


Cite this: *RSC Adv.*, 2025, 15, 15999

# Multicomponent catalyst-free regioselective synthesis and binding studies of 3-aryl-2-methylimidazo[1,2-*a*]pyrimidines with BSA using biophysical and computational techniques†

Ranjana Aggarwal,<sup>ID</sup> \*<sup>ab</sup> Manisha Sharma,<sup>ID</sup> <sup>a</sup> Garima Sumran<sup>ID</sup> <sup>c</sup> and Parvin Kumar<sup>ID</sup> <sup>a</sup>

A facile and environmentally benign protocol for regioselective synthesis of diversely substituted imidazo[1,2-*a*]pyrimidines **5a–h** has been described *via* multicomponent reaction of unsymmetrical  $\beta$ -diketones **1**, *N*-bromosuccinimide **2** and 2-aminopyrimidine **4** in DCM. The reaction proceeds through *in situ* formation of  $\alpha$ -bromo- $\beta$ -diketones **3** and their ensuing condensation with 2-aminopyrimidine without the need of any organic or inorganic catalyst. The structure of the regioisomeric product was characterized by <sup>1</sup>H, <sup>13</sup>C NMR, heteronuclear 2D NMR and HRMS studies. The present protocol offers several advantages such as avoidance of metal-based and toxic catalysts, broad substrate scope with respect to substitutions on  $\beta$ -diketones, operational simplicity, easy work-up and high yields. Computational molecular docking studies were carried out to examine the interaction of imidazo[1,2-*a*]pyrimidines with bovine serum albumin (BSA). Moreover, different spectroscopic approaches *viz.* UV-visible, steady-state fluorescence and competitive displacement assays were carried out to investigate the binding mechanisms of imidazo[1,2-*a*]pyrimidines (**5c**, **5e** and **5h**) with BSA. The results thus obtained revealed that imidazo[1,2-*a*]pyrimidines showed moderate binding with BSA through a static quenching mechanism and compound **5e** had more affinity to bind in site I of BSA.

Received 12th March 2025  
Accepted 30th April 2025

DOI: 10.1039/d5ra01795e

rsc.li/rsc-advances

## 1. Introduction

Nitrogen containing heterocyclic compounds are constituents of many biologically important molecules, including several endogenous metabolites, natural products and pharmaceuticals.<sup>1,2</sup> Imidazopyrimidines containing a bridgehead nitrogen atom are an important class of fused heterocycles due to their structural resemblance to purine bases present in DNA and RNA.<sup>3</sup> In particular, imidazo[1,2-*a*]pyrimidines are considered as important scaffolds in medicinal and pharmaceutical science, because of their broad spectrum of biological activities such as cytotoxic,<sup>4</sup> antimicrobial,<sup>5</sup> anti-inflammatory,<sup>6</sup> antiviral,<sup>7</sup> anti-tubercular,<sup>8</sup> antileishmanial,<sup>9</sup> and antimalarial activity.<sup>10</sup> Imidazo[1,2-*a*]pyrimidine derivatives also acted as acetylcholinesterase inhibitors<sup>11</sup> for Alzheimer's disease

treatment,  $\alpha$ -glucosidase inhibitors to treat diabetes mellitus,<sup>12</sup> dual inhibitors of both ACE2 and spike protein to prevent SARS-CoV-2 infection,<sup>13</sup> GABA receptor ligands for treatment of anxiety disorders,<sup>14</sup> and p38 MAP kinase inhibitors for treatment of inflammatory conditions such as rheumatoid arthritis and Crohn's disease.<sup>15</sup> Additionally, a photophysical study of imidazo[1,2-*a*]pyrimidines demonstrated that these derivatives serve as photosensitizers in intracellular environments that generate singlet oxygen upon irradiation for killing cancer cells,<sup>16</sup> and act as a fluorescent chemo sensor to detect zinc ions,<sup>17</sup> and an electrochemical study showed that imidazo[1,2-*a*]pyrimidines are proven effective corrosion inhibitors for copper.<sup>18</sup> Moreover, many drugs containing imidazo[1,2-*a*]pyrimidine scaffold as a core unit such as faspiron, taniplon, and divaplon are available in the market (Fig. 1).

Serum albumins, major constituent proteins of blood, are a type of biological receptor that are intricate in transport and distribution of endogenous and exogenous chemicals including thyroid hormones, metals, fatty acids, steroids, and other molecules throughout the bloodstream.<sup>19</sup> Serum albumins also play a crucial role in maintaining the extracellular fluid by influencing the plasma oncotic pressure.<sup>20</sup> The drug–protein interaction significantly influences pharmacokinetic properties such as absorption, delivery, metabolism, and excretion

<sup>a</sup>Department of Chemistry, Kurukshetra University, Kurukshetra-136119, Haryana, India

<sup>b</sup>Council of Scientific and Industrial Research-National Institute of Science Communication and Policy Research, New Delhi 110012, India. E-mail: ranjana67in@yahoo.com; ranjanaaggarwal67@gmail.com; Tel: +91-9896740740

<sup>c</sup>Department of Chemistry, D. A. V. College (Lahore), Ambala City, Haryana 134 003, India

† Electronic supplementary information (ESI) available. See DOI: <https://doi.org/10.1039/d5ra01795e>

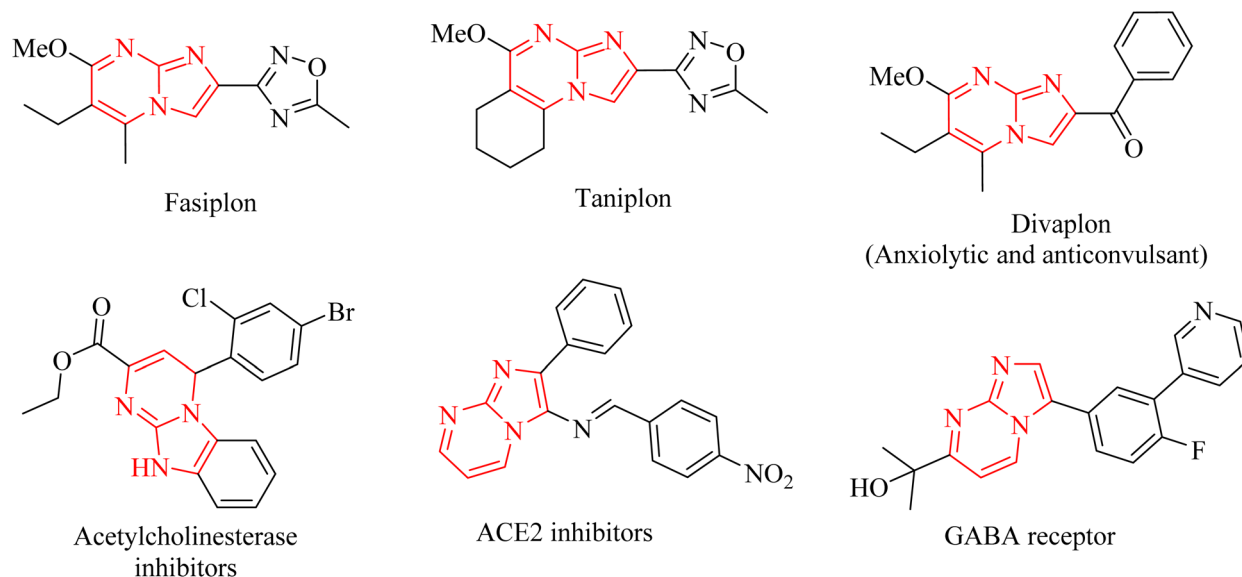



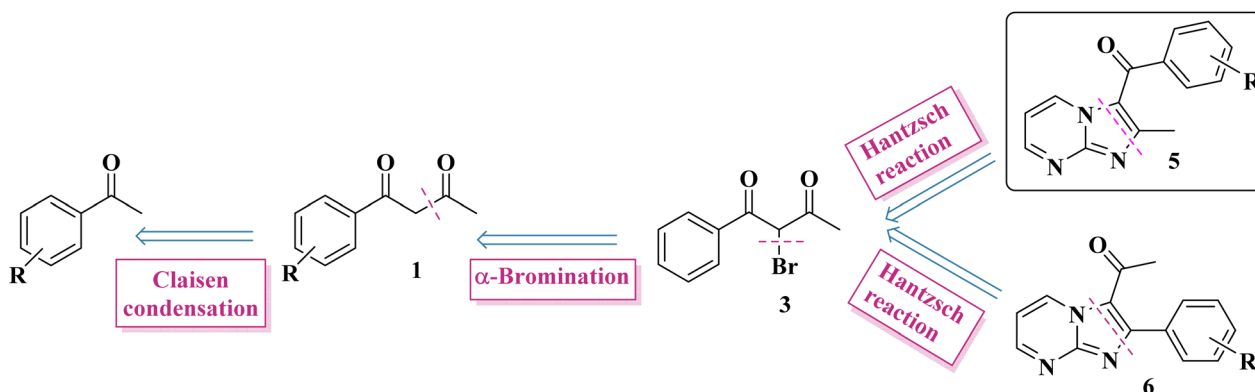
Fig. 1 Representative molecules containing imidazo[1,2-*a*]pyrimidine scaffold.

(ADME) and pharmacodynamics properties of drugs.<sup>21</sup> In order to achieve the desired therapeutic effect, drugs must bind to serum albumin with optimum binding affinity. Therefore, investigation of the molecular interactions between proteins and small organic molecules and drugs is very crucial to design and develop novel and more efficient drugs with improved plasma solubility/stability.

Bovine serum albumin (BSA) is one of the most extensively researched proteins as primary molecular targets due to its capability to reversibly bind to a wide variety of drugs.<sup>22,23</sup> BSA is more widely used as a crucial model protein to study the interaction of small molecules with serum albumins due to its cost effectiveness, wide availability, good stability and 76% structural homology with Human serum albumin (HSA).<sup>24</sup> BSA's primary structure contains 583 amino acids with 17 disulfide bonds and a free cysteine residue.<sup>25</sup> It has three homologous  $\alpha$ -helical domains (I–III), each containing two subdomains A and B.<sup>26</sup> BSA exhibits intrinsic fluorescence specifically due to Trp134 and Trp213 residues which are located on the surface of

subdomain IB and in the hydrophobic cavity of subdomain IIA, respectively<sup>27</sup> and the quenching of this fluorescence by small molecules is used in BSA-ligand interaction studies.

In view of their biological potential, to date different synthetic strategies have been developed for the construction of functionalized imidazo[1,2-*a*]pyrimidine derivatives.<sup>28</sup> The traditional and most commonly used method for the synthesis of imidazo[1,2-*a*]pyrimidines is Hantzsch-type synthesis which involves the cyclo-condensation of  $\alpha$ -functionalized carbonyl compounds with 2-aminopyrimidine in the presence of various catalysts such as neutral  $\text{Al}_2\text{O}_3$ ,<sup>29</sup> KI,  $\text{K}_2\text{CO}_3$ ,<sup>30</sup> PTSA,<sup>31</sup>  $\text{NaHCO}_3$ ,<sup>32</sup> *etc.* For  $\alpha$ -bromination of 1,3-diketones, *N*-bromosuccinimide (NBS) was utilized over other reagents *e.g.* molecular bromine, sodium hypobromite, tetraalkylammonium tribromide,  $\text{CuBr}_2$  with Koser's reagent,  $\text{HBr-H}_2\text{O}_2$ , due to its easily accessible, easy to handle, inexpensive, and selective brominating property.<sup>33</sup> In recent years, three component reactions of aldehydes, 2-aminopyrimidine and nitriles/isonitriles/alkyne or their derivatives have been reported to produce imidazo[1,2-*a*]



Scheme 1 Retrosynthetic approach for the construction of imidazo[1,2-*a*]pyrimidine.



pyrimidines.<sup>17,28</sup> Other method involves the reaction between 2-aminopyrimidine, ketone and [hydroxy(tosyloxy)iodo]benzene (HTIB) in one pot.<sup>34</sup> More recently, Chanda *et al.*<sup>35</sup> described water and isopropanol mediated catalyst-free microwave assisted protocol for the synthesis of imidazo[1,2-*a*]pyrimidines which shows progress towards green synthetic methods. However, most of these methods often suffer from a drawback of harsh reaction conditions calling for high temperature, hazardous high boiling organic solvents, long reaction times, decomposition of substrates during overheating and lower yields of the products. The necessity of thermal activation for the imidazo[1,2-*a*]pyrimidines ring closure is one of the main and most disappointing limitations on the way to medicinal utilization of this heterocyclic motif. Therefore, the development of energy and environment-efficient greener synthetic methodologies for the construction of these versatile bicyclic heterocycles is always in demand.

Synthetic strategy for functionalized imidazo[1,2-*a*]pyrimidine through a classical Hantzsch reaction between the  $\alpha$ -bromo-1,3-diketones and 2-aminopyrimidine is shown in Scheme 1. The reaction of unsymmetrical diketones **1** with 2-aminopyrimidine **4** in the presence of NBS may lead to the formation of two possible regioisomers: 3-aryl-2-methyl-imidazo[1,2-*a*]pyrimidines **5** and 3-acetyl-2-aryl-imidazo[1,2-*a*]pyrimidines **6**, by different permutations, as depicted in Scheme 1. Differently substituted acetophenones on reaction with ethyl acetate in the presence of sodium ethoxide were converted into corresponding unsymmetrical 1,3-diketones through Claisen condensation following the literature protocol.

