Fig. 5 Important HMBC correlations of compound **9c**.



Fig. 6 EI-MS spectrum of compound **9c**.Fig. 7 General structural features of compounds **9a–k**.

enzyme and their results are tabulated in Table 2. All these compounds exhibited very potent inhibitory activities against this enzyme, as evident from their lower IC_{50} (μM) values as compared to the standard, kojic acid, having IC_{50} value of $16.8320 \pm 1.1600 \mu\text{M}$. Though the rendered activity is an accumulative attribute of a whole molecule, yet a limited structure–activity relationship (SAR) was recognized by examining the effect of different aryl entities on the inhibitory potential, as it was the only varying part in all these molecules. The general structural parts of the studied benzamides are featured in Fig. 7.

2.3 Structure–activity relationship

Amongst three di-methylated regio-isomers, compound **9c** in which the methyl groups were present at both *ortho* positions exhibited better inhibitory activity ($IC_{50} = 0.008 \pm 0.009 \mu\text{M}$) as compared to other isomers in which methyl groups were on

different positions (2,4-dimethylphenyl, 2,5-dimethylphenyl). These analogues **9a** and **9b** possessed comparatively high IC_{50} values; 1.277 ± 0.083 and $0.371 \pm 0.062 \mu\text{M}$, indicating that the presence of the methyl groups at symmetrical *ortho*-positions in former molecule, probably increased its appropriate interactions with the active site of the enzyme (Fig. 8).

Similarly, when the inhibitory potential of other two di-methylated isomers was compared in which the one small-sized methyl group was fixed at 3-position, it was observed that **9d** with adjacent methyl groups (3,4-dimethylphenyl), had a lesser activity ($IC_{50} = 0.419 \pm 0.162 \mu\text{M}$), as compared to **9e** ($IC_{50} = 0.025 \pm 0.011 \mu\text{M}$). Indeed, the molecule **9e** was the second most potent among the series bearing the methyl groups now at *meta* but again the symmetrical positions in aryl part, which is presumed to augment the appropriate interactions with the active site of the enzyme and increase its inhibitory activity (Fig. 9).

The three *mono*-methylated compounds **9f**, **9g** and **9h** having $IC_{50} = 0.098 \pm 0.028$, 0.299 ± 0.081 and $0.622 \pm 0.099 \mu\text{M}$, showed variable activities owing to change in the position of the methyl group in the aryl part. The molecule **9f** in which the methyl group was at *ortho*-position and closer to the benzamido group displayed superb activity while a decreasing trend was observed when the methyl group was present away from the benzamido group, in *meta* and *para*-isomer (Fig. 10).



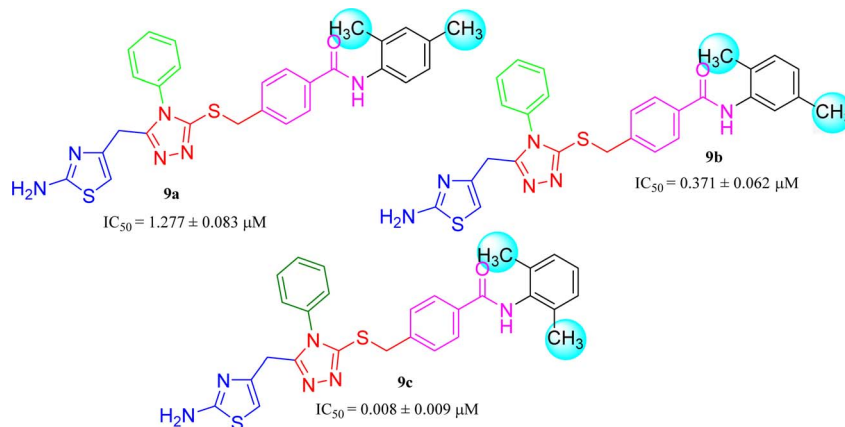


Fig. 8 SAR of compounds 9a, 9b, and 9c.

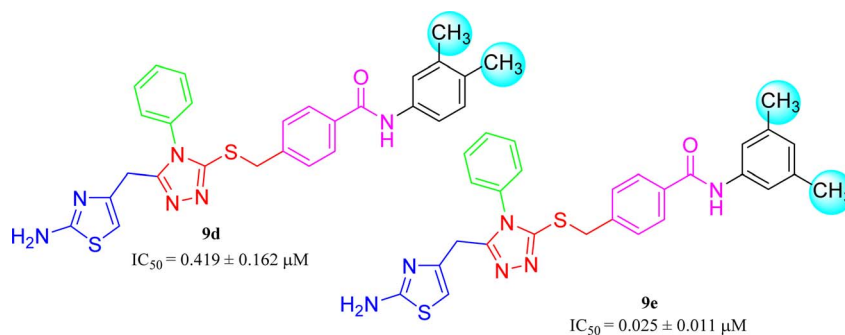


Fig. 9 SAR of compounds 9d and 9e.

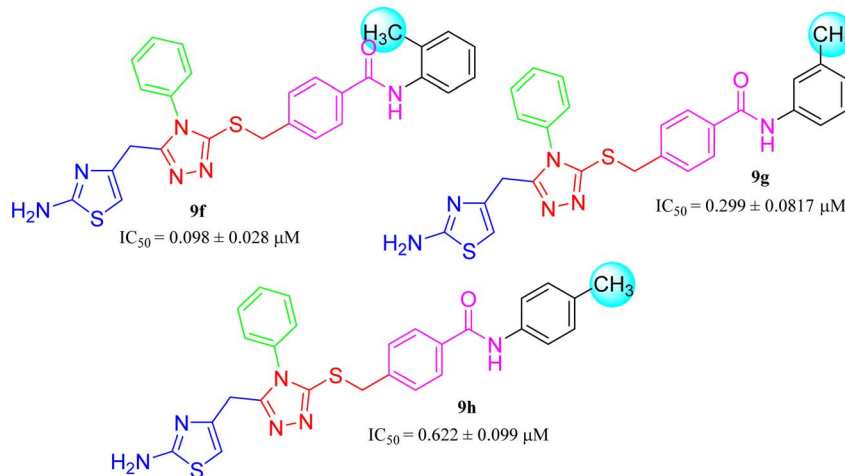


Fig. 10 SAR of compounds 9f, 9g and 9h.

The compound **9i** has a non-polar ethyl group at the *ortho*-position while **9j** has the same ethyl group at *para*-position (Fig. 11). A similar trend was observed in these *mono*-substituted molecules, as the *ortho*-isomer ($IC_{50} = 0.027 \pm 0.015 \mu\text{M}$), probably due to a compactness in size, possessed better inhibitory activity as compared to the respective *para*-isomer ($IC_{50} = 0.089 \pm 0.077 \mu\text{M}$). Anyhow, these two compounds

displayed impressive inhibition relative to *mono*-methylated molecules, presented in the aforementioned set. This order was also consistent to some extent with the +I observations, as the ethyl group has more +I effect than methyl group.

9k ($IC_{50} = 0.044 \pm 0.061 \mu\text{M}$) has medium-sized polar ethoxy group at *meta* position. Owing to varied nature of this group, having -I, and +R effects, it was presumed to exhibit good



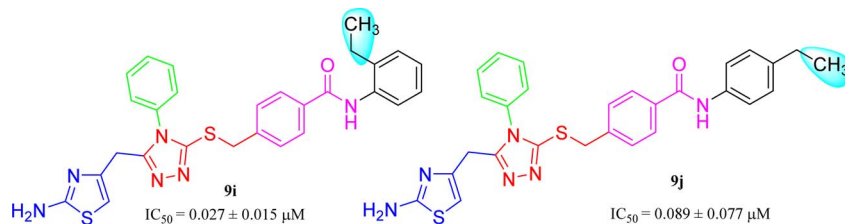


Fig. 11 SAR of compounds 9i and 9j.

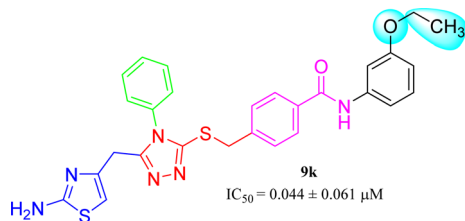


Fig. 12 SAR of compound 9k.

interactions with the enzyme and the molecule overall exhibited a considerably good inhibitory potential (Fig. 12).

2.4 Kinetic analysis

To understand the inhibitory mechanism of synthetic compounds on the tyrosinase inhibition, kinetic study was performed. Based on our IC_{50} results, the most potent compound 9c was selected to determine their inhibition type and inhibition constant. The kinetic results of the enzyme by the Lineweaver–Burk plot of $1/V$ versus $1/[S]$ in the presence of different inhibitor concentrations furnished a series of straight lines (Fig. 13A). The results of 9c showed that compound intersected within the second quadrant. The analysis showed that V_{max} decreased to new increasing doses of inhibitors on the

other hand K_m remained the same. This behavior indicated that 9c inhibited the tyrosinase non-competitively to form the enzyme–inhibitor complex. The secondary plot of slope against the concentration of inhibitors showed enzyme inhibitor dissociation constant (K_i) (Fig. 13B).

2.5 Hemolytic activity

The synthesized compounds, 9a–k, were also exposed to hemolytic assay to ascertain their harmless utility as medicinal scaffolds. Cytotoxicity was profiled as percentage of hemolysis and the results are shown in Table 1. It was obvious from the results that all the derivatives of this series had modest toxicity towards red blood cell membrane as compared to the positive control, Triton X ($94.11 \pm 0.01\%$). The minimum toxicity was rendered by 9h ($2.64 \pm 0.04\%$) in the series. In general, it was rational that most of the molecules exhibited very moderate cytotoxicity and thus can be deliberated as safe therapeutic agents for further applications in drug designing program.

2.6 Docking studies of 9a–k into mushroom tyrosinase enzyme

The docked complexes of all the compounds 9a–k against mushroom tyrosinase were analyzed separately and evaluated because of minimum energy values (kcal mol^{-1}) and ligand

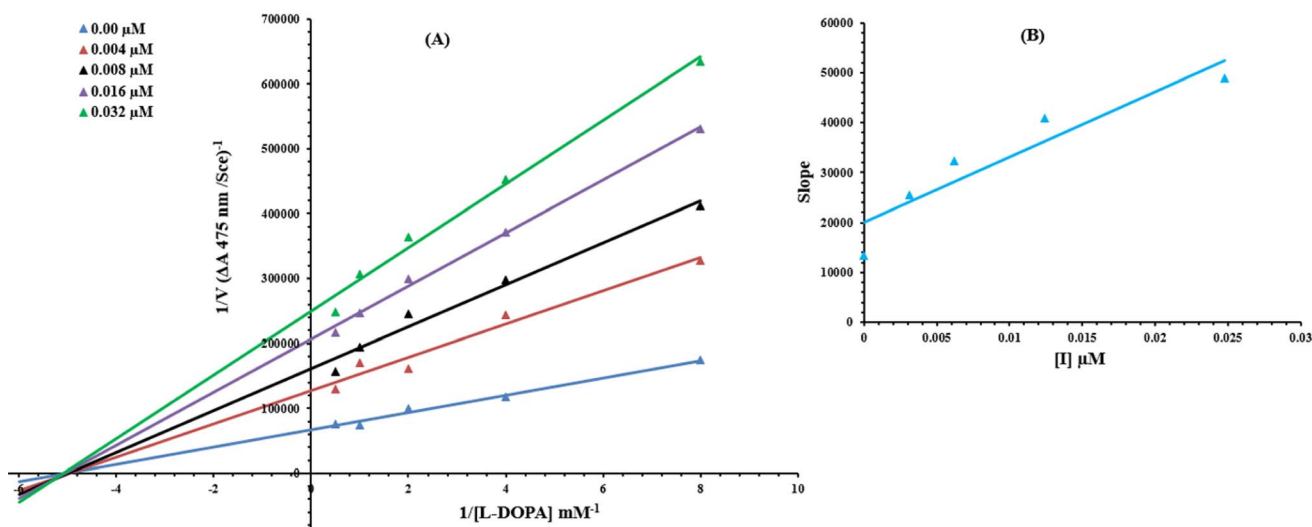
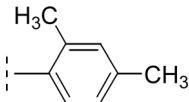
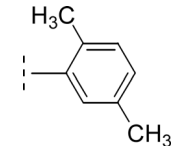
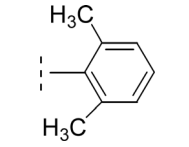
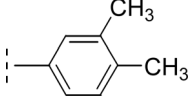
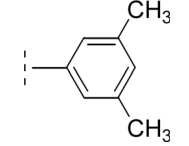
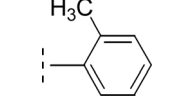
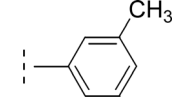
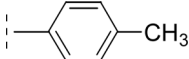
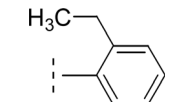
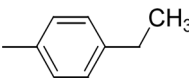
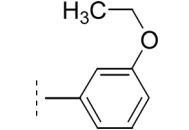


Fig. 13 Lineweaver–Burk plots for inhibition of tyrosinase in the presence of compound 9c. (A) Concentrations of 9c were 0.00, 0.004, 0.008, 0.016 and 0.032 μM , respectively. Substrate L-DOPA concentrations were 0.125, 0.25, 0.5, 1 and 2 mM, respectively. (B) The insets represent the plot of the slope versus inhibitor 9c concentrations to determine inhibition constant. The lines were drawn using linear least squares fit.

Table 1 Tyrosinase inhibitory and hemolytic activity of bi-heterocyclic benzamides, 9a–k^{ab}

Compounds	Aryl part	Mushroom tyrosinase activity IC ₅₀ ± SEM (μM)	Hemolysis (%) (mean ± SEM)
9a		1.277 ± 0.083	6.47 ± 0.02
9b		0.371 ± 0.062	11.76 ± 0.03
9c		0.008 ± 0.009	9.41 ± 0.02
9d		0.419 ± 0.162	8.82 ± 0.04
9e		0.025 ± 0.011	6.47 ± 0.02
9f		0.098 ± 0.028	12.94 ± 0.03
9g		0.299 ± 0.081	6.52 ± 0.02
9h		0.622 ± 0.099	2.64 ± 0.04
9i		0.027 ± 0.015	9.41 ± 0.05
9j		0.089 ± 0.077	8.52 ± 0.05
9k		0.044 ± 0.061	10.35 ± 0.03
Kojic acid		16.832 ± 1.162	—
Triton X		—	94.11 ± 0.01

^a SEM = standard error of the mean; values are expressed in mean ± SEM. ^b PBS hemolysis = 0.00 ± 0.01%.

binding interactions (hydrogen/hydrophobic) pattern. Results showed that all compounds showed good binding energy values while binding to the target protein (Table 2). Prior research

showed that the standard error for Autodock is testified as 2.5 kcal mol⁻¹. However, in all docking complexes the predicted energy values difference was less than standard energy value.



Table 2 Docking energy values of docked complexes

Ligands-protein docking complexes	Binding affinity (kcal mol ⁻¹)
9a	-6.9
9b	-8
9c	-7.6
9d	-6.2
9e	-6.6
9f	-8.3
9g	-6.5
9h	-6.6
9i	-8.5
9j	-7.1
9k	-6.8

Although, the basic nucleus of all the synthesized compounds was similar, therefore most of ligands possess good efficient energy values and have no large energy fluctuations difference.

2.6.1 Binding analyses of synthesized compounds against mushroom tyrosinase. The ligands-protein binding analyses showed that **9c** confined in the active binding pocket of target protein as mentioned in Fig. 14. The docking result of **9c**-receptor docked complex showed that hydrogen bonds were observed at Asn-260 residue. The hydrogen atom of **9c** form hydrogen bond against Asn-260 with bond length 2.78 Å. Our docking results showed good correlation with published research which strengthens our work and efficacy. The 2D conformations and binding pose and interactions with binding residues of all the candidate molecules are mentioned in (Fig. S21–S30†).