Recently, multicomponent reactions (MCRs) have gained considerable interest as a sustainable approach offering several advantages such as efficiency, selectivity, high atom economy, reduced waste generation and short reaction time.<sup>36</sup> Currently our research group is exploring regioselective reaction of substituted unsymmetrical  $\alpha$ -bromo-1,3-diketones with various binucleophiles,<sup>37</sup> therefore, it was planned to examine study of regioselectivity pattern of the reaction between  $\alpha$ -bromo-1,3-diketones and 2-aminopyrimidine to construct the functionalized imidazo[1,2-*a*]pyrimidine scaffold using MCR approach. To the best of our knowledge, the reaction between 2-aminopyrimidine and unsymmetrical  $\alpha$ -bromo-1,3-diketones has not been proposed yet. Intrigued by these findings and in continuation of our ongoing efforts towards the synthesis of fused nitrogen-bridged heterocyclic compounds using green synthetic protocols,<sup>37–39</sup> herein we describe a highly efficient, catalyst-free regioselective approach for the synthesis of functionalized imidazo[1,2-*a*]pyrimidine derivatives involving *in situ* bromination of unsymmetrical 1,3-diketones by NBS and condensation with 2-aminopyrimidine under mild reaction condition and characterization of regioisomer on the basis of heteronuclear 2D NMR [<sup>1</sup>H–<sup>13</sup>C] heteronuclear multiple bond correlation, HMBC, [<sup>1</sup>H–<sup>13</sup>C] heteronuclear multiple quantum coherence, HSQC, [<sup>1</sup>H–<sup>15</sup>N] HMBC] spectroscopic studies. The molecular docking studies have been executed to determine interaction mode of imidazo[1,2-*a*]pyrimidine derivatives with BSA. Binding studies of representative compounds with BSA

have been investigated using UV-visible absorption spectroscopy and fluorescence emission spectroscopy.

## 2. Result and discussion

### 2.1 Chemistry

In order to synthesize imidazo[1,2-*a*]pyrimidines, a model reaction was carried out by stirring an equimolar amount of 1-phenylbutane-1,3-dione **1** with NBS **2** to form 2-bromo-1-phenylbutane-1,3-dione **3** *in situ* and then followed by the subsequent addition of 2-aminopyrimidine **4** in acetone at room temperature. Thin layer chromatography (TLC) examination of the reaction mixture denoted that the reaction was complete within 24 h, resulting in the formation of a single regioisomer out of two possible regioisomers with only 25% yield (entry 1, Table 1).

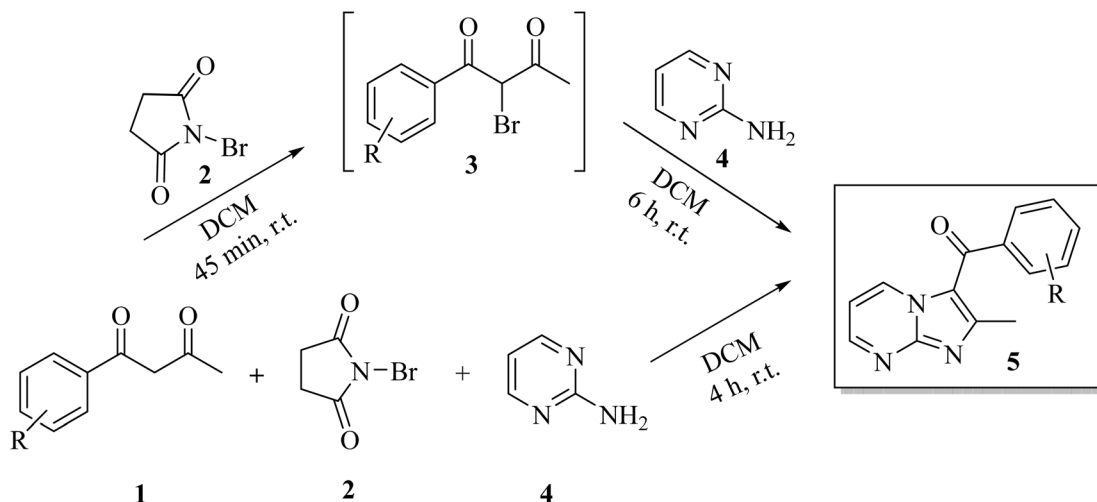
Further to optimize the reaction condition, we explored the model reaction in different polar protic (MeOH, EtOH, AcOH); polar aprotic (DCM, chloroform, acetone, MeCN) and nonpolar aprotic (toluene) solvents at different temperatures (room temperature up to reflux) to determine the most appropriate reaction conditions (entry 2–8, Table 1). Among the solvents screened, DCM improved the yield significantly to 75% at room temperature (entry 5, Table 1). To further improve the yield, reaction was also carried out in DCM by the gradual increase of temperature from r.t. to refluxing temperature. However, there was no further improvisation of the product yield. In addition, the reaction was performed at r.t. under solvent-free conditions which furnished the desired product in trace amount (entry 9, Table 1). However, on heating under solvent-free conditions, the desired product was obtained but only with 60% yield (entry 10, Table 1).

Having optimized conditions identified for sequential synthesis, we explored further one-pot multicomponent domino process by stirring an equimolar mixture of 2-aminopyrimidine **4**, 1-phenylbutane-1,3-dione **1** and NBS **2** in DCM at r.t. The reaction proceeded smoothly to yield the single product

**Table 1** Screening of solvents and optimization of reaction temperature for synthesis of imidazo[1,2-*a*]pyrimidine from 2-bromo-1-phenylbutane-1,3-dione **3** and 2-aminopyrimidine **4**

Entry	Solvent <sup>a</sup>	Temperature <sup>a</sup>	Time	Yield <sup>b</sup> (%)
1	Acetone	r.t.	24 h	25
2	MeOH	Reflux	6 h	45
3	EtOH	Reflux	6 h	52
4	AcOH	Reflux	4 h	48
5	DCM	r.t.	6 h 45 min	75
6	DCM	Reflux	2.5 h	65
7	MeCN	Reflux	6 h	55
8	Toluene	Reflux	5 h	NR <sup>c</sup>
9	Solvent-free	r.t.	3 h	Trace
10	Solvent-free	90 °C	1.5 h	60
11	DCM <sup>d</sup>	r.t.	4 h	80

<sup>a</sup> Reaction conditions: unsymmetrical  $\beta$ -diketone (**1a**, 1 mmol), NBS (**2**, 1 mmol), 2-aminopyrimidine (**4**, 1 mmol) and an appropriate solvent (10.0 mL). <sup>b</sup> Isolated yield. <sup>c</sup> NR, no reaction. <sup>d</sup> Domino reaction.



Scheme 2 Sequential and multicomponent regioselective synthesis of 3-arylimidazo[1,2-*a*]pyrimidines.

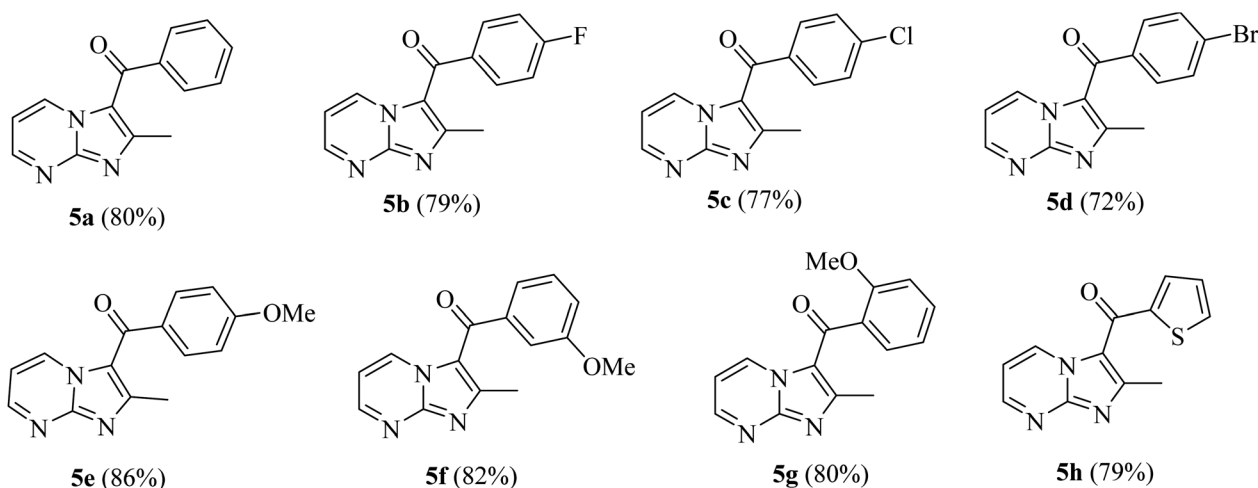
in 4 h as indicated by TLC out of two possible regioisomers. It is noteworthy to mention that this domino approach increased the yield (entry 11 vs. 5) and the product was isolated in 80% yield in short reaction time (4 h vs. 6 h 45, entry 11 vs. 5) than the sequential synthesis protocol. Hence, this one-pot MCRs approach was chosen as the model method (Scheme 2).

To get an insight into the extensibility and efficiency of the developed protocol, differently substituted unsymmetrical  $\beta$ -diketones containing electron donating and electron withdrawing substituents were reacted with 2-aminopyrimidine under optimized conditions. All the reaction combinations underwent successful condensation and yielded the corresponding products in high yields with high regioselectivity. Generally,  $\beta$ -diketones bearing electron-donating groups provided better yields of products than  $\beta$ -diketones bearing

electron-withdrawing groups. Differently synthesized imidazo[1,2-*a*]pyrimidines are outlined in Table 2.

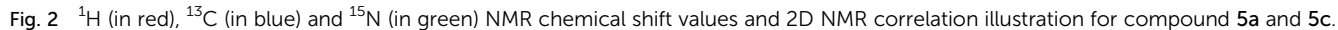
The successful conversion of reactants into products was confirmed through spectroscopic ( $^1\text{H}$ -NMR,  $^{13}\text{C}$ -NMR, IR) data analysis. The IR spectrum of the attained product **5a** displayed a sharp absorption band at  $1648\text{ cm}^{-1}$  corresponding to carbonyl group stretch indicating the product formation. The  $^1\text{H}$  NMR spectrum of compound **5a** displayed a singlet of three proton intensity at  $\delta$  2.21 ppm corresponding to the methyl group, three signals each as doublet of doublet of one proton intensity at 7.62, 9.02 and 9.65 ppm attributable to H-6, H-5, H-7 protons of pyrimidine nucleus, respectively, and three signals for two, one and two proton intensity at 7.62, 7.73, 7.80 ppm corresponding to phenyl protons.  $^{13}\text{C}$  NMR spectrum of compound **5a** displayed signals at  $\delta$  15.44 and 186.20 ppm

Table 2 Substrate scope for unsymmetrical  $\beta$ -diketones<sup>a,b</sup>



<sup>a</sup> Reaction conditions:  $\beta$ -diketones **1** (1.0 mmol), NBS **2** (1.0 mmol) and 2-aminopyrimidine **4** (1.0 mmol) were stirred in DCM at room temperature for 4–5 h. <sup>b</sup> Isolated yields.





with  $\alpha$ -bromo- $\beta$ -diketones to give the desired product **5a**. Additionally, the mass spectra of **5a** exhibited a peak at  $m/z = 238.0890$ , representing the molecular ion,  $[\text{M} + \text{H}]^+$ , which

Chemical shifts ( $\delta$ in ppm)	gs-HMQC correlation	gs-HMBC correlation	Assignments
186.20	—	7.80 (H2'/H6')	CO
156.73	9.02 (5H)	7.62 (H6) 9.65 (H7) 146.82 (C9)	C5
147.60	—	2.21 (2-CH <sub>3</sub> )	C2
146.82	—	9.02 (H5) 9.65 (H7)	C9
138.81	—	7.62 (H3'/H5')	C1'
138.07	9.65 (7H)	7.62 (H6) 9.02 (H5) 146.82 (C9)	C7
133.52	7.73 (H4')	7.62 (H3'/H5')	C4'
129.38	7.62 (H3'/H5')	138.81 (C1')	C3'/C5'
129.27	7.80 (H2'/H6')	7.73 (H4') 186.20 (CO)	C2'/C6'
119.67	—	2.21(2-CH <sub>3</sub> )	C3
113.66	7.62 (6H)	9.65 (7H) 9.02 (5H)	C6
15.44	2.21 (2-CH <sub>3</sub> )	C2 (147.60) C3 (119.67)	2-CH <sub>3</sub>

aligns with the expected composition for the title compound ( $C_{14}H_{11}N_3O$ ).

In order to unambiguously assign the correct regioisomeric structure to reaction product as **5** out of **5** and **6**, multinuclear 2D NMR experiments [ $(^1H-^{13}C)$  HMBC,  $(^1H-^{13}C)$  HSQC and  $(^1H-^{15}N)$  HMBC] were carried out for compound **5a**. The 2D NMR correlation results and  $^1H$ ,  $^{13}C$  and  $^{15}N$  chemical shifts for compound **5a** are shown in Fig. 2 and Table 3.