3. Experimental

3.1 General

All the reagents were purchased from Sigma Aldrich and Alfa Aesar (Germany). Analytical grade solvents were delivered by local

suppliers. Melting points were measured on Griffin and George apparatus by using capillary tubes and were found uncorrected. Initial purity of compound was documented by pre-coated silica gel aluminum plates in thin layer chromatography (TLC). Ethyl acetate and *n*-hexane were used as solvent system in ratio (30 : 70). The spots were recognized by UV₂₅₄. IR peaks were documented on a Jasco-320-A spectrometer by using KBr pellet method. By using BBO probe in Bruker Advance III 600 As-cend spectrometer, signals were recorded in DMSO-*d*₆ of ¹H-NMR (δ , ppm) at 600 MHz and ¹³C-NMR at 150 MHz. Foss Heraeus CHN-O-rapid instrument was used for elemental analyses and theoretical values were within $\pm 0.4\%$. Spectra of EI-MS was obtained by JEOL JMS-600H instrument with data processing system. Value of chemical shift (δ) is given in ppm and coupling constant (*J*) in Hz.

3.2 Synthesis of 5-[(2-amino-1,3-thiazol-4-yl)methyl]-4-phenyl-4*H*-1,2,4-triazole-3-thiol (5)

Synthesis of 5-[(2-amino-1,3-thiazol-4-yl)methyl]-4-phenyl-4*H*-1,2,4-triazole-3-thiol (5). Starting from ethyl 2-(2-amino-1,3-thiazol-4-yl)acetate (1), the compound 5 was synthesized by following the reported protocols by our group.³⁷

3.3 General synthesis of 4-[[5-[(2-amino-1,3-thiazol-4-yl)methyl]-4-phenyl-4*H*-1,2,4-triazol-3-yl]sulfanyl)methyl]-*N*-(aryl)benzamides (9a–k)

N,N-Dimethyl formamide (DMF, 3 mL) was taken in 100 mL round bottom flask and 5-[(2-amino-1,3-thiazol-4-yl)methyl]-4-phenyl-4*H*-1,2,4-triazole-3-thiol (0.2 g, 5) was dissolved in it. Added one pinch of LiH in this solution and mixture was stirred for half an hour at room temperature. After that equimolar amount of different electrophiles, *N*-aryl-4-chlorobenzamide (8a–k, one in each reaction), were added in each respective reaction and stirred for 12–24 hours. The completion of reaction was monitored by TLC. The *n*-hexane and ethyl acetate were used to make a solvent system in the ratio of 70 : 30. Single spot

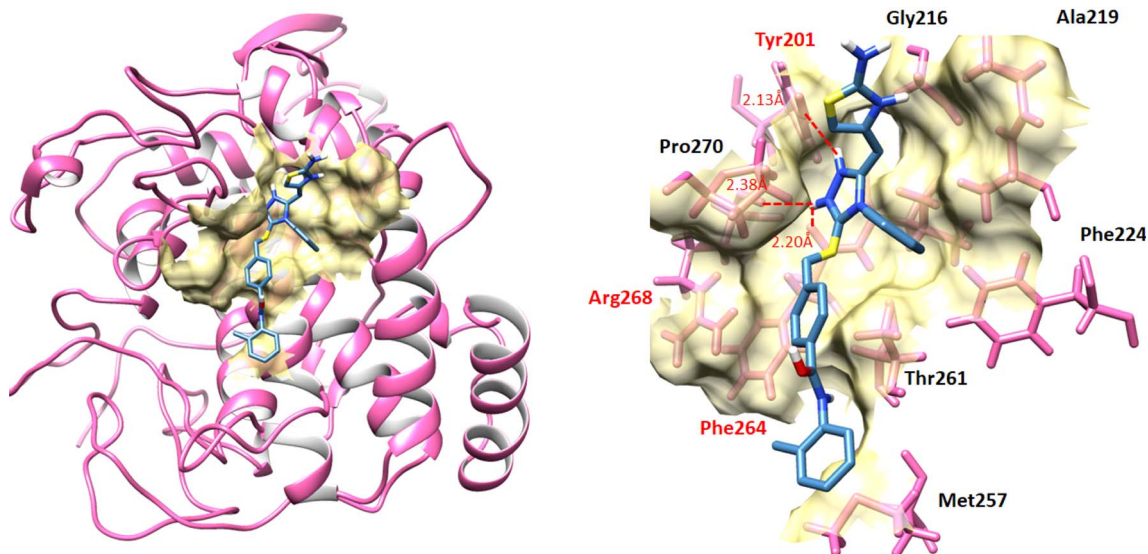


Fig. 14 Binding interaction pattern between **9c** and target protein.



of product was the sign of completion of reaction. The product was precipitated out by adding excess ice-cold distilled water. The targeted products, **9a-k**, were obtained through filtration, excess washing with distilled water and dried respectively for further use.

3.3.1 4-[(5-[(2-Amino-1,3-thiazol-4-yl)methyl]-4-phenyl-4H-1,2,4-triazol-3-yl)sulfanyl)methyl]-N-(2,4-dimethylphenyl)benzamide (9a). Mol. formula: $C_{28}H_{26}N_6OS_2$; mol. mass: 526 g mol⁻¹; IR (KBr, ν/cm^{-1}): 3346 (N-H str.), 3052 (C-H str. of aromatic ring), 2920 (-CH₂- str.), 1670 (C=O str.), 1562 (C=C str. of aromatic ring), 1519 (C=N str.), 1167 (C-N-C bond str.), 621 (C-S str.); ¹H-NMR (600 MHz, DMSO-d₆, δ/ppm): 9.77 (s, 1H, -CO-NH-1'''), 7.88 (br.d, $J = 7.8$, 2H, H-2''' & H-6'''), 7.52-7.47 (m, 3H, H-3'', H-4'' & H-5''), 7.43 (br.d, $J = 7.9$, 2H, H-3''' & H-5'''), 7.22 (br.d, $J = 7.4$, 2H, H-2'' & H-6''), 7.17 (br.d, $J = 7.81$, 1H, H-6'''), 7.07 (br.s, 1H, H-3'''), 7.01 (br.d, $J = 7.8$ 1H, H-5'''), 6.85 (br.s, 2H, H₂N-2), 5.83 (s, 1H, H-5), 4.38 (s, 2H, CH₂-8'''), 3.76 (s, 2H, CH₂-6), 2.28 (s, 3H, CH₃-4'''), 2.17 (s, 3H, CH₃-2'''); ¹³C-NMR (150 MHz, DMSO-d₆, δ/ppm): 168.21 (C-2), 164.88 (C-7'''), 153.50 (C-5'), 149.31 (C-3'), 145.57 (C-4), 140.88 (C-4'''), 135.05 (C-1'''), 133.70 (C-1'''), 133.52 (C-2'''), 133.51 (C-4'''), 132.91 (C-1''), 130.78 (C-5'''), 129.70 (C-4''), 129.47 (C-3'' & C-5''), 128.91 (C-3''' & C-5'''), 127.65 (C-2'' & C-6''), 127.16 (C-2'' & C-6''), 126.52 (C-3'''), 126.46 (C-6'''), 102.19 (C-5), 35.67 (C-8'''), 27.52 (C-6), 20.51 (CH₃-4'''), 17.76 (CH₃-2'''). Anal. calc. for $C_{28}H_{26}N_6OS_2$ (526.67): C, 63.85; H, 4.98; N, 15.96. Found: C, 63.81; H, 4.93; N, 15.92.