The  $(^1H-^{13}C)$  HMBC spectrum of **5a** displayed cross peak of carbonyl carbon at  $\delta$  186.20 ppm with H2'/H6' ( $\delta$  7.80) protons of aryl ring confirming the presence of CO with aryl/heteroaryl ring. Similarly  $(^1H-^{13}C)$  HMBC spectrum showed cross peaks of methyl protons ( $\delta$  2.21 ppm) with C-2 ( $\delta$  147.60 ppm) and C-3 ( $\delta$  119.67 ppm) which indicated the presence of methyl substituent at position-2 of imidazo[1,2-*a*]pyrimidine nucleus. Further,  $(^1H-^{15}N)$  HMBC of compound **5a** also showed a cross peak of methyl protons ( $\delta$  2.21) with N-1 ( $\delta$  -215.9), thus confirming the presence of methyl substituent at position-2 of imidazo[1,2-*a*]pyrimidinyl scaffold. Had the structure been 3-acetyl-2-arylimidazo[1,2-*a*]pyrimidine **6a**, then the correlation between the methyl protons with N-1, C-3 and the cross peak of carbonyl carbon with the H2'/H6' proton of the aryl part would have been absent. Thus, the structure can certainly be assigned as 1-phenyl-1-(2-methylimidazo[1,2-*a*]pyrimidin-3-yl) methanone **5a**. Similar correlation results of  $(^1H-^{13}C)$  HMBC and  $(^1H-^{15}N)$  HMBC were observed for the compound **5c** as shown in Fig. 2.

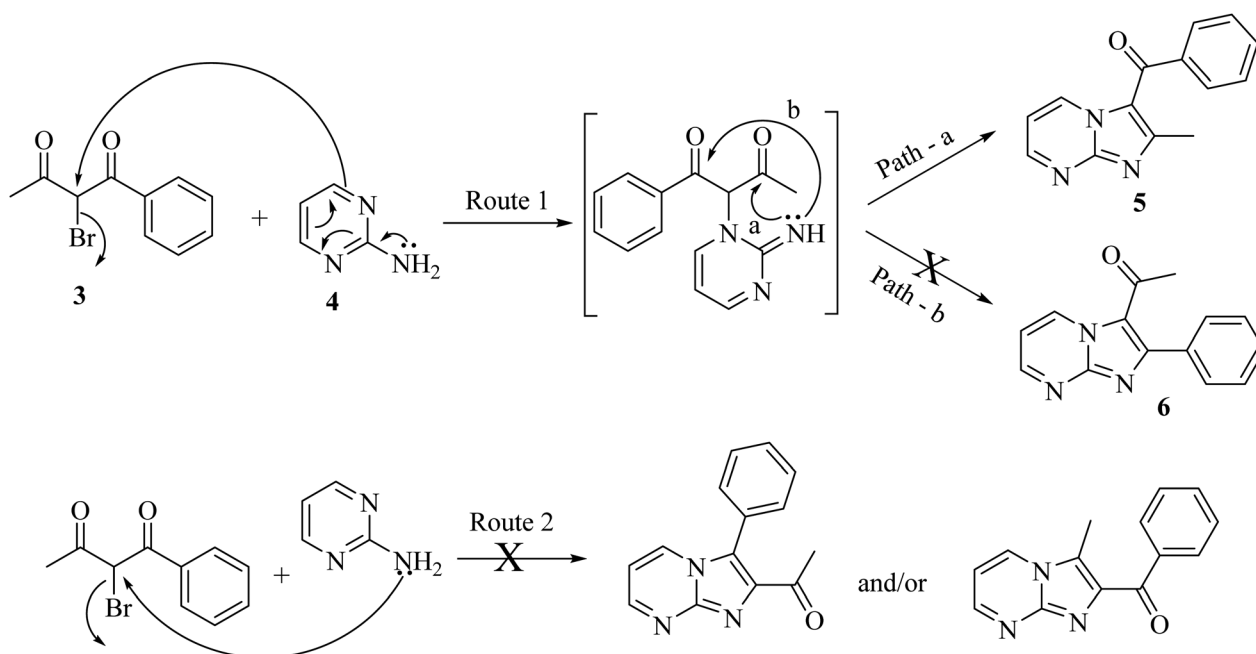
The plausible mechanism for regioselective synthesis of imidazo[1,2-*a*]pyrimidine is depicted in Scheme 3. In the first step, ring nitrogen of 2-aminopyrimidine attacks nucleophilically on  $\alpha$ -bromo-1,3-diketones to give the intermediate that is cyclized by nucleophilic addition of imine nitrogen to either of the carbonyl carbons to form regioisomer 2-alkyl/arylimidazo

[1,2-*a*]pyrimidine **5** or **6** (Route 1). Due to more electrophilic character and less steric hindrance on carbonyl carbon adjacent to methyl, imine nitrogen underwent nucleophilic addition on this carbonyl carbon (Path-a) followed by removal of water to give 2-alkylimidazo[1,2-*a*]pyrimidine **5** as the final product.

## 2.2 Biological studies

**2.2.1. Molecular docking studies.** Molecular docking is a computational tool that predicts the conformation and orientation of organic molecules within the binding site of biomolecular targets (protein, nucleic acids, *etc.*) by studying the root mean square deviances ranging from 1.5 to 2.0 Å, depending on the experimental poses. All novel synthesized imidazo[1,2-*a*]pyrimidine derivatives **5a–h** and phenylbutazone (PBZ) and ibuprofen (IBP) as site-I and site-II markers, respectively, were studied by *in silico* molecular docking studies in order to better study the diverse non-covalent molecular interaction, calculation of binding free energy, and figuring out the appropriate binding site between these compounds with BSA.

Outcomes from molecular docking simulation studies showed that imidazo[1,2-*a*]pyrimidine derivatives **5a–h** exhibited binding interactions with the BSA protein mainly through hydrophobic interaction as well as through hydrogen bonding, electrostatic and van der Waals interaction with binding free energy in the range of  $-8.25$  to  $-9.04$  kcal mol $^{-1}$  (Table 4). Among the synthesized imidazo[1,2-*a*]pyrimidine derivatives, compound **5e** with 4-methoxy substitution at phenyl ring binds with the albumin protein (BSA, PDB ID: 4f5s) in AutoDock more efficiently than other derivatives with a maximum docking score of  $-9.04$  kcal mol $^{-1}$  (Table 4). Results of docking analysis using BIOVIA Discovery Studio Visualizer (DSV) have shown that **5e** interacts with BSA in the active pocket of chain A. Compound **5e**



Scheme 3 Plausible mechanism for regioselective synthesis of imidazo[1,2-*a*]pyrimidine.



Table 4 Receptor–ligand interactions between binding pocket of BSA (PDB ID: 4f5s) and imidazo[1,2-*a*]pyrimidines

Compounds	Binding energy (kcal mol <sup>-1</sup> )	Interacted residues
5a	−8.69	Tyr160 <sup>a,b,c,d</sup> , Tyr137 <sup>b,c</sup> , Arg185 <sup>e</sup> , Leu115 <sup>e</sup> , Ile181 <sup>d,e</sup> , Pro117 <sup>d</sup> , Met184 <sup>e</sup> , Val188 <sup>e</sup> , Lys136 <sup>f</sup> , Glu140 <sup>f</sup> , Ile141 <sup>f</sup> , Leu122 <sup>f</sup>
5b	−8.51	Tyr160 <sup>a,d</sup> , Tyr137 <sup>b,g</sup> , Ile181 <sup>d</sup> , Met184 <sup>d,h</sup> , Arg185 <sup>d,e</sup> , Val188 <sup>e</sup> , Pro117 <sup>e</sup> , Leu115 <sup>e</sup> , Ile141 <sup>i</sup> , Leu189 <sup>f</sup>
5c	−8.50	Tyr137 <sup>a,b</sup> , Met184 <sup>c,h</sup> , Arg185 <sup>k,e</sup> , Val188 <sup>e</sup> , Ile181 <sup>e</sup> , Leu115 <sup>e</sup> , Pro117 <sup>e</sup> , Lys114 <sup>e</sup> , Leu122 <sup>f</sup> , Tyr160 <sup>f</sup> , Ile141 <sup>f</sup>
5d	−8.95	Tyr160 <sup>b,d</sup> , Met184 <sup>d</sup> , Arg185 <sup>d,e,g,i</sup> , Ile181 <sup>d,e</sup> , Pro117 <sup>e</sup> , Leu115 <sup>e</sup> , Tyr137 <sup>f</sup> , Ile141 <sup>f</sup> , Val188 <sup>f</sup> , Glu182 <sup>f</sup> , Glu140 <sup>f</sup> , Lys136 <sup>f</sup>
5e	−9.04	Tyr137 <sup>a,b</sup> , Lys136 <sup>d</sup> , Pro117 <sup>d</sup> , Tyr160 <sup>b,d</sup> , Met184 <sup>e</sup> , Ile181 <sup>d,e</sup> , Arg185 <sup>e,g</sup> , Val188 <sup>e</sup> , Leu115 <sup>e</sup> , Phe133 <sup>j</sup>
5f	−8.76	Tyr160 <sup>a,d</sup> , Tyr137 <sup>c</sup> , Met184 <sup>d</sup> , Ile181 <sup>d,e</sup> , Arg185 <sup>d,e,g,k</sup> , Pro117 <sup>e</sup> , Glu140 <sup>f</sup> , Lys136 <sup>f</sup> , Leu115 <sup>d,i</sup> , Leu122 <sup>f</sup> , Ile141 <sup>f</sup> , Glu182 <sup>f</sup>
5g	−8.28	Tyr160 <sup>a,c,d</sup> , Tyr137 <sup>a,c,d</sup> , Val188 <sup>e</sup> , Met184 <sup>e</sup> , Arg185 <sup>e,g</sup> , Ile181 <sup>d,e</sup> , Ile141 <sup>d</sup> , Pro117 <sup>d,e</sup> , Leu115 <sup>d,i</sup> , Leu122 <sup>f</sup> , Glu140 <sup>f</sup>
5h	−8.25	Tyr160 <sup>b,g,h,l</sup> , Arg185 <sup>e,g</sup> , Ile181 <sup>d,e</sup> , Val188 <sup>e</sup> , Tyr137 <sup>b,h</sup> , Met184 <sup>h</sup> , Leu115 <sup>e</sup> , Pro117 <sup>d,e</sup>
PBZ	−8.81	Tyr137 <sup>a</sup> , Tyr160 <sup>a</sup> , Pro117 <sup>d</sup> , Ile181 <sup>d</sup> , Val188 <sup>e</sup> , Arg185 <sup>e</sup> , Glu140 <sup>f</sup> , Met184 <sup>f</sup> , Leu178 <sup>f</sup> , Glu182 <sup>f</sup> , Leu115 <sup>i</sup> , Ile141 <sup>i</sup>
IBP	−7.46	Leu115 <sup>d,e</sup> , Ile181 <sup>d</sup> , Arg185 <sup>d,g,k</sup> , Lys114 <sup>d,j</sup> , Pro117 <sup>e</sup> , Glu182 <sup>f</sup> , Tyr160 <sup>f</sup> , Tyr137 <sup>f</sup> , Arg144 <sup>f</sup>

<sup>a</sup>  $\pi$ -donor hydrogen bonding. <sup>b</sup>  $\pi$ - $\pi$  T-shaped. <sup>c</sup>  $\pi$ - $\pi$  stacked. <sup>d</sup> Alkyl. <sup>e</sup>  $\pi$ -alkyl. <sup>f</sup> van der Waals. <sup>g</sup> hydrogen bonding. <sup>h</sup>  $\pi$ -sulfur. <sup>i</sup>  $\pi$ -sigma. <sup>j</sup> Carbon–hydrogen bond. <sup>k</sup>  $\pi$ -cation. <sup>l</sup> Amide  $\pi$ -stacking.

interacts with various amino acids (Tyr137, Lys136, Pro117, Tyr160, Met184, Ile181, Val188, Leu115, Phe133) through hydrogen bonding and various hydrophobic interactions as shown in the 2D and 3D plots (Fig. 3). Also, compound 5e formed a conventional hydrogen bond with Arg185 via its carbonyl group. The lowest binding free energy conformer of compounds 5a–h, PBZ and IBP with BSA is shown in Fig. 4a–j.

Docking analysis showed that almost all the derivatives exhibit strong interactions with the A chain of the protein with few exceptions where derivatives 5a and 5b were found to interact with the protein through chain B. Compound 5e was selected for further *in vitro* analysis with BSA protein using various spectroscopic techniques due to the best docking score. To compare the results, two compounds bearing substituents of different nature, 5c with 4-chlorophenyl moiety and 5h with 2-thienyl group, were chosen.

### 2.2.2. Binding studies of imidazo[1,2-*a*]pyrimidines with BSA

**2.2.2.1 UV-visible absorption spectroscopy studies.** Recently UV-visible spectroscopy has been a valuable and imperative tool for studying the conformational changes of proteins during

interactions between drugs and proteins due to its good sensitivity, convenience of usage and versatility.<sup>40</sup> Furthermore, this technique can also be used to investigate the type of quenching (static/dynamic) between ligand and protein. When BSA interacts with ligands, the biomolecule's electronic environment is perturbed, which results in a wavelength shift or alteration in the intensity in UV-visible spectra.<sup>41</sup> During static quenching, a significant change in absorption spectra is observed on subsequent addition of ligand, while the absorption spectra remain unchanged in dynamic quenching.

Among the synthesized compounds, 5e having electron donating 4-OMePh ring, 5h having 2-thienyl ring and 5c having electron withdrawing 4-ClPh ring were selected for UV-visible spectral studies with BSA. Absorption spectra of BSA were recorded in the absence and presence of 5e, 5h, and 5c at a varying concentration ranging from 0 to 40, 0 to 28 and 0 to 36  $\mu$ M, respectively, in the wavelength range of 250–350 nm at room temperature. The UV-visible absorption spectra of BSA display characteristic peaks at about 278 nm, which could result from  $\pi$ - $\pi^*$  transition due to the presence of three aromatic amino acids (tryptophan, tyrosine and phenylalanine) on the

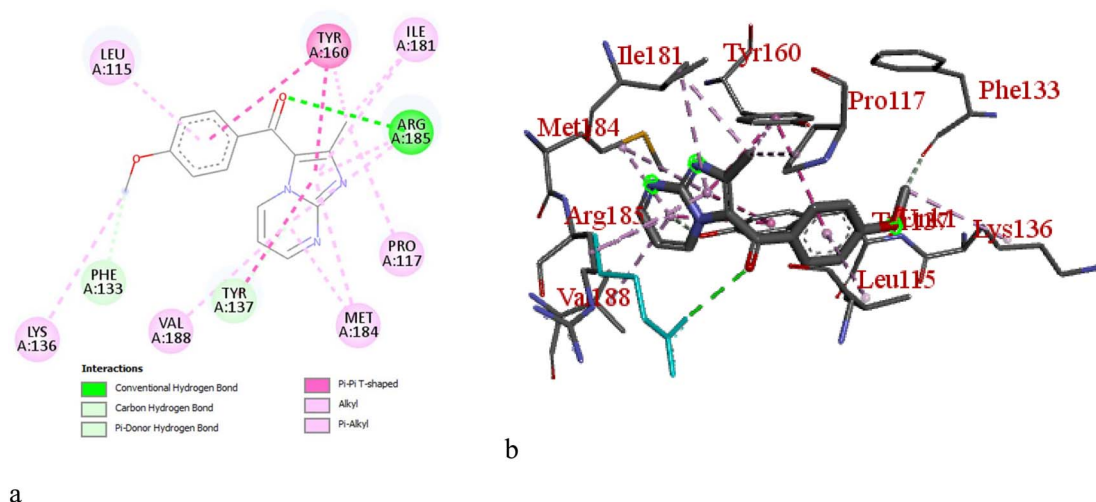


Fig. 3 (a and b) 2D and 3D poses of the interaction of ligand 5e with BSA protein.



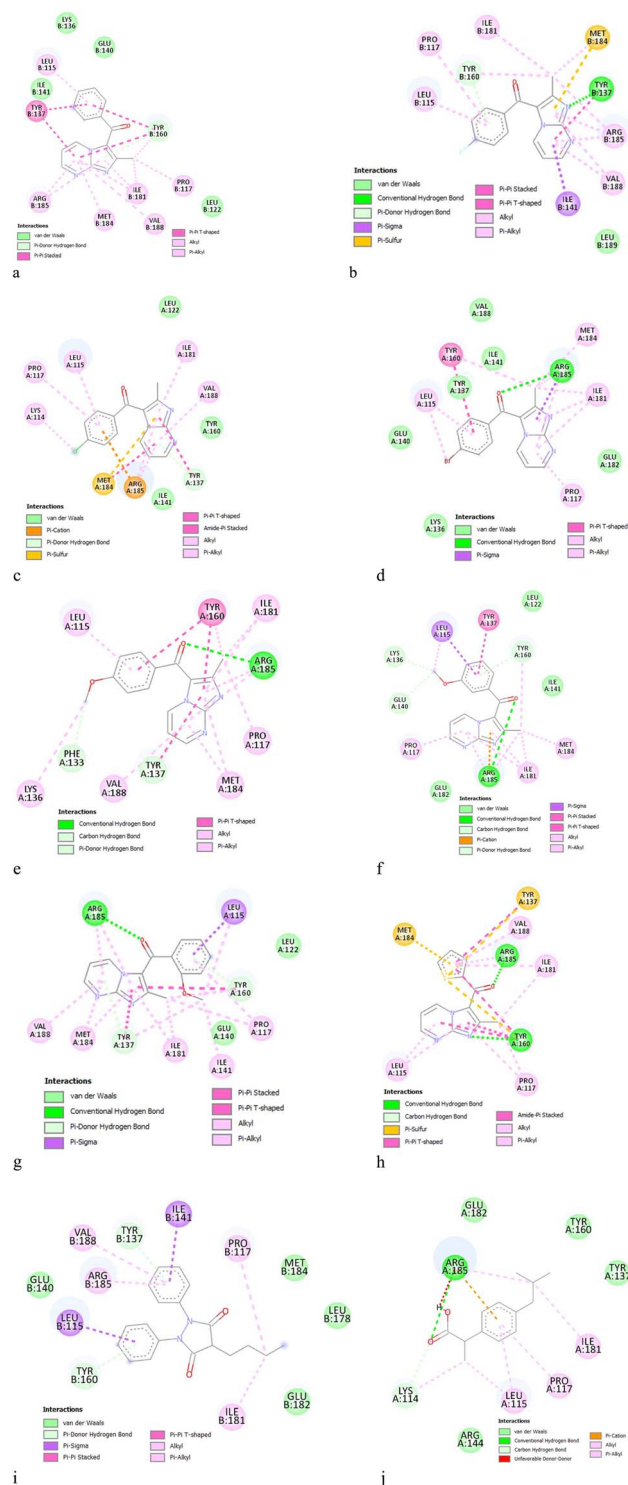


Fig. 4 (a–j) 2D diagram showing of the interaction of ligand 5a–h, phenylbutazone (PBZ) and ibuprofen (IBP) with BSA protein.

protein chain surface.<sup>42</sup> In the presence of increasing concentration of **5e**, **5h** and **5c**, the intensity of the BSA spectra continues to rise due to the changes in the environment surrounding aromatic amino acids Trp, Phe and Tyr (Fig. 5a–c). This perturbation in the absorption spectra of BSA indicates the preferable interactions between BSA and compounds studied.

The hyperchromic effect in the absorption spectra of BSA on subsequent addition of **5e**, **5h** and **5c** indicated a possible static interaction between BSA and imidazo[1,2-*a*]pyrimidine ligands.

The stoichiometric ratio in which BSA protein interacts with a ligand was determined employing Job's method of continuous variation. Ten different solutions were prepared with different volumes of BSA and **5e** fixed at a constant concentration of 15  $\mu\text{M}$ . In Job's plot, the absorbance at  $\lambda_{\text{max}}$  (278 nm) was measured and absorbance *versus* mole fraction of ligand **5e** was plotted (Fig. 5d). The maxima in Job's plot was observed at 0.5 on the axis which indicates 1 : 1 binding stoichiometry between BSA and ligand **5e**.<sup>43</sup>

**2.2.3. Fluorescence quenching studies.** Fluorescence spectroscopy is a sensitive tool to explore the mode of interaction, binding mechanism, binding constant and binding sites of ligands with biomolecules. Binding studies of BSA and ligand **5e**, **5h** and **5c** were also studied by steady state fluorescence. Intrinsic fluorescence intensity was measured using a constant concentration of BSA (15  $\mu\text{M}$ ) and varying concentration of compounds **5e**, **5h** and **5c** (ranging from 0 to 56  $\mu\text{M}$ ) at room temperature (25  $^{\circ}\text{C}$ ). The emission spectra were recorded in the spectral range of 300–430 nm by fixing the excitation wavelength at 280 nm. A progressive decrease in the fluorescence intensity of BSA spectra after the gradual addition of compounds indicates that imidazo[1,2-*a*]pyrimidines quenched the fluorescence of BSA due to binding interactions. Fig. 6 illustrates the fluorescence intensity of BSA at 343 nm, exhibiting a slight red shift (for *e.g.* 10 nm in case of **5e**) as a result of quenching.

Further to understand the interaction between ligand **5e** and BSA, a fixed concentration of compound **5e** (0.1 mM) was titrated with increasing BSA concentration (0–40  $\mu\text{M}$ ) at room temperature (25  $^{\circ}\text{C}$ ) and the effects on emission profiles was monitored following excitation at 367 nm (Fig. 6d). Free ligand (**5e**) exhibits maximum in absorption peak at 367 nm. Therefore, the emission spectra were recorded in the spectral range of 387–700 nm by fixing the excitation wavelength at 367 nm. Ligand **5e** exhibited a strong fluorescence emission at 454 nm in the absence of BSA. As expected, emission intensity increases at 481 nm as the concentration of BSA is increased (Fig. 6d). The emission increases linearly as a function of BSA concentration until about 28  $\mu\text{M}$ , where it reaches a plateau that may be attributed to a self-quenching phenomenon.<sup>44</sup> Red shift in emission maxima was observed in the presence of BSA.

The static and dynamic quenching process can be identified by calculating the quenching constant  $K_q$  using the Stern–Volmer equation<sup>45</sup> (eqn (1)).

$$\frac{F_0}{F} = 1 + K_{sv}[Q] = 1 + K_q\tau_0[Q]. \quad (1)$$

where  $F_0$  denotes the fluorescence intensity of BSA,  $F$  refers to the fluorescence intensity of BSA in the presence of ligands (**5e**, **5h** and **5c**) at the various concentrations (0–56  $\mu\text{M}$ ).  $[Q]$  is a molar concentration of ligand.  $K_{sv}$  and  $K_q$  are the Stern–Volmer constant and quenching constant, respectively.  $\tau_0$  denotes the average fluorescence lifetime of BSA. Quenching



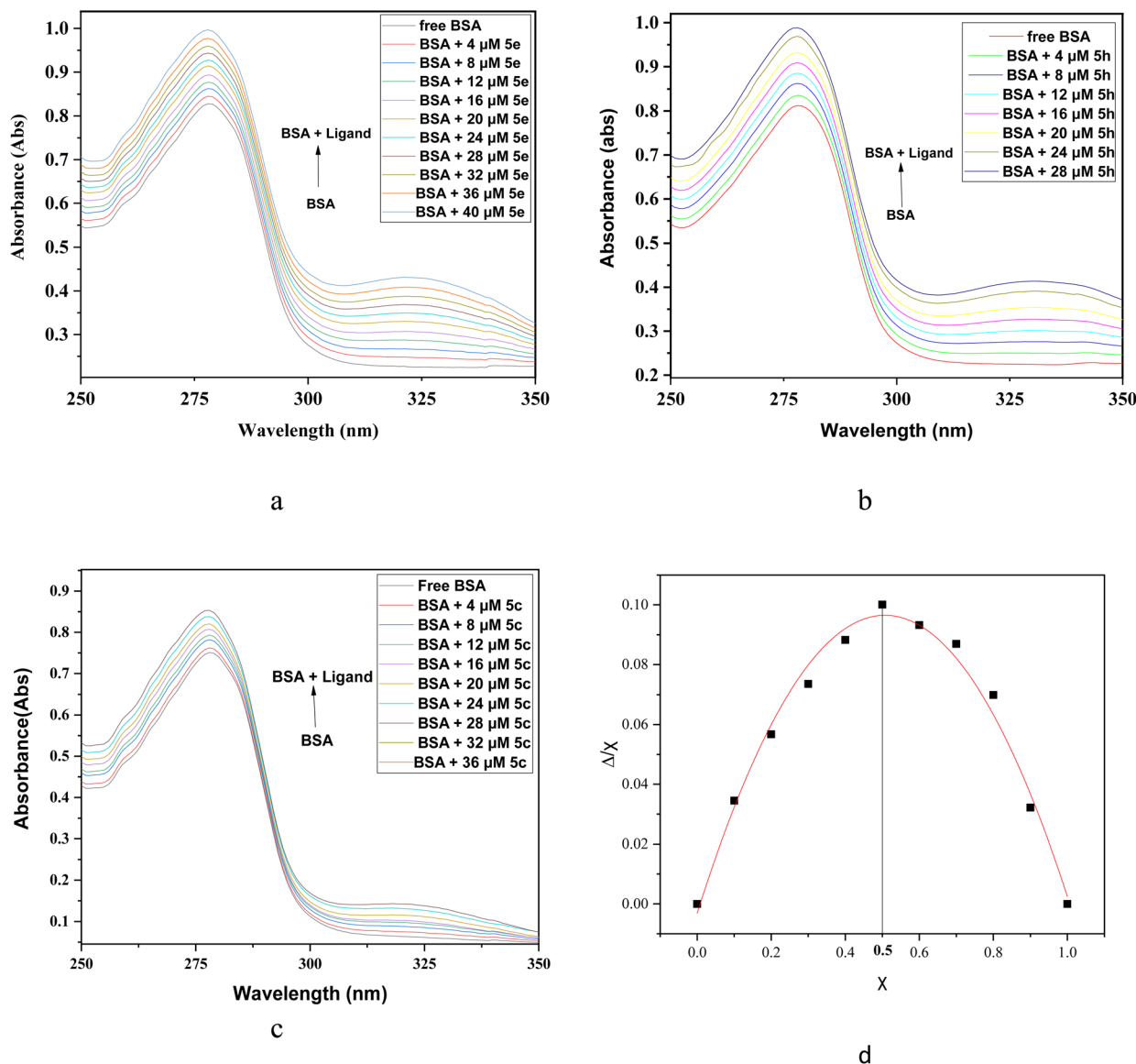


Fig. 5 UV-visible spectra of BSA-ligand complex system at increasing concentrations of ligand at constant BSA concentration of 15  $\mu\text{M}$  in physiological pH 7.2 of Tris-HCl buffer at room temperature. (a) BSA spectra at variable concentration of **5e** (0–40  $\mu\text{M}$ ); (b) BSA spectra at variable concentration of **5h** (0–28  $\mu\text{M}$ ); (c) BSA spectra at variable concentration of **5c** (0–36  $\mu\text{M}$ ). (d) Job's plot for BSA-**5e** complex system.

constant  $K_q$  was calculated by employing average lifetime  $\tau_0 = 10^{-8}$  s.