3.3.2 4-[(5-[(2-Amino-1,3-thiazol-4-yl)methyl]-4-phenyl-4H-1,2,4-triazol-3-yl)sulfanyl)methyl]-N-(2,5-dimethylphenyl)benzamide (9b). Mol. formula: $C_{28}H_{26}N_6OS_2$; mol. mass.: 526 g mol⁻¹; IR (KBr, ν/cm^{-1}): 3353 (N-H str.), 3050 (C-H str. of aromatic ring), 2926 (-CH₂- str.), 1676 (C=O str.), 1569 (C=C str. of aromatic ring), 1527 (C=N str.), 1165 (C-N-C bond str.), 633 (C-S str.); ¹H-NMR (600 MHz, DMSO-d₆, δ/ppm): 9.80 (s, 1H, -CO-NH-1'''), 7.87 (br.d, $J = 7.8$, 2H, H-2''' & H-6'''), 7.51-7.47 (m, 3H, H-3'', H-4'' & H-5''), 7.44 (br.d, $J = 7.9$, 2H, H-3''' & H-5'''), 7.22 (br.d, $J = 7.0$, 2H, H-2'' & H-6''), 7.15-7.14 (m, 2H, H-3'''' & H-6'''), 6.98 (br.d, $J = 7.5$, 1H, H-4'''), 6.85 (br.s, 2H, H₂N-2), 5.84 (s, 1H, H-5), 4.39 (s, 2H, CH₂-8'''), 3.77 (s, 2H, CH₂-6), 2.28 (s, 3H, CH₃-5'''), 2.17 (s, 3H, CH₃-2'''); ¹³C-NMR (150 MHz, DMSO-d₆, δ/ppm): 168.22 (C-2), 164.90 (C-7'''), 153.52 (C-5'), 149.34 (C-3'), 145.52 (C-4), 140.92 (C-4'''), 136.04 (C-1'''), 135.01 (C-5'''), 133.50 (C-1'''), 132.87 (C-1''), 130.50 (C-2'''), 130.07 (C-3'''), 129.72 (C-4''), 129.48 (C-3'' & C-5''), 128.92 (C-3''' & C-5'''), 127.65 (C-2'' & C-6''), 127.14 (C-2'' & C-6''), 127.06 (C-4'''), 125.28 (C-6'''), 102.23 (C-5), 35.67 (C-8'''), 27.50 (C-6), 20.45 (CH₃-5'''), 17.39 (CH₃-2'''). Anal. calc. for $C_{28}H_{26}N_6OS_2$ (526.67): C, 63.85; H, 4.98; N, 15.96. Found: C, 63.82; H, 4.94; N, 15.95.

3.3.3 4-[(5-[(2-Amino-1,3-thiazol-4-yl)methyl]-4-phenyl-4H-1,2,4-triazol-3-yl)sulfanyl)methyl]-N-(2,6-dimethylphenyl)benzamide (9c). Mol. formula: $C_{28}H_{26}N_6OS_2$; mol. mass.: 526 g mol⁻¹; IR (KBr, ν/cm^{-1}): 3355 (N-H str.), 3060 (C-H str. of aromatic ring), 2928 (-CH₂- str.), 1662 (C=O str.), 1571 (C=C str. of aromatic ring), 1527 (C=N str.), 1174 (C-N-C bond str.), 638 (C-S str.); ¹H-NMR (600 MHz, DMSO-d₆, δ/ppm): 9.75 (s, 1H, -CO-NH-1'''), 7.91 (br.d, $J = 7.6$, 2H, H-2''' & H-6'''), 7.50-7.48 (m, 3H, H-3'', H-4'' & H-5''), 7.45 (br.d, $J = 7.9$, 2H, H-3''' & H-5'''), 7.23 (br.d, $J = 6.7$, 2H, H-2'' & H-6''), 7.13 (br.s, 3H, H-3'''' & H-4'''' &

H-5'''), 6.85 (br.s, 2H, H₂N-2), 5.85 (s, 1H, H-5), 4.39 (s, 2H, CH₂-8'''), 3.77 (s, 2H, CH₂-6), 2.18 (s, 6H, CH₃-2'''' & CH₃-6'''''); ¹³C-NMR (150 MHz, DMSO-d₆, δ/ppm): 168.21 (C-2), 164.61 (C-7'''), 153.52 (C-5'), 149.32 (C-3'), 145.56 (C-4), 140.97 (C-4'''), 135.59 (C-2'''' & C-6'''), 135.21 (C-1'''), 133.29 (C-1'''), 132.90 (C-1''), 129.70 (C-4''), 129.45 (C-3'' & C-5''), 128.97 (C-3''' & C-5'''), 127.67 (C-2'' & C-6''), 127.56 (C-3'''' & C-5'''), 127.15 (C-2'' & C-6''), 126.65 (C-4'''), 102.19 (C-5), 35.67 (C-8'''), 27.52 (C-6), 17.99 (CH₃-2'''' & CH₃-6'''). Anal. calc. for $C_{28}H_{26}N_6OS_2$ (526.67): C, 63.85; H, 4.98; N, 15.96. Found: C, 63.89; H, 4.92; N, 15.98.

3.3.4 4-[(5-[(2-Amino-1,3-thiazol-4-yl)methyl]-4-phenyl-4H-1,2,4-triazol-3-yl)sulfanyl)methyl]-N-(3,4-dimethylphenyl)benzamide (9d). Mol. formula: $C_{28}H_{26}N_6OS_2$; mol. mass.: 526 g mol⁻¹; IR (KBr, ν/cm^{-1}): 3358 (N-H str.), 3040 (C-H str. of aromatic ring), 2912 (-CH₂- str.), 1652 (C=O str.), 1551 (C=C str. of aromatic ring), 1511 (C=N str.), 1149 (C-N-C bond str.), 612 (C-S str.); ¹H-NMR (600 MHz, DMSO-d₆, δ/ppm): 10.05 (s, 1H, -CO-NH-1'''), 7.85 (br.d, $J = 7.6$, 2H, H-2''' & H-6'''), 7.54 (br.s, 1H, H-2'''), 7.50-7.47 (m, 4H, H-3'', H-4'', H-5'' & H-6'''), 7.44 (br.d, $J = 7.8$, 2H, H-3''' & H-5'''), 7.24 (br.d, $J = 6.7$, 2H, H-2'' & H-6''), 7.09 (br.d, $J = 8.0$, 1H, H-5'''), 6.84 (br.s, 2H, H₂N-2), 5.84 (s, 1H, H-5), 4.39 (s, 2H, CH₂-8'''), 3.77 (s, 2H, CH₂-6), 2.22 (s, 3H, CH₃-4'''), 2.19 (s, 3H, CH₃-3'''); ¹³C-NMR (150 MHz, DMSO-d₆, δ/ppm): 168.20 (C-2), 164.81 (C-7'''), 153.49 (C-5'), 149.31 (C-3'), 145.56 (C-4), 140.82 (C-4'''), 136.74 (C-3'''), 136.09 (C-1'''), 133.97 (C-1'''), 132.91 (C-1''), 131.39 (C-4'''), 129.71 (C-4''), 129.48 (C-3'' & C-5''), 128.86 (C-3''' & C-5'''), 127.64 (C-2'' & C-6''), 127.16 (C-2'' & C-6''), 121.60 (C-5'''), 117.90 (C-2'''), 102.20 (C-5), 35.66 (C-8'''), 27.51 (C-6), 19.59 (CH₃-3'''), 18.78 (CH₃-4'''). Anal. calc. for $C_{28}H_{26}N_6OS_2$ (526.67): C, 63.85; H, 4.98; N, 15.96. Found: C, 63.80; H, 4.99; N, 15.99.

3.3.5 4-[(5-[(2-Amino-1,3-thiazol-4-yl)methyl]-4-phenyl-4H-1,2,4-triazol-3-yl)sulfanyl)methyl]-N-(3,5-dimethylphenyl)benzamide (9e). Mol. formula: $C_{28}H_{26}N_6OS_2$; mol. mass.: 526 g mol⁻¹; IR (KBr, ν/cm^{-1}): 3339 (N-H str.), 3040 (C-H str. of aromatic ring), 2934 (-CH₂- str.), 1677 (C=O str.), 1562 (C=C str. of aromatic ring), 1518 (C=N str.), 1165 (C-N-C bond str.), 620 (C-S str.); ¹H-NMR (600 MHz, DMSO-d₆, δ/ppm): 10.00 (s, 1H, -CO-NH-1'''), 7.85 (br.d, $J = 7.6$, 2H, H-2''' & H-6'''), 7.50-7.45 (m, 3H, H-3'', H-4'' & H-5''), 7.45 (br.d, $J = 7.7$, 2H, H-3''' & H-5'''), 7.40 (br.s, 2H, H-2'''' & H-6'''), 7.23 (br.d, $J = 6.8$, 2H, H-2'' & H-6''), 6.85 (br.s, 2H, H₂N-2), 6.75 (br.s, 1H, H-4'''), 5.84 (s, 1H, H-5), 4.39 (s, 2H, CH₂-8'''), 3.77 (s, 2H, CH₂-6), 2.20 (s, 6H, CH₃-3'''' & CH₃-5'''''); ¹³C-NMR (150 MHz, DMSO-d₆, δ/ppm): 168.21 (C-2), 164.97 (C-7'''), 153.50 (C-5'), 149.32 (C-3'), 145.56 (C-4), 140.89 (C-4'''), 138.88 (C-1'''), 137.48 (C-3'''), 133.96 (C-1'''), 132.90 (C-1''), 129.71 (C-4''), 129.48 (C-3'' & C-5''), 128.88 (C-3''' & C-5'''), 127.66 (C-2'' & C-6''), 127.15 (C-2'' & C-6''), 125.16 (C-4'''), 118.11 (C-2'''), 102.21 (C-5), 35.65 (C-8'''), 27.51 (C-6), 21.08 (CH₃-C-3'''' & C-5'''). Anal. calc. for $C_{28}H_{26}N_6OS_2$ (526.67): C, 63.85; H, 4.98; N, 15.96. Found: C, 63.84; H, 4.96; N, 15.98.

3.3.6 4-[(5-[(2-Amino-1,3-thiazol-4-yl)methyl]-4-phenyl-4H-1,2,4-triazol-3-yl)sulfanyl)methyl]-N-(2-methylphenyl)benzamide (9f). Mol. formula: $C_{27}H_{24}N_6OS_2$; mol. mass.: 512 g mol⁻¹; IR (KBr, ν/cm^{-1}): 3361 (N-H str.), 3068 (C-H str. of aromatic ring), 2940 (-CH₂- str.), 1688 (C=O str.), 1541 (C=C str. of aromatic ring), 1531 (C=N str.), 1178 (C-N-C bond str.), 634 (C-



S str.); ¹H-NMR (600 MHz, DMSO-d₆, δ/ppm): 9.86 (s, 1H, -CO-NH-1'''), 7.90 (br.d, *J* = 8.1, 2H, H-2''' & H-6'''), 7.50–7.47 (m, 3H, H-3'', H-4'' & H-5'''), 7.46 (br.d, *J* = 8.2, 2H, H-3''' & H-5'''), 7.34 (br.d, *J* = 7.6, 1H, H-6'''), 7.27 (br.d, *J* = 7.3, 1H, H-3'''), 7.25–7.21 (m, 3H, H-2'', H-6'' & H-5'''), 7.16 (dis.t, *J* = 7.3, 1H, H-4'''), 6.86 (br.s, 2H, H₂N-2), 5.85 (s, 1H, H-5), 4.40 (s, 2H, CH₂-8'''), 3.78 (s, 2H, CH₂-6), 2.23 (s, 3H, CH₃-2'''); ¹³C-NMR (150 MHz, DMSO-d₆, δ/ppm): 168.22 (C-2), 164.89 (C-7'''), 153.52 (C-5'), 149.32 (C-3'), 145.57 (C-4), 140.98 (C-4'''), 133.69 (C-1'''), 133.52 (C-1'''), 133.46 (C-2'''), 132.90 (C-1''), 130.26 (C-3'''), 129.71 (C-4''), 129.47 (C-3'' & C-5''), 128.94 (3''' & C-5'''), 127.70 (C-2'' & C-6'''), 127.16 (C-2'' & C-6''), 126.57 (C-5'''), 125.96 (C-4'''), 125.29 (C-6'''), 102.20 (C-5), 35.66 (C-8'''), 27.52 (C-6), 17.84 (CH₃-2'''). Anal. calc. for C₂₇H₂₄N₆OS₂ (512.64): C, 63.26; H, 4.72; N, 16.39. Found: 63.31; H, 4.70; N, 16.36.

3.3.7 4-[(5-[(2-Amino-1,3-thiazol-4-yl)methyl]-4-phenyl-4*H*-1,2,4-triazol-3-yl)sulfanyl)methyl]-*N*-(3-methylphenyl)benzamide (9g). Mol. formula: C₂₇H₂₄N₆OS₂; mol. mass.: 512 g mol⁻¹; IR (KBr, ν/cm⁻¹): 3341 (N-H str.), 3053 (C-H str. of aromatic ring), 2921 (-CH₂- str.), 1671 (C=O str.), 1562 (C=C str. of aromatic ring), 1516 (C=N str.), 1164 (C-N-C bond str.), 622 (C-S str.); ¹H-NMR (600 MHz, DMSO-d₆, δ/ppm): 10.14 (s, 1H, -CO-NH-1'''), 7.86 (br.d, *J* = 8.1, 2H, H-2''' & H-6'''), 7.60 (br.s, 1H, H-2'''), 7.55 (br.d, *J* = 7.1, 1H, H-6'''), 7.52–7.49 (m, 3H, H-3'', H-4'' & H-5'''), 7.45 (br.d, *J* = 8.2, 2H, H-3''' & H-5'''), 7.24–7.21 (m, 3H, H-2'', H-6'' & H-5'''), 6.92 (br.d, *J* = 7.2, 1H, H-4'''), 6.85 (br.s, 2H, H₂N-2), 5.84 (s, 1H, H-5), 4.39 (s, 2H, CH₂-8'''), 3.77 (s, 2H, CH₂-6), 2.30 (s, 3H, CH₃-3'''); ¹³C-NMR (150 MHz, DMSO-d₆, δ/ppm): 168.21 (C-2), 165.06 (C-7'''), 153.50 (C-5'), 149.32 (C-3'), 145.54 (C-4), 140.94 (C-4'''), 138.95 (C-1'''), 137.71 (C-3'''), 133.91 (C-1'''), 132.89 (C-1''), 129.72 (C-4''), 129.49 (C-3'' & C-5''), 128.89 (C-3''' & C-5'''), 128.39 (C-5'''), 127.68 (C-2'' & C-6'''), 127.15 (C-2'' & C-6''), 124.35 (C-4'''), 120.87 (C-2'''), 117.52 (C-6'''), 102.22 (C-5), 35.64 (C-8'''), 27.50 (C-6), 21.16 (CH₃-3'''). Anal. calc. for C₂₇H₂₄N₆OS₂ (512.64): C, 63.26; H, 4.72; N, 16.39. Found: 63.29; H, 4.77; N, 16.34.