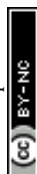
In order to find the effectiveness of binding of compounds with BSA, fluorescent data were utilized to compute quenching constant ( $K_q$ ) and Stern-Volmer constant ( $K_{sv}$ ) by plotting emission spectral data at  $\lambda_{\text{max}}$  (343 nm) at room temperature by employing the Stern-Volmer equation (eqn (1)). Based on the linear fit graph of fluorescence intensity ( $F_0$ ) of BSA in the absence of ligand to the fluorescence intensity ( $F$ ) of BSA in the presence of ligand at varying concentration was plotted against increasing concentration of ligand  $[Q]$  (Fig. 7), quenching constant ( $K_q$ ) and Stern-Volmer constant ( $K_{sv}$ ) were calculated and shown in Table 5. The values of quenching constant  $K_q$  are  $2.0 \times 10^{12}$ ,  $4.9 \times 10^{12}$ , and  $5.0 \times 10^{12} \text{ M}^{-1} \text{ s}^{-1}$  for compounds (**5c**, **5e** and **5h**, respectively) which were found to be higher than

maximum scatter collision quenching constant<sup>46</sup> ( $2 \times 10^{10} \text{ M}^{-1} \text{ s}^{-1}$ ), indicating the quenching of fluorophore (BSA) by our compounds is initiated by the static mechanism with formation complex at the ground state. According to  $K_q$  values, the capability of the compounds to quench the emission intensity of BSA follows the sequence **5h** > **5e** > **5c**.

**2.2.3.1 Identification of binding constant and number of binding sites.** Further, Modified Stern-Volmer equation<sup>46</sup> (eqn (2)) was used to calculate binding constant  $K_b$  and number of binding sites ( $n$ ).

$$\frac{\log(F_0 - F)}{F} = n \log[Q] + \log K_b. \quad (2)$$

From the plot of  $\log(F_0 - F)/F$  against  $\log[Q]$  (Fig. 8), the binding constant  $K_b$  and the number of binding sites  $n$  have



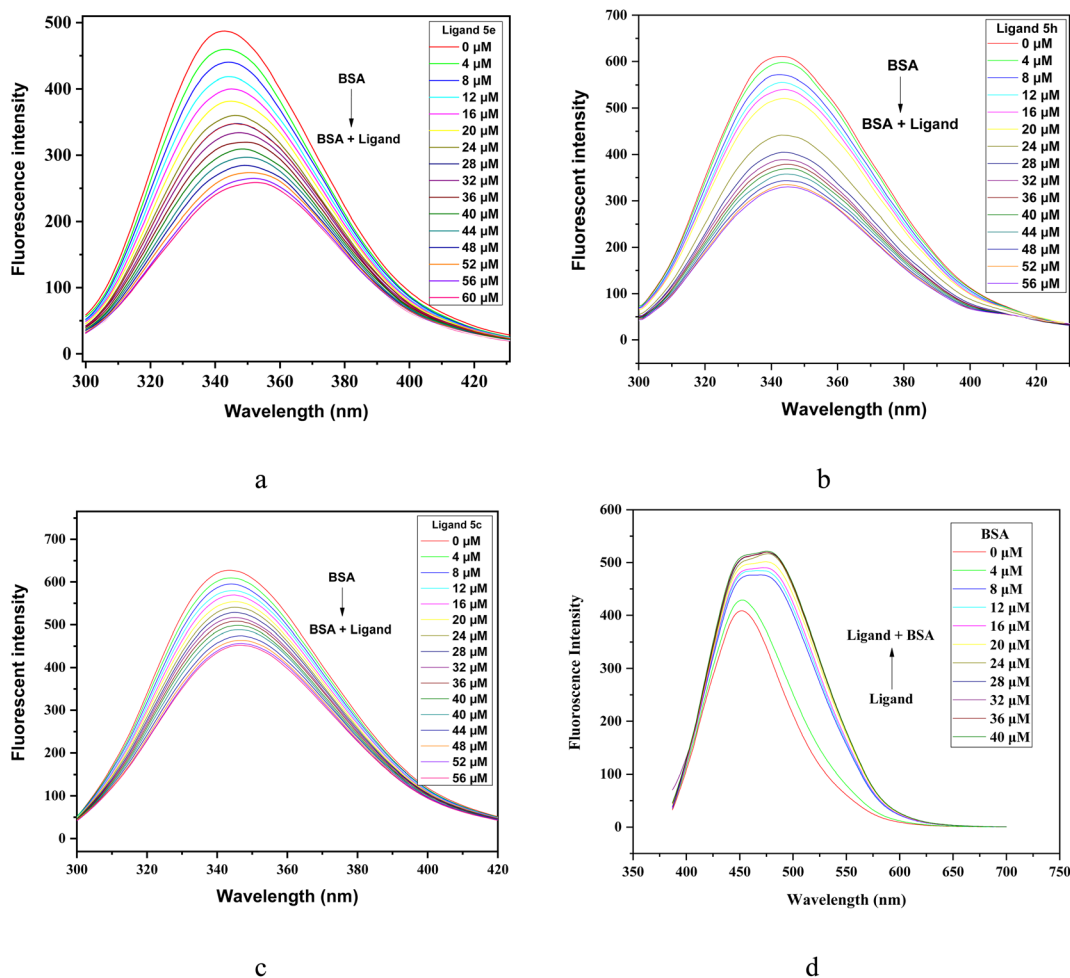


Fig. 6 Emission spectra of BSA (15  $\mu\text{M}$ ) and in presence of increasing concentration of compounds (a) 5e (0–60  $\mu\text{M}$ ), (b) 5h (0–56  $\mu\text{M}$ ), (c) 5c (0–56  $\mu\text{M}$ ) and (d) emission spectra of ligand 5e (0.1 mM) and in presence of increasing concentration of BSA (0–40  $\mu\text{M}$ ).

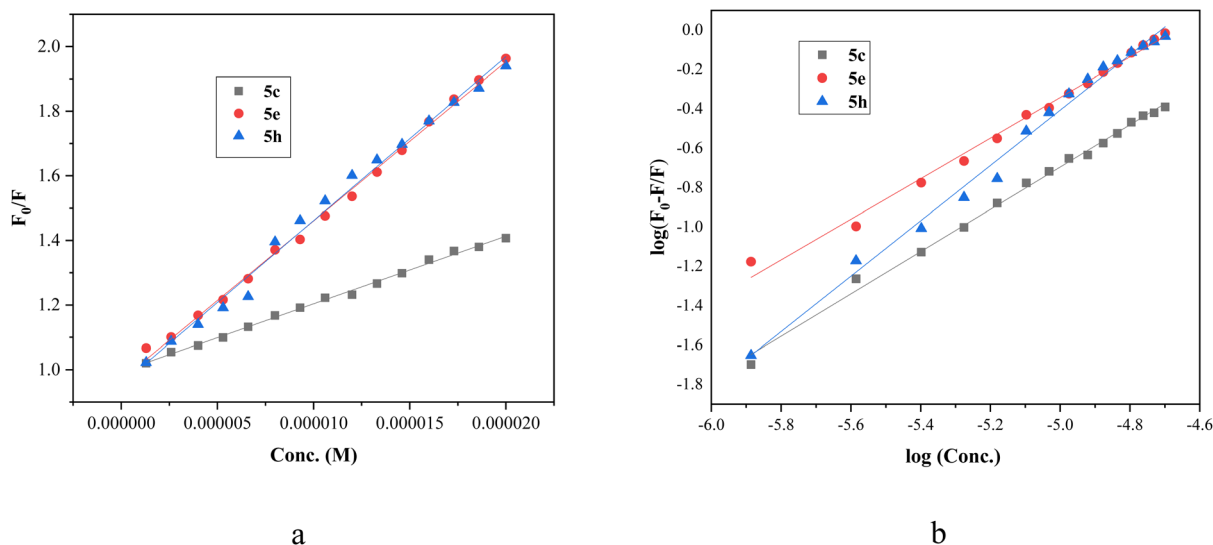


Fig. 7 (a) Stern–Volmer plot of BSA quenching by compounds 5e, 5h and 5c. (b) Double logarithmic plot employed to determine binding parameters for compounds 5e, 5h and 5c.



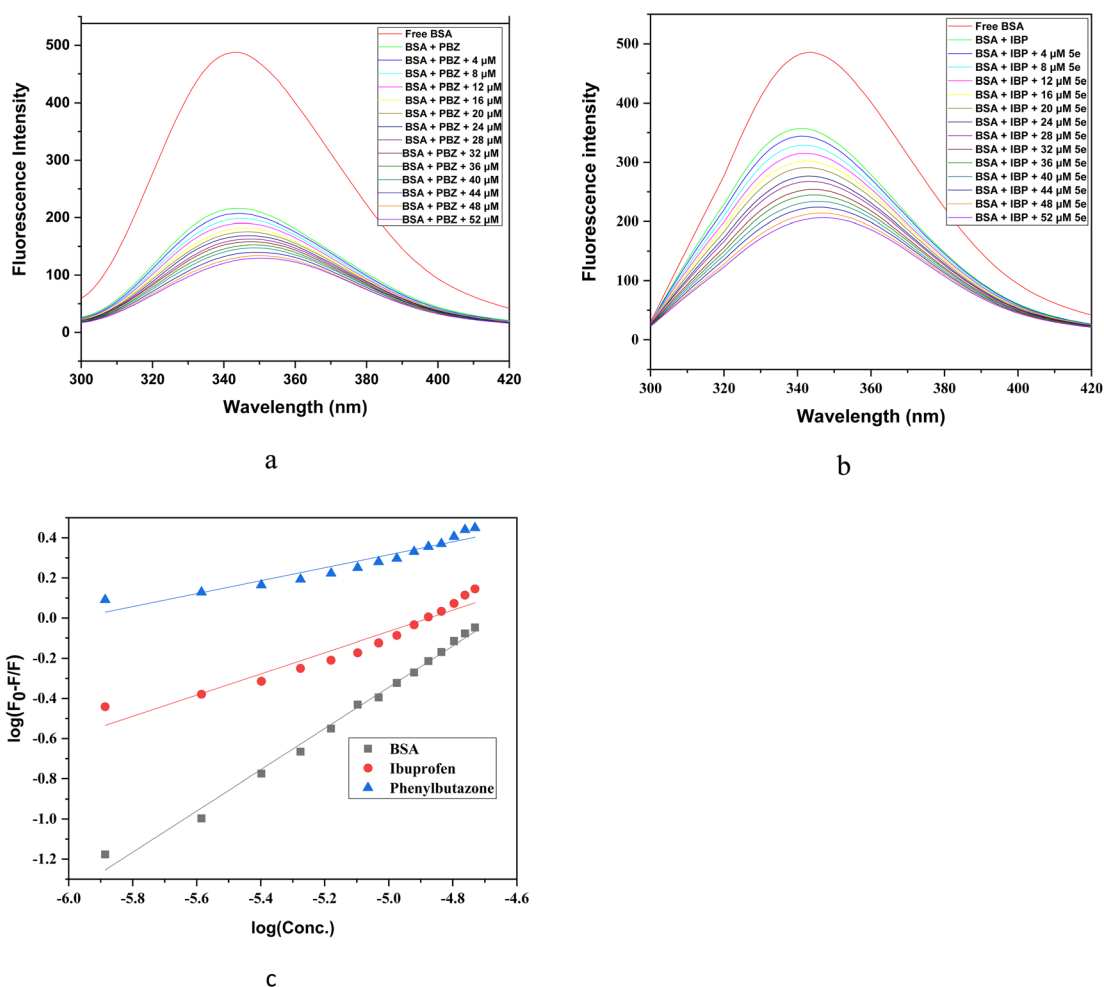
**Table 5** Stern–Volmer constant, binding constant, quenching constant and number of binding sites for the interaction of BSA with compound 5c, 5e and 5h

Compd	$K_{SV} \times 10^4 \text{ (M}^{-1}\text{)}$	$K_q \times 10^{12} \text{ (M}^{-1} \text{ s}^{-1}\text{)}$	$\text{Log } K_b$	$K_b \text{ (M}^{-1}\text{)}$	n	$\Delta G^\circ \text{ (kcal mol}^{-1}\text{)}$
5c	$2.0 \pm 0.03$	$2.0 \pm 0.03$	$4.6 \pm 0.1$	$4.7 \times 10^4$	1.0	−6.27
5e	$4.9 \pm 0.07$	$4.9 \pm 0.07$	$4.8 \pm 0.1$	$6.4 \times 10^4$	$1.0 \pm 0.02$	−6.54
5h	$5.0 \pm 0.1$	$5.0 \pm 0.1$	$6.6 \pm 0.1$	$4.2 \times 10^6$	$1.4 \pm 0.03$	−9.00

been obtained from the intercept and slope, respectively. The results are mentioned in Table 5. The value of  $n$  unveils the presence of a single binding site in BSA for imidazopyrimidines. The desired value for binding constant is contemplated to be in the range  $10^4$ – $10^6 \text{ M}^{-1}$ . The value of  $K_b$  observed for 5c, 5e and 5h are in this range, suggesting that the ligand binding to BSA protein is reversible and allows the transport and release of the ligand to the target site. Results indicate that the compounds bind to BSA in the order of  $5h > 5e > 5c$ . Furthermore, the negative value of standard Gibbs free energy change ( $\Delta G^\circ$ ) from eqn (3) suggests the spontaneity of the binding process that resulted into BSA–imidazopyrimidine complex.

$$G^\circ = -RT \ln K_b \quad (3)$$

**2.2.4. Competitive displacement assay.** Structural information regarding different ligands' interactions with protein is often investigated using site markers which are small molecules with specific binding locations in the BSA structure. Ligands can bind to BSA protein through two major sites of binding, recognized as Sudlow's Site I and Sudlow's Site II which are located in the hydrophobic cavities in subdomains IIA and IIIA, respectively. Site-I is known to bind with markers like phenylbutazone, warfarin and dansylamide while site-II illustrates



**Fig. 8** Fluorescence spectra of (a) BSA-PBZ and (b) BSA-IBP complex in the absence and presence of increasing concentration of 5e (0–62 μM) at 298 K. (c) Double logarithmic plot employed to determine binding constants for compound 5e the absence (DMSO) and presence of site markers, ibuprofen and phenylbutazone.