3.3.8 4-[(5-[(2-Amino-1,3-thiazol-4-yl)methyl]-4-phenyl-4*H*-1,2,4-triazol-3-yl)sulfanyl)methyl]-*N*-(4-dimethylphenyl)benzamide (9h). Mol. formula: C₂₇H₂₄N₆OS₂; mol. mass.: 512 g mol⁻¹; IR (KBr, ν/cm⁻¹): 3349 (N-H str.), 3054 (C-H str. of aromatic ring), 2922 (-CH₂- str.), 1672 (C=O str.), 1561 (C=C str. of aromatic ring), 1517 (C=N str.), 1165 (C-N-C bond str.), 623 (C-S str.); ¹H-NMR (600 MHz, DMSO-d₆, δ/ppm): 10.13 (s, 1H, -CO-NH-1'''), 7.85 (br.d, *J* = 8.1, 2H, H-2''' & H-6'''), 7.63 (br.d, *J* = 7.3, 2H, H-2'''' & H-6''''), 7.50–7.47 (m, 3H, H-3'', H-4'' & H-5'''), 7.44 (br.d, *J* = 8.2, 2H, H-3''' & H-5'''), 7.22 (dd, *J* = 1.7, 8.1, 2H, H-2'' & H-6'''), 7.14 (br.d, *J* = 8.2, 2H, H-3''' & H-5'''), 6.84 (br.s, 2H, H₂N-2), 5.83 (s, 1H, H-5), 4.39 (s, 2H, CH₂-8'''), 3.76 (s, 2H, CH₂-6), 2.27 (s, 3H, CH₃-4'''); ¹³C-NMR (150 MHz, DMSO-d₆, δ/ppm): 168.20 (C-2), 164.94 (C-7'''), 153.50 (C-5'), 149.31 (C-3'), 145.56 (C-4), 140.86 (C-4'''), 136.53 (C-1'''), 133.95 (C-1'''), 132.91 (C-1''), 132.59 (C-4'''), 129.71 (C-4''), 129.48 (C-3'' & C-5''), 128.90 (C-3''' & C-5'''), 128.88 (C-3'''), 127.66 (C-2'' & C-6'''), 127.16 (C-2'' & C-6''), 120.20 (C-2'''), 102.20 (C-5), 35.65 (C-8'''), 27.51 (C-6), 20.45 (CH₃-4'''). Anal. calc. for C₂₇H₂₄N₆OS₂ (512.64): C, 63.26; H, 4.72; N, 16.39. Found: 63.23; H, 4.75; N, 16.36.

3.3.9 4-[(5-[(2-Amino-1,3-thiazol-4-yl)methyl]-4-phenyl-4*H*-1,2,4-triazol-3-yl)sulfanyl)methyl]-*N*-(2-ethylphenyl)benzamide (9i). Mol. formula: C₂₈H₂₆N₆OS₂; mol. mass.: 526 g mol⁻¹; IR (KBr, ν/cm⁻¹): 3354 (N-H str.), 3039 (C-H str. of aromatic ring), 2919 (-CH₂- str.), 1680 (C=O str.), 1570 (C=C str. of aromatic ring), 1537 (C=N str.), 1162 (C-N-C bond str.), 625 (C-S str.); ¹H-NMR (600 MHz, DMSO-d₆, δ/ppm): 9.88 (s, 1H, -CO-NH-1'''), 7.90 (br.d, *J* = 8.2, 2H, H-2''' & H-6'''), 7.52–7.49 (m, 4H, H-3'', H-4'', H-5'' & H-6'''), 7.46 (br.d, *J* = 8.2, 2H, H-3''' & H-5'''), 7.31–7.29 (m, 1H, H-3'''), 7.24–7.23 (m, 4H, H-2'', H-6'', H-4'' & H-5'''), 6.86 (br.s, 2H, H₂N-2), 5.85 (s, 1H, H-5), 4.40 (s, 2H, CH₂-8'''), 3.78 (s, 2H, CH₂-6), 2.61 (q, 2H, *J* = 7.5, CH₃ - CH₂ - 2'''), 1.12 (t, 3H, *J* = 7.5, CH₃ - CH₂ - 2'''); ¹³C-NMR (150 MHz, DMSO-d₆, δ/ppm): 168.22 (C-2), 165.27 (C-7'''), 153.52 (C-5'), 149.33 (C-3'), 145.56 (C-4), 140.98 (C-4'''), 139.82 (C-1'''), 135.67 (C-2'''), 133.44 (C-1'''), 132.90 (C-1''), 129.71 (C-4''), 129.47 (C-3'' & C-5''), 128.99 (C-3''' & C-5'''), 128.47 (C-3'''), 127.65 (C-2'' & C-6'''), 127.55 (C-5'''), 127.16 (C-2'' & C-6''), 126.49 (C-4'''), 125.98 (C-6'''), 102.21 (C-5), 35.66 (C-8'''), 27.52 (C-6), 23.96 (CH₃ - CH₂ - 2'''), 14.11 (CH₃ - CH₂ - 2'''). Anal. calc. for C₂₈H₂₆N₆OS₂ (526.67): C, 63.85; H, 4.98; N, 15.96. Found: C, 63.80; H, 4.95; N, 15.93.

3.3.10 4-[(5-[(2-Amino-1,3-thiazol-4-yl)methyl]-4-phenyl-4*H*-1,2,4-triazol-3-yl)sulfanyl)methyl]-*N*-(4-ethylphenyl)benzamide (9j). Mol. formula: C₂₈H₂₆N₆OS₂; mol. mass.: 542 g mol⁻¹; IR (KBr, ν/cm⁻¹): 3344 (N-H str.), 3050 (C-H str. of aromatic ring), 2927 (-CH₂- str.), 1677 (C=O str.), 1567 (C=C str. of aromatic ring), 1526 (C=N str.), 1180 (C-N-C bond str.), 631 (C-S str.); ¹H-NMR (600 MHz, DMSO-d₆, δ/ppm): 9.87 (s, 1H, -CO-NH-1'''), 7.89 (d, *J* = 7.9, 2H, H-2''' & H-6'''), 7.51–7.49 (m, 3H, H-3'', H-4'' & H-5'''), 7.31–7.28 (m, 2H, H-2'''' & H-6''''), 7.24–7.23 (m, 4H, H-2'', H-6'', H-3''' & H-5'''), 6.86 (br.s, 2H, H₂N-2), 5.84 (s, 1H, H-5), 4.39 (s, 2H, CH₂-8'''), 3.77 (s, 2H, CH₂-6), 2.61 (q, 2H, *J* = 7.4, CH₃ - CH₂ - 4'''), 1.12 (t, 3H, *J* = 7.5, CH₃ - CH₂ - 4'''); ¹³C-NMR (150 MHz, DMSO-d₆, δ/ppm): 168.21 (C-2), 165.26 (C-7'''), 153.51 (C-5'), 149.31 (C-3'), 145.56 (C-4), 140.97 (C-4'''), 139.82 (C-4'''), 135.67 (C-1'''), 133.44 (C-1'''), 132.91 (C-1''), 129.70 (C-4''), 129.47 (C-3'' & C-5''), 128.95 (C-3''' & C-5'''), 128.47 (C-3'''), 127.65 (C-2'' & C-6'''), 127.16 (C-2'' & C-6''), 125.98 (C-2'''), 102.20 (C-5), 35.66 (C-8'''), 27.52 (C-6), 23.96 (CH₃ - CH₂ - 4'''), 14.10 (CH₃ - CH₂ - 4'''). Anal. calc. for C₂₈H₂₆N₆O₂S₂ (542.67): C, 63.85; H, 4.98; N, 15.96. Found: C, 63.90; H, 4.96; N, 15.91.

3.3.11 4-[(5-[(2-Amino-1,3-thiazol-4-yl)methyl]-4-phenyl-4*H*-1,2,4-triazol-3-yl)sulfanyl)methyl]-*N*-(3-ethoxyphenyl)benzamide (9k). Mol. formula: C₂₈H₂₆N₆O₂S₂; mol. mass.: 542 g mol⁻¹; IR (KBr, ν/cm⁻¹): 3361 (N-H str.), 3069 (C-H str. of aromatic ring), 2942 (-CH₂- str.), 1684 (C=O str.), 1547 (C=C str. of aromatic ring), 1505 (C=N str.), 1141 (C-N-C bond str.), 608 (C-S str.); ¹H-NMR (600 MHz, DMSO-d₆, δ/ppm): 10.18 (s, 1H, -CO-NH-1'''), 7.95 (br.s, 1H, H-2'''), 7.87 (d, *J* = 8.1, 2H, H-2'' & H-6'''), 7.51–7.49 (m, 3H, H-3'', H-4'' & H-5'''), 7.46 (br.d, *J* = 8.4, 2H, H-3''' & H-5'''), 7.34 (br.d, *J* = 7.9, 1H, H-6'''), 7.24–7.22 (m, 3H, H-2'', H-6'' & H-5'''), 6.66 (dd, *J* = 1.7, 8.1, 1H, H-4'''), 6.85 (br.s, 2H, H₂N-2), 5.84 (s, 1H, H-5), 4.40 (s, 2H, CH₂-8'''), 3.77 (s, 2H, CH₂-6), 4.01 (q, 2H, *J* = 6.9, O - CH₂ - CH₃ - 3'''), 1.34 (t, 3H, *J* = 6.9, O - CH₂ - CH₃ - 3'''); ¹³C-NMR (150 MHz, DMSO-d₆, δ/ppm): 168.22 (C-2), 165.16 (C-7'''), 158.63 (C-3'''),



153.51 (C-5'), 149.32 (C-3'), 145.55 (C-4), 141.01 (C-4'''), 140.24 (C-1'''), 133.89 (C-1'''), 132.90 (C-1''), 129.72 (C-4''), 129.49 (C-3'' & C-5''), 129.32 (C-5'''), 128.90 (C-3''' & C-5'''), 127.71 (C-2''' & C-6'''), 127.15 (C-2'' & C-6''), 112.41 (C-6'''), 109.70 (C-4'''), 106.47 (C-2'''), 102.23 (C-5), 62.91(O - CH₂ - CH₃ - 3'''), 35.64 (C-8'''), 27.51 (C-6), 14.63 (O - CH₂ - CH₃ - 3'''). Anal. calc. for C₂₈H₂₆N₆O₂S₂ (542.67): C, 61.97; H, 4.83; N, 15.49. Found: C, 61.93; H, 4.80; N, 15.44.

3.4 Mushroom tyrosinase inhibition assay

The mushroom tyrosinase (Sigma Chemical, USA) inhibition was performed following our previously reported methods.^{38–40} In detail, 140 μL of phosphate buffer (20 mM, pH 6.8), 20 μL of mushroom tyrosinase (30 U mL⁻¹) and 20 μL of the inhibitor solution were placed in the wells of a 96-well microplate. After pre-incubation for 10 minutes at room temperature, 20 μL of L-DOPA (3,4-dihydroxyphenylalanine, Sigma Chemical, USA) (0.85 mM) was added and the assay plate was further incubated at 25 °C for 20 minutes. Afterward, the absorbance of dopachrome was measured at 475 nm using a microplate reader (OPTI Max, Tunable). Kojic acid was used as a reference inhibitor and phosphate buffer was used as a negative control. The amount of inhibition by the test compounds was expressed as the percentage of concentration necessary to achieve 50% inhibition (IC₅₀). Each concentration was analyzed in three independent experiments. IC₅₀ values were calculated by nonlinear regression using GraphPad Prism 5.0.

The % inhibition of tyrosinase was calculated as following:

$$\text{Inhibition (\%)} = [(B - S)/B] \times 100$$

here, the *B* and *S* are the absorbance's for the blank and samples.

3.5 Kinetic analysis of the inhibition of tyrosinase

Based upon IC₅₀ results, the most potent **9c** compound was designated for the kinetic analysis. A series of experiments were performed to determine the inhibition kinetics of **9c** by following the already reported methods.^{41,42} The inhibitor concentrations for **9c** were 0.00, 0.004, 0.008, 0.016 and 0.032 μM. Substrate L-DOPA concentrations were between 0.125 and 2 mM in all kinetic studies. Pre-incubation and measurement time was the same as discussed in the mushroom tyrosinase inhibition assay protocol. Maximal initial velocity was determined from the initial linear portion of absorbance up to five minutes after addition of enzyme at a 30 s interval. The inhibition type of the enzyme was assayed by Lineweaver–Burk plots of the inverse of velocities (1/*V*) versus the inverse of substrate concentration 1/[L-DOPA] mM⁻¹. The EI dissociation constant *K*_i was determined by the secondary plot of 1/*V* versus inhibitors concentrations.

3.6 Hemolytic activity

Bovine blood sample was collected in EDTA that was diluted with saline (0.9% NaCl), and centrifuge at 1000 × *g* for 10 min. The erythrocytes separated diluted in phosphate buffer saline of pH 7.4 and a suspension was made. Add 20 μL of synthetic compounds

solution (10 mg mL⁻¹) in 180 μL of RBCs suspension and incubate for 30 min at room temperature. PBS was used as negative control and Triton 100-X was taken as positive control.^{43,44} The percentage of hemolysis was taken as by using formula:

$$\begin{aligned} (\%) \text{ of hemolysis} = \\ \frac{\text{absorbance of sample} - \text{absorbance of negative control}}{\text{absorbance of positive control}} \times 100 \end{aligned}$$

3.7 Molecular docking process

The crystal structure of mushroom tyrosinase was accessed from the protein data bank (PDB) having PDBID 2Y9X (<https://www.rcsb.org/>). Energy minimization of mushroom tyrosinase was carried out by using conjugate gradient algorithm and Amber force field in UCSF Chimera 1.10.1.⁴⁵ The stereo-chemical properties, Ramachandran graph and values⁴⁶ of mushroom tyrosinase structure were assessed by Molprobit server,⁴⁷ while the hydrophobicity graph was generated by Discovery Studio 4.1 Client (<https://discover.3ds.com/discovery-studio-visualizer-download>). The protein architecture and statistical percentage values of helices, beta-sheets, coils and turns were accessed by using online tool VADAR 1.8 (<https://vadar.wishartlab.com/>).

4. Conclusions

A structurally distinctive series of novel molecules, amalgamated with a thiazole, a phenyl triazole and a benzamide moiety, was synthesized and recognized with very superb mushroom tyrosinase inhibition as compared to the standard used and most of previously reported inhibitors. It was postulated from their SAR studies that molecules, particularly bearing medium-sized non-polar group with +I effect at *ortho*-position in aryl part of the compound, generally inhibited the mushroom tyrosinase in an excellent manner. All these molecules also showed mild cytotoxicity towards red blood cell membranes. Therefore, it was concluded generally that these hybrid molecules might be pondered as creditable medicinal scaffolds for the treatment of tyrosinase related ailments, particularly skin disorders.

Conflicts of interest

The authors declare no conflict of interest.

Acknowledgements

Special thanks are paid to Higher Education Commission (HEC) of Pakistan for the financial support for this study.

References

- 1 A. M. Abdella, A. M. Abdelmoniem, I. A. Abdelhamid and A. H. M. Elwahy, *J. Heterocycl. Chem.*, 2020, 57, 1476.
- 2 B. D. Vanjare, P. G. Mahajan, N. C. Dige, H. Raza, M. Hassan, Y. Han, S. Ja Kim, S. Yum Seo and Ki. Hwan Lee, *Mol. Diversity*, 2021, 25, 2089.