**Table 6** Binding constants for the BSA-5e complex in the presence of site markers at 298 K

System	Log $K_b$	$K_b$ ( $M^{-1}$ )
BSA + 5e	4.790	$6.2 \times 10^4$
BSA + PBZ + 5e	1.932	$0.0085 \times 10^4$
BSA + IBP + 5e	2.573	$0.374 \times 10^4$

binding towards markers like ibuprofen, diazepam, flufenamic acid and dansylglycine.<sup>47</sup> Thus, in order to identify the binding site for ligand 5e on BSA, a displacement assay was performed selecting phenylbutazone (PBZ) and ibuprofen (IBP) as the site marker ligands.

In this experiment, the fluorescence spectra of BSA–PBZ and BSA–IBP complexes (1 : 1) were recorded in the absence and presence of the subsequent increasing amounts of ligand 5e (0–52  $\mu$ M) (Fig. 8a and b). A decrement in the emission intensity was observed by the increase in the concentration of 5e, and the data thus obtained was utilized to plot a graph of  $\log(F_0 - F)/F$  against  $\log[Q]$  for complex 5e in the absence and presence of site markers as shown in Fig. 8c. The binding constant  $K_b$  for phenylbutazone and ibuprofen was found to be  $0.0085 \times 10^4$  and  $0.374 \times 10^4$ , respectively (Table 6). It can be observed that the  $K_b$  and  $\log K_b$  values of the compound 5e with BSA significantly decrease in the presence of phenylbutazone and ibuprofen, compared to the  $K_b$  value in the absence of any site markers, indicating that there are competitive interactions between compound 5e and the two site makers with BSA.<sup>48</sup> Therefore, it can be inferred that the compound 5e may interact with BSA at both site I and site II. Nevertheless, the value of  $K_b$  is much lower in the presence of phenylbutazone, suggesting compound 5e mainly binds in Sudlow's Site I of BSA.

### 3. Conclusions

In summary, we have developed a simple and efficient approach for the synthesis of 2,3-di substituted imidazo[1,2-*a*]pyrimidines from  $\alpha$ -bromo- $\beta$ -diketones and 2-aminopyrimidine which could serve as promising scaffolds for the development of novel bioactive molecules. The structure of regioisomeric product was characterized by  $^1H$ ,  $^{13}C$ , HMBC, HMQC, IR spectral, and mass spectrometric studies. The salient features of this approach include catalyst-free mild reaction conditions, inexpensive reagents, broad substrate scope, high yields, and easy workup procedure in a single synthetic operation. Spectroscopic methods and molecular docking studies were used to explore binding interaction of imidazo[1,2-*a*]pyrimidines with BSA. UV-visible study confirmed the interaction and 1 : 1 binding stoichiometry. Fluorescence analyses revealed that compounds bind moderately with BSA through static quenching mechanism and binding constant revealed that compound 5h exhibits greater binding affinity than 5c and 5e. Negative  $\Delta G^\circ$  values suggested that the binding process was spontaneous. Furthermore, site-marker competitive binding assay indicated that compound 5e interacts with BSA at both site I and site II and site I is the main binding site. The present study illustrates that the nature of substituents modifies binding properties of imidazo

[1,2-*a*]pyrimidines to BSA and it is expected to influence design, development and synthesis of novel imidazo[1,2-*a*]pyrimidine derivatives with significant medical applications.

## 4. Experimental

### 4.1 General

All the chemicals and solvents used in the present study were purchased from commercial suppliers of Hi-media and Avera, India, and used without further purification. To monitor the progress of the reaction and purity of the products, TLC experiments were performed on 0.2 mm Merck precoated silica gel 60 F254-coated aluminum plates. The mixture of ethyl acetate and petroleum ether was used as the mobile phase and spots were visualized under UV light at 254 nm. Melting points were determined using an electrical digital Melting Point Apparatus (MEPA) in an open capillary tube and were uncorrected. IR spectra were recorded on Buck Scientific IR M-500 instrument in KBr pellets ( $\nu_{max}$  in  $cm^{-1}$ ).  $^1H$  and  $^{13}C$  NMR spectra were recorded on a Jeol ECZS-400 instrument at 400 and 100 MHz, respectively, using DMSO- $d_6$  as a solvent and tetramethylsilane (TMS) as an internal standard (the chemical shift in  $\delta$  scale and coupling constants (*J*) were expressed in parts per million (ppm) and hertz, respectively). High-resolution mass spectra (HRMS) were measured in the ESI+ mode at CIL GJU, Hisar. 2D correlation spectroscopy, ( $^1H$ - $^{13}C$ ) gs-HSQC, ( $^1H$ - $^{13}C$ ) gs-HMBC of the samples were carried out at Kurukshetra University, Kurukshetra. The UV-vis spectra were recorded on a UV-vis spectrophotometer 117 (Systronic, India) with 1 cm path-length cell. Fluorescence spectra were recorded on the Shimadzu-5301pc spectro-fluorophotometer (Kyoto, Japan).

**4.1.1. General procedure for the synthesis of 3-aryl-2-methylimidazo[1,2-*a*]pyrimidines (5a-h).** A mixture of appropriate unsymmetrical  $\beta$ -diketone 1a-h (1.0 mmol), NBS 2 (1.0 mmol) and 2-aminopyrimidine 4 (1.0 mmol) in dichloromethane (10 mL) was taken in a conical flask and stirred at room temperature for about 5–6 h. Progress of the reaction was monitored by TLC (ethyl acetate: petroleum ether; 60 : 40, v/v). After the completion of reaction (as indicated by TLC), the solvent was evaporated on the rotary evaporator and the resulting mixture was neutralized with a saturated solution of sodium bicarbonate and extracted with ethyl acetate. The gummy mass obtained was triturated with petroleum ether. Then acetone (5 mL) was added, stirred at room temperature for 15–20 minutes and the residue thus obtained was filtered, washed with cold ethanol, dried, and recrystallized from ethanol to obtain the target 3-arylimidazo[1,2-*a*]pyrimidine (5a-h).

**4.1.1.1 1-Phenyl-1-(2-methylimidazo[1,2-*a*]pyrimidin-3-yl) methanone (5a).** White solid; m.p. 187  $^\circ$ C; yield: 0.190 g (80%); IR (KBr)  $\nu_{max}$  ( $cm^{-1}$ ): 1648 (C=O).

$^1H$  NMR (400 MHz, DMSO- $d_6$ ):  $\delta$  9.64–9.66 (d, 1H, *J* = 6.8 Hz, 7 H), 9.02 (d, 1H, *J* = 4.0 Hz, 5-H), 7.79–7.81 (d, 2H, *J* = 7.6 Hz, Ph-2',6'-H), 7.73 (dd, 1H, *J*<sub>o</sub> = 6.8, *J*<sub>m</sub> = 4.4 Hz, Ph-4'-H), 7.60–7.64 (m, 3H, Ph-3',5', 6-H), 2.21 (s, 3H, 2-CH<sub>3</sub>).

$^{13}C$  NMR (100 MHz, DMSO- $d_6$ ):  $\delta$  186.20, 156.99, 147.59, 146.82, 138.81, 138.08, 133.52, 129.38, 129.27, 119.67, 113.66, 15.44.



HRMS (ESI)  $m/z$ : 238.0890  $[M + H]^+$ .

Elemental analysis: calcd. for  $C_{14}H_{11}N_3O$ : C, 70.87; H, 4.67; N, 17.71% found: C, 70.65; H, 4.74; N, 17.48%.

4.1.1.2 1-(4'-Fluorophenyl)-1-(2-methylimidazo[1,2-a]pyrimidin-3-yl)methanone (**5b**). White solid; m.p. 206 °C; yield: 0.202 g (79%); IR (KBr)  $\nu_{\max}$  ( $\text{cm}^{-1}$ ): 1650 (C=O).

$^1\text{H}$  NMR (400 MHz, DMSO- $d_6$ ):  $\delta$  9.60–9.62 (dd, 1H,  $J = 7.2$ ,  $J = 2.0$  Hz, 7-H), 8.99–9.01 (dd, 1H,  $J = 4.4$ ,  $J = 2.0$  Hz, 5-H), 7.87–7.91 (dt, 2H,  $J_o = 8.8$ ,  $J_{(m)HF} = 5.6$  Hz, Ph-2',6'-H), 7.59–7.62 (dd, 1H,  $J = 6.8$  Hz,  $J = 4.4$  Hz, 6-H), 7.43–7.48 (t, 2H,  $J_o = 8.8$  Hz, Ph-3',5'-H), 2.24 (s, 3H, 2-CH<sub>3</sub>).

$^{13}\text{C}$  NMR (100 MHz, DMSO- $d_6$ ):  $\delta$  184.85, 156.91, 147.04, 138.05, 135.44, 135.41, 132.48, 132.38, 119.67, 116.68, 116.46, 113.56, 15.65.

HRMS (ESI)  $m/z$ : 256.0836  $[M + H]^+$ .

Elemental analysis: calcd. for  $C_{14}H_{10}FN_3O$ : C, 65.88; H, 3.95; N, 16.46% found: C, 65.81; H, 3.81; N, 16.63%.

4.1.1.3 1-(4'-Chlorophenyl)-1-(2-methylimidazo[1,2-a]pyrimidin-3-yl)methanone (**5c**). White solid; m.p. 224 °C; yield: 0.210 g (77%); IR (KBr)  $\nu_{\max}$  ( $\text{cm}^{-1}$ ): 1650 (C=O).

$^1\text{H}$  NMR (400 MHz, DMSO- $d_6$ ):  $\delta$  9.61–9.63 (d, 1H,  $J = 6.8$  Hz, 7-H), 9.01 (d, 1H,  $J = 3.2$  Hz, 5-H), 7.81–7.83 (d, 2H,  $J = 8.4$  Hz, Ph-2',6'-H), 7.68–7.70 (d, 2H,  $J = 8.4$  Hz, Ph-3',5'-H), 7.61–7.64 (dd, 1H,  $J = 6.4$ ,  $J = 4.4$  Hz, 6-H), 2.24 (s, 3H, 2-CH<sub>3</sub>).

$^{13}\text{C}$  NMR (100 MHz, DMSO- $d_6$ ):  $\delta$  184.96, 156.98, 147.98, 146.98, 138.31, 138.05, 137.50, 131.28, 129.53, 119.56, 113.63, 15.63.

HRMS (ESI)  $m/z$ : 272.0469  $[M + H]^+$ ; 274.0463  $[M + 1 + 2]^+$ ; (3 : 1).

Elemental analysis: calcd. for  $C_{14}H_{10}ClN_3O$ : C, 61.89; H, 3.71; N, 15.47% found: C, 61.78; H, 3.63; N, 15.25%.

4.1.1.4 1-(4'-Bromophenyl)-1-(2-methylimidazo[1,2-a]pyrimidin-3-yl)methanone (**5d**). White solid; m.p. 248 °C; yield: 0.227 g (72%); IR (KBr)  $\nu_{\max}$  ( $\text{cm}^{-1}$ ): 1650 (C=O).

$^1\text{H}$  NMR (400 MHz, DMSO- $d_6$ ):  $\delta$  9.61–9.63 (dd, 1H,  $J = 6.8$ ,  $J = 1.2$  Hz, 7-H), 8.97 (dd, 1H,  $J = 4.0$ ,  $J = 1.2$  Hz, 5-H), 7.82–7.84 (d, 2H,  $J = 8.4$  Hz, Ph-2',6'-H), 7.71–7.73 (d, 2H,  $J = 8.4$  Hz, Ph-3',5'-H), 7.56–7.58 (dd, 1H,  $J = 6.8$ ,  $J = 4.4$  Hz, 6-H), 2.21 (s, 3H, 2-CH<sub>3</sub>).

$^{13}\text{C}$  NMR (100 MHz, DMSO- $d_6$ ):  $\delta$  185.19, 156.39, 149.45, 147.70, 138.11, 137.81, 132.42, 131.27, 127.15, 119.54, 113.15, 16.14.

HRMS (ESI)  $m/z$ : 316.9978  $[M + H]^+$ ; 318.9959  $[M + 1 + 2]^+$ ; (1 : 1).

Elemental analysis: calcd. for  $C_{14}H_{10}BrN_3O$ : C, 53.19; H, 3.19; N, 13.29% found: C, 52.98; H, 3.24; N, 13.43%.

4.1.1.5 1-(4'-Methoxyphenyl)-1-(2-methylimidazo[1,2-a]pyrimidin-3-yl)methanone (**5e**). White solid; m.p. 190 °C; yield: 0.230 g (86%); IR (KBr)  $\nu_{\max}$  ( $\text{cm}^{-1}$ ): 1645 (C=O).

$^1\text{H}$  NMR (400 MHz, DMSO- $d_6$ ):  $\delta$  9.52–9.54 (dd, 1H,  $J = 7.0$ ,  $J = 2.4$  Hz, 7-H), 9.01–9.03 (dd, 1H,  $J = 4.8$ ,  $J = 2.0$  Hz, 5-H), 7.82–7.84 (d, 2H,  $J = 8.8$  Hz, Ph-2',6'-H), 7.61–7.64 (dd, 1H,  $J = 6.0$ ,  $J = 4.8$  Hz, 6-H), 7.14–7.16 (d, 2H,  $J = 9.2$  Hz, Ph-3',5'-H), 3.90 (s, 3H, Ph-4'-OMe), 2.31 (s, 3H, 2-CH<sub>3</sub>).