- 3 A. T. Balaban, D. C. Oniciu and A. R. Katritzky, *Chem. Rev.*, 2004, **104**, 2777.
- 4 N. H. Karam, J. H. Tomma and A. H. Al-Dujaili, *Synthesis*, 2013, **3**, 162.
- 5 T. A. Farghaly, A. M. A. Alnaja, H. A. El-Ghamry and M. R. Shaaban, *Bioorg. Chem.*, 2020, **102**, 104103.
- 6 R. E. Khidre and I. A. M. Radini, *Sci. Rep.*, 2021, **11**, 1.
- 7 J. Rudolph, H. Theis, R. Hanke, R. Endermann, L. Johannsen and F. U. Geschke, *J. Med. Chem.*, 2001, **44**, 619.
- 8 A. Badorc, M. F. Bordes, P. de Cointet, P. Savi, A. Bernat, A. Lalé, M. Petitou, J. P. Maffrand and J. M. Herbert, *J. Med. Chem.*, 1997, **40**, 3393.
- 9 C. I. Lino, I. G. de Souza, B. M. Borelli, T. T. S. Matos, I. N. S. Teixeira, J. P. Ramos and E. M. S. Fagundes, *Eur. J. Med. Chem.*, 2018, **151**, 248.
- 10 G. M. Reddy, J. R. Garcia, V. H. Reddy, A. M. de Andrade, A. Camilo Jr, R. A. P. Ribeiro and S. R. de Lazaro, *Eur. J. Med. Chem.*, 2016, **123**, 508.
- 11 S. Sinha, M. Doble and S. L. Manju, *Eur. J. Med. Chem.*, 2018, **158**, 34.
- 12 S. O. Pember, G. L. Mejia, T. J. Price and R. J. Pasteris, *Bioorg. Med. Chem. Lett.*, 2016, **26**, 2965.
- 13 T. I. de Santana, M. de O. Barbosa, P. A. T. de M. Gomes, A. C. N. da Cruz, T. G. da Silva and A. C. L. Leite, *Eur. J. Med. Chem.*, 2018, **144**, 874.
- 14 K. M. Amin, D. E. A. Rahman and Y. A. Al-Eryani, *Bioorg. Med. Chem.*, 2008, **16**, 5377.
- 15 S. R. Patil, A. Asrondkar, V. Patil, J. N. Sangshetti, F. A. K. Khan, M. G. Damale, R. H. Patil, A. S. Bobade and D. B. Shinde, *Bioorg. Med. Chem. Lett.*, 2017, **27**, 3845.
- 16 R. Sathyanarayana and B. Poojary, *J. Chin. Chem. Soc.*, 2020, **67**, 459.
- 17 J. Lin, S. Zhou, J. X. Xu, W. Q. Yao, G. F. Hao and Y. T. Li, *J. Agric. Food Chem.*, 2020, **68**, 6792.
- 18 S. Akin, E. A. Demir, A. Colak, Y. Kolcuoglu, N. Yildirim and O. Bekircan, *J. Mol. Struct.*, 2019, **1175**, 280.
- 19 M. Mahdavi, A. Ashtari, M. Khoshneviszadeh, S. Ranjbar, A. Dehghani, T. Akbarzadeh, B. Larijani, M. Khoshneviszadeh and M. Saeedi, *Chem. Biodiversity*, 2018, **15**, 1800120.
- 20 Z. Peng, G. Wang, Q. H. Zeng, Y. Li, Y. Wu, H. Liu, J. Jing Wang and Y. Zhao, *Food Chem.*, 2021, **341**, 128265.
- 21 S. Parvez, M. Kang, H. S. Chung and H. Bae, *Phytother. Res.*, 2007, **21**, 805.
- 22 E. A. Demir, A. Colak, A. Kalfa, A. Yasar, O. Bekircan and M. Y. Akatin, *Turk. J. Biochem.*, 2019, **44**, 473.
- 23 A. Goodspeed, L. M. Heiser, J. W. Gray and J. C. Costello, *Mol. Cancer Res.*, 2016, **14**, 3.
- 24 T. Pan, X. Li and J. Jankovic, *Int. J. Cancer*, 2011, **128**, 2251.
- 25 U. Ghani and N. Ullah, *Bioorg. Med. Chem.*, 2010, **18**, 4042.
- 26 T. B. Tumer, F. C. Onder, H. Ipek, T. Gungor, S. Savranoglu, T. T. Tok, A. Celik and M. Ay, *Int. Immunopharmacol.*, 2017, **43**, 129.
- 27 B. A. Bahr, J. Bendiske, Q. B. Brown, S. Munirathinam, E. Caba, M. Rudin, S. Urwyler, A. Sauter and G. Rogers, *Exp. Neurol.*, 2002, **174**, 47.
- 28 D. Y. Peng, Q. Sun, X. L. Zhu, H. Y. Lin, Q. Chen, N. Xi Yu, W. C. Yang and G. Fu Yang, *Bioorg. Med. Chem.*, 2012, **20**, 6739.
- 29 R. Xie, Y. Yao, P. Tang, G. Chen, X. Liu, F. Yun, C. Cheng, X. Wu and Q. Yuan, *Eur. J. Med. Chem.*, 2017, **34**, 1.
- 30 S. Zolghadri, A. Bahrami, M. T. H. Khan, J. Munoz-Munoz, F. G. Molina, F. G. Canovas and A. A. Saboury, *J. Enzyme Inhib. Med. Chem.*, 2019, **34**, 279–309.
- 31 A. R. S. Butt, M. A. Abbasi, S. Z. Siddiqui, H. Raza, M. Hassan, S. A. A. Shah, M. Shahid and S. Y. Seo, *Bioorg. Chem.*, 2019, **86**, 459.
- 32 S. Lee, S. Ullah, C. Park, H. W. Lee, D. Kang, J. Yang, J. Akter, Y. Park, P. Chun and H. R. Moon, *Bioorg. Med. Chem.*, 2019, **27**, 3929.
- 33 U. Ghani, *Bioorg. Chem.*, 2019, **83**, 235.
- 34 W. Xie, J. Zhang, X. Ma, W. Yang, Y. Zhou, X. Tang and Y. Zou, *Chem. Biol. Drug Des.*, 2015, **86**, 1087.
- 35 P. G. Mahajan, N. C. Dige, B. D. Vanjare, H. Raza, M. Hassan, S. Y. Seo, C. H. Kim and K. H. Lee, *J. Mol. Struct.*, 2019, **1198**, 126915.
- 36 B. D. Vanjare, P. G. Mahajan, N. C. Dige, H. Raza, M. Hassan, Y. Han, S. Ja Kim, S. Y. Seo and K. H. Lee, *Mol. Divers.*, 2021, **25**, 2089.
- 37 F. M. Khan, M. A. Abbasi, S. Z. Siddiqui, A. R. S. Butt, H. Raza, A. Zafar, S. A. A. Shah, M. Shahid and S. -Y. Seo, *J. Heterocycl. Chem.*, 2021, **58**, 1089–1103.
- 38 A. R. S. Butt, M. A. Abbasi, S. Z. Siddiqui, H. Raza, M. Hassan, S. A. A. Shah, M. Shahid and S. Y. Seo, *Bioorg. Chem.*, 2019, **86**, 459.
- 39 M. A. Abbasi, Z.-ur Rehman, A.-ur Rehman, S. Z. Siddiqui, M. Nazir, M. Hassan, H. Raza, S. A. A. Shah and S.-Y. Seo, *Acta Chim. Slov.*, 2020, **67**(2), 403.
- 40 A. R. S. Butt, M. A. Abbasi, A.-ur Rehman, S. Z. Siddiqui, H. Raza, M. Hassan, S. A. A. Shah and S.-Y. Seo, *Iran. J. Pharm. Res.*, 2021, **20**(2), 206.
- 41 H. Raza, A. R. S. Butt, M. A. Abbasi, A.-ur Rehman, S. Z. Siddiqui, M. Hassan, S. A. A. Shah and S. J. Kim, *Chem. Biodiversity*, 2023, **20**(2), e202201019.
- 42 M. A. Abbasi, H. Raza, A.-ur Rehman, S. Z. Siddiqui, S. Muhammad, F. M. Khan, S. A. A. Shah, A. G. Al-Sehemi and S. J. Kim, *Chem. Biodiversity*, 2023, **20**(9), e202300257.
- 43 P. Sharma and J. D. Sharma, *J. Ethnopharmacol.*, 2001, **74**, 239.
- 44 W. A. Powell, C. M. Catranis and C. A. Maynard, *Lett. Appl. Microbiol.*, 2000, **31**, 163.
- 45 E. F. Pettersen, T. D. Goddard, C. C. Huang, G. S. Couch, D. M. Greenblatt, E. C. Meng and T. E. Ferrin, *J. Comput. Chem.*, 2004, **25**, 1605.
- 46 V. B. Chen, W. B. Arendall, J. J. Headd, D. A. Keedy, R. M. Immormino, G. J. Kapral, L. W. Murray, J. S. Richardson and D. C. Richardson, *Acta Crystallogr., D*, 2010, **66**, 12.
- 47 S. C. Lovell, I. W. Davis, W. B. Arendall III, P. I. W. De Bakker, J. M. Word, M. G. Prisant, J. S. Richardson and D. C. Richardson, *Proteins: Struct., Funct., Genet.*, 2003, **50**, 437.