$^{13}\text{C}$  NMR (100 MHz, DMSO- $d_6$ ):  $\delta$  184.54, 164.08, 157.17, 146.21, 138.12, 132.26, 130.82, 119.82, 114.78, 113.76, 56.23, 15.07.

HRMS (ESI)  $m/z$ : 268.1003  $[M + H]^+$ .

Elemental analysis: calcd. for  $C_{15}H_{13}N_3O_2$ : C, 67.40; H, 4.90; N, 15.72% found: C, 67.28; H, 4.74; N, 15.59%.

4.1.1.6 1-(3'-Methoxyphenyl)-1-(2-methylimidazo[1,2-a]pyrimidin-3-yl)methanone (**5f**). White solid; m.p. 215 °C; yield: 0.220 g (82%); IR (KBr)  $\nu_{\max}$  ( $\text{cm}^{-1}$ ): 1645 (C=O).

$^1\text{H}$  NMR (400 MHz, DMSO- $d_6$ ):  $\delta$  9.63–9.65 (dd,  $J = 6.8$ ,  $J = 1.2$  Hz, 1H, 7-H), 9.03–9.04 (dd, 1H,  $J = 4.4$ ,  $J = 2.8$  Hz, 5-H), 7.63–7.66 (dd, 1H,  $J = 6.8$ ,  $J = 4.8$  Hz, 6-H), 7.51–7.55 (t, 1H,  $J_o = 7.6$  Hz, Ph-5'-H), 7.36–7.38 (d, 1H,  $J_o = 7.6$  Hz, Ph-6'-H), 7.29–7.31 (m, 2H, Ph-2',4'-H), 3.83 (s, 3H, Ph-3'-OMe), 2.23 (s, 3H, 2-CH<sub>3</sub>).

$^{13}\text{C}$  NMR (100 MHz, DMSO- $d_6$ ):  $\delta$  185.89, 159.88, 157.17, 147.31, 146.60, 140.15, 138.12, 130.67, 121.53, 119.66, 119.54, 113.83, 113.74, 55.99, 15.25.

HRMS (ESI)  $m/z$ : 268.1003  $[M + H]^+$ .

Elemental analysis: calcd. for  $C_{15}H_{13}N_3O_2$ : C, 67.40; H, 4.90; N, 15.72% found: C, 67.52; H, 4.66; N, 15.97%.

4.1.1.7 1-(2'-Methoxyphenyl)-1-(2-methylimidazo[1,2-a]pyrimidin-3-yl)methanone (**5g**). White solid; m.p. 229 °C; yield: 0.213 g (80%); IR (KBr)  $\nu_{\max}$  ( $\text{cm}^{-1}$ ): 1645 (C=O).

$^1\text{H}$  NMR (400 MHz, DMSO- $d_6$ ):  $\delta$  9.64–9.66 (dd, 1H,  $J = 6.8$ ,  $J = 2.0$  Hz, 7-H), 9.06–9.07 (dd, 1H,  $J = 4.4$ ,  $J = 2.0$  Hz, 5-H), 7.67–7.70 (dd, 1H,  $J = 6.8$ ,  $J = 4.4$  Hz, 6-H), 7.52–7.56 (t, 1H,  $J = 8.0$  Hz, Ph-4'-H), 7.37–7.39 (d, 1H,  $J = 7.6$  Hz, Ph-6'-H), 7.30–7.33 (m, 2H, Ph-3',5'-H), 3.84 (s, 3H, Ph-2'-OMe), 2.25 (s, 3H, 2-CH<sub>3</sub>).

$^{13}\text{C}$  NMR (100 MHz, DMSO- $d_6$ ):  $\delta$  185.89, 159.92, 157.59, 146.23, 140.03, 138.34, 130.76, 130.71, 121.68, 119.75, 119.70, 114.17, 113.81, 56.05, 15.05.

HRMS (ESI)  $m/z$ : 268.1003  $[M + H]^+$ .

Elemental analysis: calcd. for  $C_{15}H_{13}N_3O_2$ : C, 67.40; H, 4.90; N, 15.72% found: C, 67.71; H, 4.56; N, 15.46%.

4.1.1.8 1-(Thiophen-2'-yl)-1-(2-methylimidazo[1,2-a]pyrimidin-3-yl)methanone (**5h**). White solid; m.p. 244 °C; yield: 0.193 g (79%); IR (KBr)  $\nu_{\max}$  ( $\text{cm}^{-1}$ ): 1654 (C=O).

$^1\text{H}$  NMR (400 MHz, DMSO- $d_6$ ):  $\delta$  9.43–9.45 (dd, 1H,  $J = 6.8$ ,  $J = 1.2$  Hz, 7-H), 9.02–9.03 (dd, 1H,  $J = 4.0$ ,  $J = 1.2$  Hz, 5-H), 8.24–8.25 (d, 1H,  $J = 4.4$  Hz, thienyl-5'-H), 7.93–7.94 (d, 1H,  $J = 3.2$  Hz, thienyl-3'-H), 7.61–7.64 (dd, 1H,  $J = 6.8$ ,  $J = 4.4$  Hz, 6-H), 7.33–7.35 (t, 1H,  $J = 4.4$  Hz, thienyl-4'-H), 2.51 (s, 3H, 2-CH<sub>3</sub>).

$^{13}\text{C}$  NMR (100 MHz, DMSO- $d_6$ ):  $\delta$  177.02, 157.36, 145.97, 144.53, 142.68, 138.04, 137.19, 136.61, 129.26, 119.49, 113.74, 14.97.

HRMS (EI)  $m/z$ : 244.0467  $[M + H]^+$ .

Elemental analysis: calcd. for  $C_{12}H_9N_3OS$ : C, 59.24; H, 3.73; N, 17.27% found: C, 59.12; H, 3.62; N, 17.49%.

## 4.2 Binding studies

**4.2.1. Molecular docking studies.** Primarily ligand structures were drawn using ChemDraw Professional 15.0 software and the crystal structure of the BSA protein (PDB ID: 4f5s) was attained from Protein Data Bank (<https://www.rcsb.org/pdb>). The protein file was then prepared for docking by removing water molecules and packing with polar hydrogen with Kollman charges using Auto Dock Tools version 1.5.6. After preparing the PDBQT files of protein and ligands, they are



inputted in the determined grid boxes for docking. In order to perform the molecular docking analysis Auto Dock software was used with grid size (BSA grid; center\_x = 37.823, center\_y = 23.992, center\_z = 98.761; size\_x = 126, size\_y = 56, size\_z = 74; energy\_range = 4, exhaustiveness = 10) using the Lamarckian genetic algorithm for calculations, and the output results were analyzed using the BIOVIA Discovery Studio Visualizer (DSV).

**4.2.2. BSA-binding experiments.** Bovine serum albumin was purchased directly from Sigma-Aldrich Company and exploited without any absolution. For binding interaction studies, analytical grade reagents were used. BSA stock solution of 15  $\mu\text{M}$  (1 mg mL<sup>-1</sup>) concentration was prepared in 10 mM phosphate buffer saline (prepared using Na<sub>2</sub>HPO<sub>4</sub> and NaH<sub>2</sub>PO<sub>4</sub>) with pH 7.4, and then stored at 4 °C for further usage. Stock solutions of imidazo[1,2-*a*]pyrimidines (**5e**, **5c** and **5h**) of 1 mM concentration were prepared in DMSO as a solvent and further diluted with buffer as per requirement depending upon the mode of interaction study.

## Data availability

Data for this article, including experimental data, (<sup>1</sup>H, <sup>13</sup>C, [<sup>1</sup>H-<sup>13</sup>C] HMBC, and [<sup>1</sup>H-<sup>13</sup>C] HSQC) for final compounds have been included as part of the ESI.†

## Author contributions

R. A. and P. K.; conceptualization, supervision, reviewing and editing. M. S. and G. S.; methodology, investigation, computational calculations, writing and original draft preparation.

## Conflicts of interest

The author(s) confirm that this article's content has no conflict of interest.

## Acknowledgements

The authors are thankful to their respective institutions for providing the required facility to complete this work. We thank the Council of Scientific and Industrial Research (CSIR), New Delhi, India, for providing financial assistance to Manisha Sharma (Grant 09/105(0274)/2018-EMR-I) as SRF.

## References

- 1 R. Aggarwal and G. Sumran, An insight on medicinal attributes of 1,2,4-triazoles, *Eur. J. Med. Chem.*, 2020, **205**, 112652, DOI: [10.1016/j.ejmech.2020.112652](https://doi.org/10.1016/j.ejmech.2020.112652).
- 2 N. Kerru, L. Gummidi, S. Maddila, K. K. Gangu and S. B. Jonnalagadda, A Review on Recent Advances in Nitrogen-Containing Molecules and Their Biological Applications, *Molecules*, 2020, **25**(8), 1909, DOI: [10.3390/molecules25081909](https://doi.org/10.3390/molecules25081909).
- 3 F. Doganc, A. S. Aydin, E. Şahin and H. Göker, Regioselective N-alkylation of some 2 or 6-chlorinated purine analogues, *J. Mol. Struct.*, 2023, **1272**, 134200, DOI: [10.1016/j.molstruc.2022.134200](https://doi.org/10.1016/j.molstruc.2022.134200).
- 4 H. T. Abdel-Mohsen, A. Abood, K. J. Flanagan, A. Meindl, M. O. Senge and H. I. El Diwani, Synthesis, crystal structure, and ADME prediction studies of novel imidazopyrimidines as antibacterial and cytotoxic agents, *Arch. Pharm.*, 2020, **353**(3), 1900271, DOI: [10.1002/ardp.201900271](https://doi.org/10.1002/ardp.201900271).
- 5 Á. Ramírez-Trinidad, K. Carrillo-Jaimes, J. A. Rivera-Chávez and E. Hernández-Vázquez, Synthesis and cytotoxic/antimicrobial screening of 2-alkenylimidazo[1,2-*a*]pyrimidines, *Med. Chem. Res.*, 2023, **32**(1), 144–157, DOI: [10.1007/s00044-022-02997-6](https://doi.org/10.1007/s00044-022-02997-6).
- 6 M. Bayanati, M. Khoramjouy, M. Faizi, M. A. Movahed, M. Mahboubi-Rabbani and A. Zarghi, Novel Benzo[4,5]imidazo[1,2-*a*]pyrimidine derivatives as selective Cyclooxygenase-2 Inhibitors: Design, synthesis, docking studies, and biological evaluation, *Med. Chem. Res.*, 2023, **32**(3), 495–505, DOI: [10.1007/s00044-023-03022-0](https://doi.org/10.1007/s00044-023-03022-0).
- 7 S. Alqarni, L. Cooper, J. Galvan Achi, *et al.*, Synthesis, Optimization, and Structure–Activity Relationships of Imidazo[1,2-*a*]pyrimidines as Inhibitors of Group 2 Influenza A Viruses, *J. Med. Chem.*, 2022, **65**(20), 14104–14120, DOI: [10.1021/acs.jmedchem.2c01329](https://doi.org/10.1021/acs.jmedchem.2c01329).
- 8 N. M. Panchani and H. S. Joshi, Green and Catalyst-Free Synthesis of Some New Benzo[4,5]imidazo[1,2-*a*]pyrimidine Derivatives as Antimicrobial and Antitubercular Agents, *Russ. J. Org. Chem.*, 2022, **58**(4), 604–611, DOI: [10.1134/S1070428022040200](https://doi.org/10.1134/S1070428022040200).
- 9 R. Kumar, R. Singh, A. das Chagas Almeida, *et al.*, Imidazo[1,2-*a*]pyrimidine as a New Antileishmanial Pharmacophore against Leishmania amazonensis Promastigotes and Amastigotes, *ACS Omega*, 2023, **8**(43), 40613–40621, DOI: [10.1021/acsomega.3c05441](https://doi.org/10.1021/acsomega.3c05441).
- 10 P. Prasad, A. G. Kalola and M. P. Patel, Microwave assisted one-pot synthetic route to imidazo[1,2-*a*]pyrimidine derivatives of imidazo/triazole clubbed pyrazole and their pharmacological screening, *New J. Chem.*, 2018, **42**(15), 12666–12676, DOI: [10.1039/C8NJ00670A](https://doi.org/10.1039/C8NJ00670A).
- 11 E. Koti Reddy, C. Remya, A. M. Sajith, K. V. Dileep, C. Sadasivan and S. Anwar, Functionalised dihydroazo pyrimidine derivatives from Morita–Baylis–Hillman acetates: synthesis and studies against acetylcholinesterase as its inhibitors, *RSC Adv.*, 2016, **6**(81), 77431–77439, DOI: [10.1039/C6RA12507G](https://doi.org/10.1039/C6RA12507G).
- 12 F. Peytam, G. Takalloobanafshi, T. Saadattalab, *et al.*, Design, synthesis, molecular docking, and in vitro  $\alpha$ -glucosidase inhibitory activities of novel 3-amino-2,4-diarylbenzo[4,5]imidazo[1,2-*a*]pyrimidines against yeast and rat  $\alpha$ -glucosidase, *Sci. Rep.*, 2021, **11**(1), 11911, DOI: [10.1038/s41598-021-91473-z](https://doi.org/10.1038/s41598-021-91473-z).
- 13 M. Azzouzi, Z. E. Ouafi, O. Azougagh, *et al.*, Design, synthesis, and computational studies of novel imidazo[1,2-*a*]pyrimidine derivatives as potential dual inhibitors of hACE2 and spike protein for blocking SARS-CoV-2 cell entry, *J. Mol. Struct.*, 2023, **1285**, 135525, DOI: [10.1016/j.molstruc.2023.135525](https://doi.org/10.1016/j.molstruc.2023.135525).



- 14 S. C. Goodacre, L. J. Street, D. J. Hallett, *et al.*, Imidazo[1,2-*a*]pyrimidines as Functionally Selective and Orally Bioavailable GABA A  $\alpha 2/\alpha 3$  Binding Site Agonists for the Treatment of Anxiety Disorders, *J. Med. Chem.*, 2006, **49**(1), 35–38, DOI: [10.1021/jm051065l](#).
- 15 K. C. Rupert, J. R. Henry, J. H. Dodd, *et al.*, Imidazopyrimidines, potent inhibitors of p38 MAP kinase, *Bioorg. Med. Chem. Lett.*, 2003, **13**(3), 347–350, DOI: [10.1016/S0960-894X\(02\)01020-X](#).
- 16 M. L. S. O. Lima, C. B. Braga, T. B. Becher, *et al.*, Fluorescent Imidazo[1,2-*a*]pyrimidine Compounds as Biocompatible Organic Photosensitizers that Generate Singlet Oxygen: A Potential Tool for Phototheranostics, *Chem.–Eur. J.*, 2021, **27**(20), 6213–6222, DOI: [10.1002/chem.202004957](#).
- 17 M. Rawat and D. S. Rawat, Copper oxide nanoparticle catalysed synthesis of imidazo[1,2-*a*]pyrimidine derivatives, their optical properties and selective fluorescent sensor towards zinc ion, *Tetrahedron Lett.*, 2018, **59**(24), 2341–2346, DOI: [10.1016/j.tetlet.2018.05.005](#).
- 18 S. Pareek, D. Jain, S. Hussain, *et al.*, A new insight into corrosion inhibition mechanism of copper in aerated 3.5 wt.% NaCl solution by eco-friendly imidazopyrimidine dye: experimental and theoretical approach, *Chem. Eng. J.*, 2019, **358**, 725–742, DOI: [10.1016/j.cej.2018.08.079](#).
- 19 K. Karami, N. Jamshidian and M. Zakariazadeh, Synthesis, characterization and molecular docking of new C,N-palladacycles containing pyridinium-derived ligands: DNA and BSA interaction studies and evaluation as anti-tumor agents, *Appl. Organomet. Chem.*, 2019, **33**(3), 4728, DOI: [10.1002/aoc.4728](#).
- 20 S. Siddiqui, F. Ameen, I. Jahan, S. M. Nayeem and M. Tabish, A comprehensive spectroscopic and computational investigation on the binding of the anti-asthmatic drug triamcinolone with serum albumin, *New J. Chem.*, 2019, **43**(10), 4137–4151, DOI: [10.1039/C8NJ05486J](#).
- 21 T. A. Wani, A. H. Bakheit, S. Zargar, *et al.*, Toxicity Study and Binding Analysis of Newly Synthesized Antifungal N-(4-aryl/cyclohexyl)-2-(pyridine-4-yl carbonyl) hydrazinecarbothioamide Derivative with Bovine Serum Albumin, *Int. J. Mol. Sci.*, 2023, **24**(5), 4942, DOI: [10.3390/jms24054942](#).
- 22 H. Dezhampanah and A. M. Moghaddam Pour, Multi technique investigation on interaction between 5-(2-thiazolylazo)-2,4,6-triaminopyrimidine and HSA and BSA, *J. Biomol. Struct. Dyn.*, 2022, **40**(18), 8143–8154, DOI: [10.1080/07391102.2021.1906751](#).
- 23 V. S. A. K. Das, Y. Bylappa, A. Nag and M. Dolai, A dual-functional rhodamine B and azo-salicylaldehyde derivative for the simultaneous detection of copper and hypochlorite: synthesis, biological applications and theoretical insights, *Anal. Methods*, 2024, **16**(47), 8164–8178, DOI: [10.1039/D4AY01758G](#).
- 24 P. Singla, V. Luxami, R. Singh, V. Tandon and K. Paul, Novel pyrazolo[3,4-*d*]pyrimidine with 4-(1H-benzimidazol-2-yl)-phenylamine as broad spectrum anticancer agents: Synthesis, cell based assay, topoisomerase inhibition, DNA intercalation and bovine serum albumin studies, *Eur. J. Med. Chem.*, 2017, **126**, 24–35, DOI: [10.1016/j.ejmech.2016.09.093](#).
- 25 J. Liu, Y. He, D. Liu, *et al.*, Characterizing the binding interaction of astilbin with bovine serum albumin: a spectroscopic study in combination with molecular docking technology, *RSC Adv.*, 2018, **8**(13), 7280–7286, DOI: [10.1039/C7RA13272G](#).
- 26 Y. Lv, Q. Liang, Y. Li, X. Liu, D. Zhang and X. Li, Study of the binding mechanism between hydroxytyrosol and bovine serum albumin using multispectral and molecular docking, *Food Hydrocoll.*, 2022, **122**, 107072, DOI: [10.1016/j.foodhyd.2021.107072](#).
- 27 S. Khatun and Q. F. A. Riyazuddeen, In-vitro binding analysis of bovine serum albumin with sulindac/chlorpromazine: Spectroscopic, calorimetric and computational approaches, *J. Mol. Liq.*, 2020, **299**, 112124, DOI: [10.1016/j.molliq.2019.112124](#).
- 28 R. Goel, V. Luxami and K. Paul, Synthetic approaches and functionalizations of imidazo[1,2-*a*]pyrimidines: an overview of the decade, *RSC Adv.*, 2015, **5**(99), 81608–81637, DOI: [10.1039/C5RA14795F](#).
- 29 Y. Vara, E. Aldaba, A. Arrieta, J. L. Pizarro, M. I. Arriortua and F. P. Cossio, Regiochemistry of the microwave-assisted reaction between aromatic amines and  $\alpha$ -bromoketones to yield substituted 1H-indoles, *Org. Biomol. Chem.*, 2008, **6**(10), 1763, DOI: [10.1039/b719641e](#).
- 30 K. M. Bongor, R. J. van den Berg, L. H. Heitman, *et al.*, Synthesis and evaluation of homo-bivalent GnRHR ligands, *Bioorg. Med. Chem.*, 2007, **15**(14), 4841–4856, DOI: [10.1016/j.bmc.2007.04.065](#).
- 31 A. Herath, R. Dahl and N. D. P. Cosford, Fully Automated Continuous Flow Synthesis of Highly Functionalized Imidazo[1,2-*a*] Heterocycles, *Org. Lett.*, 2010, **12**(3), 412–415, DOI: [10.1021/ol902433a](#).
- 32 M. Mantipally, M. R. Gangireddy, R. Gundla, V. N. Badavath, S. R. Mandha and V. C. Maddipati, Rational design, molecular docking and synthesis of novel homopiperazine linked imidazo[1,2-*a*]pyrimidine derivatives as potent cytotoxic and antimicrobial agents, *Bioorg. Med. Chem. Lett.*, 2019, **29**(16), 2248–2253, DOI: [10.1016/j.bmcl.2019.06.031](#).
- 33 I. Pravst, M. Zupan and S. Stavber, Solvent-free bromination of 1,3-diketones and  $\beta$ -keto esters with NBS, *Green Chem.*, 2006, **8**(11), 1001–1005, DOI: [10.1039/B608446J](#).
- 34 R. Aggarwal and G. Sumran, A facile [hydroxy(tosyloxy)iodo] benzene mediated synthesis of 2-arylimidazo[1,2-*a*]pyrimidines and their conversion into 3-bromo-2-arylimidazo[1,2-*a*]pyrimidines, *Indian J. Chem., Sect. B: Org. Chem. Incl. Med. Chem.*, 2006, **45**(12), 2690–2695, DOI: [10.1002/chin.200714170](#).
- 35 R. N. Rao, B. MM, B. Maiti, R. Thakuria and K. Chanda, Efficient Access to Imidazo[1,2-*a*]pyridines/pyrazines/pyrimidines via Catalyst-Free Annulation Reaction under Microwave Irradiation in Green Solvent, *ACS Comb. Sci.*, 2018, **20**(3), 164–171, DOI: [10.1021/acscombsci.7b00173](#).
- 36 B. Das, N. Bhunia and M. Lingaiah, A Simple and Efficient Metal-Free Synthesis of Tetrasubstituted Pyrroles by



- Iodine-Catalyzed Four-Component Coupling Reaction of Aldehydes, Amines, Dialkyl Acetylenedicarboxylates, and Nitromethane<sup>1</sup>, *Synthesis*, 2011, **2011**(21), 3471–3474, DOI: [10.1055/s-0030-1260228](https://doi.org/10.1055/s-0030-1260228).
- 37 R. Aggarwal, M. Hooda, P. Kumar, *et al.*, Visible-Light-Prompted Synthesis and Binding Studies of 5,6-Dihydroimidazo[2,1-*b*]thiazoles with BSA and DNA Using Biophysical and Computational Methods, *J. Org. Chem.*, 2022, **87**(6), 3952–3966, DOI: [10.1021/acs.joc.1c02471](https://doi.org/10.1021/acs.joc.1c02471).
  - 38 G. Sumran and R. Aggarwal, A convenient [hydroxy(tosyloxy) iodo]benzene-mediated one-pot synthesis of 2-arylimidazo [2,1-*b*]benzothiazoles]], *J. Sulfur Chem.*, 2015, **36**(2), 170–177, DOI: [10.1080/17415993.2014.996221](https://doi.org/10.1080/17415993.2014.996221).
  - 39 R. Aggarwal, M. Hooda, P. Kumar and M. C. Torralba, Visible-light-mediated regioselective synthesis of novel thiazolo[3,2-*b*] [1,2,4]triazoles: Advantageous synthetic application of aqueous conditions, *Org. Biomol. Chem.*, 2022, **20**(3), 584–595, DOI: [10.1039/d1ob02194j](https://doi.org/10.1039/d1ob02194j).
  - 40 S. Siddiqui, F. Ameen, S. ur Rehman, T. Sarwar and M. Tabish, Studying the interaction of drug/ligand with serum albumin, *J. Mol. Liq.*, 2021, **336**, 116200, DOI: [10.1016/j.molliq.2021.116200](https://doi.org/10.1016/j.molliq.2021.116200).
  - 41 R. Patel, N. Maurya, M. U. D. Parray, *et al.*, Esterase activity and conformational changes of bovine serum albumin toward interaction with mephedrone: Spectroscopic and computational studies, *J. Mol. Recognit.*, 2018, **31**(11), 2734, DOI: [10.1002/jmr.2734](https://doi.org/10.1002/jmr.2734).
  - 42 A. Anish Babu, K. Karthick, R. Subramanian and K. Swarnalatha, Exploring the structural interaction of BSA with amine functionalized ruthenium(II) metal complex, *J. Biomol. Struct. Dyn.*, 2020, **38**(13), 4032–4039, DOI: [10.1080/07391102.2019.1683071](https://doi.org/10.1080/07391102.2019.1683071).
  - 43 B. Pastrello, G. C. dos Santos, L. C. da Silva-Filho, A. R. de Souza, N. H. Morgon and V. F. Ximenes, Novel aminoquinoline-based solvatochromic fluorescence probe: Interaction with albumin, lysozyme and characterization of amyloid fibrils, *Dyes Pigments*, 2020, **173**, 107874, DOI: [10.1016/j.dyepig.2019.107874](https://doi.org/10.1016/j.dyepig.2019.107874).
  - 44 J. Lazniewska, M. Agostino, S. M. Hickey, *et al.*, Spectroscopic and Molecular Docking Study of the Interaction between Neutral Re(I) Tetrazolate Complexes and Bovine Serum Albumin, *Chem.-Eur. J.*, 2021, **27**(44), 11406–11417, DOI: [10.1002/chem.202101307](https://doi.org/10.1002/chem.202101307).
  - 45 M. H. Gehlen, The centenary of the Stern-Volmer equation of fluorescence quenching: From the single line plot to the SV quenching map, *J. Photochem. Photobiol. C Photochem. Rev.*, 2020, **42**, 100338, DOI: [10.1016/j.jphotochemrev.2019.100338](https://doi.org/10.1016/j.jphotochemrev.2019.100338).
  - 46 K. Paliwal, P. Haldar, P. K. S. Antharjanam and M. Kumar, Mixed Ligand Mononuclear Copper(II) Complex as a Promising Anticancer Agent: Interaction Studies with DNA/HSA, Molecular Docking, and In Vitro Cytotoxicity Studies, *ACS Omega*, 2022, **7**(25), 21961–21977, DOI: [10.1021/acsomega.2c02354](https://doi.org/10.1021/acsomega.2c02354).
  - 47 A. V. Ardhapure, V. Gayakhe, S. Bhilare, *et al.*, Extended fluorescent uridine analogues: synthesis, photophysical properties and selective interaction with BSA protein, *New J. Chem.*, 2020, **44**(34), 14744–14754, DOI: [10.1039/D0NJ02803G](https://doi.org/10.1039/D0NJ02803G).
  - 48 R. Ma, H. Pan, T. Shen, *et al.*, Interaction of Flavonoids from Woodwardia unigemmata with Bovine Serum Albumin (BSA): Application of Spectroscopic Techniques and Molecular Modeling Methods, *Molecules*, 2017, **22**(8), 1317, DOI: [10.3390/molecules22081317](https://doi.org/10.3390/molecules22081317).

