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## PAPER

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## 1. Introduction

Computational biology and bioinformatics play a major role in designing drug molecules and have the potential to speed up the drug discovery process. Molecular docking of the drug molecule with the receptor gives important information about drug–receptor interactions and is commonly used to find out the binding orientation of drug candidates to their protein targets in order to predict the affinity and activity.<sup>1</sup> Medicinal chemistry is a specialized science that has evolved to encompass a broad range of disciplines concerned with the identification, synthesis and development of drug-like compounds for therapeutic use. It needs a wide range of expertise, developed through years of training, dedication and learning from best practice in order to produce drugs that are good enough to enter clinical trials with patients.<sup>2</sup>

Quinazolinones and their derivatives are building blocks of approximately 150 naturally occurring alkaloids isolated from plants, animals and microorganisms. The quinazolinone nucleus has diverse pharmacological activities, including

# A correlation study of biological activity and molecular docking of Asp and Glu linked bishydrazones of quinazolinones†

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The present investigation involves the synthesis and spectroscopic and biological activity studies of the bishydrazones of quinazolinones derived from aspartic acid and glutamic acid. The antioxidant activities of the compounds were evaluated using DPPH, DMPD and ABTS radical scavenging assays whose results revealed that the IC<sub>50</sub> of compounds **6**, **7**, **11**, **12**, **20**, **21**, **25** and **26** was lower than those of the standard references. The anti-inflammatory activity was evaluated with a haemolysis assay using a human blood erythrocytes suspension and the results demonstrated that compounds **8**, **9**, **13**, **14**, **22**, **23**, **27** and **28** were excellent anti-inflammatory agents. In addition, the antibacterial and antifungal activities against various clinical pathogens of human origin revealed that compounds **7**, **9**, **12**, **14**, **21**, **23**, **26** and **28** possessed potent antimicrobial properties. Furthermore, to understand the correlation between biological activity and drug-receptor interaction, molecular docking was performed on the active sites of tyrosine kinase (PDB ID: 2HCK), cyclooxygenase-2 (PDB ID: 1CX2) and glucosamine-6-phosphate (GlcN-6-P) synthase (PDB ID: 2VF5) which showed good binding profiles with the targets that can potentially hold the title compounds. The correlation study revealed that compounds containing EDGs (-OH,  $-OCH_3$ ) were excellent antioxidants, compounds with EWGs (-Cl,  $-NO_2$ ) exhibited good anti-inflammatory activity and compounds bearing -OH and  $-NO_2$  groups were very good antimicrobials.

> antimicrobial,<sup>3</sup> anti-inflammatory,<sup>4</sup> antimalarial,<sup>5</sup> anticonvulsant,<sup>6</sup> antihypertensive,<sup>7</sup> anti-diabetic,<sup>8</sup> PARP inhibitory<sup>9</sup> and anticancer activity.<sup>10</sup> On the other hand, hydrazones are present in many bioactive heterocyclic compounds and are of wide interest because of their diverse biological and clinical applications. Moreover, hydrazones of quinazolinone derivatives exhibited enhanced biological activity, *viz.* antimicrobial,<sup>11</sup> antiinflammatory,<sup>12</sup> antiviral,<sup>13</sup> analgesic<sup>14</sup> and anticancer activity.<sup>15</sup> The structures of the biologically active quinazolinonehydrazones are shown in Fig. 1.

> Besides these, amino acids are endogenous substances which could be used for the modification of drug skeletons to promote the absorption of drugs.<sup>16</sup> In addition, the toxicity of drugs also could be reduced via the introduction of amino acids.17 Conjugation of amino acids/peptides with heterocycles,18,19 especially with a quinazolinone moiety,20-22 enhances the overall therapeutic properties of the molecule. Motivated by the aforementioned literature and the recent encouraging results from our research group,23-25 we have directed systematic efforts towards the synthesis of new bis-hydrazones of Asp and Glu linked quinazolinone derivatives followed by the study of their antimicrobial, antioxidant and anti-inflammatory activity. In addition, the results obtained were correlated with molecular docking studies in order to obtain some information about the binding efficiency and target interactions. The above are all collectively presented in this communication.

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Fig. 1 The structures of the biologically active quinazolinone-hydrazones

## 2. Results and discussion

#### 2.1. Chemistry

The synthesis of the desired compounds was achieved according to the steps illustrated in Scheme 1. 3-(4-Oxo-3,4dihydroquinazolin-2-yl)propanoic acid (QZN 1) and 4-(4-oxo-3,4-dihydroguinazolin-2-yl)butanoic acid (OZN 2) were synthesized by literature known methods.26-28 Conjugation of QZN 1/QZN 2 with p-TsOH·NH2-Asp(OBzl)-OBzl/HCl·NH2-Glu(OCH<sub>3</sub>)-OCH<sub>3</sub> was carried out using EDCI/HOBt as the coupling agent and NMM as the base to obtain 1, 2, 15 and 16. Next, these conjugates were refluxed with an excess of hydrazine hydrate to obtain the corresponding bis-hydrazides of the quinazolinones, 3, 4, 17 and 18. These bis-hydrazides were allowed to react with various substituted aldehydes to obtain the bishydrazones of the quinazolinone derivatives, 5-14 and 19-28. All the compounds were obtained in good yields. The structures of all the newly synthesized compounds were confirmed by IR, <sup>1</sup>H-NMR, <sup>13</sup>C-NMR and mass spectral analyses. The conjugation of the quinazolinones with the amino acids was confirmed by the appearance of an –NH peak at  $\delta$  8.54–8.15 and the absence of a COOH signal at  $\delta$  12.25 in the <sup>1</sup>H NMR spectrum. In the IR spectra, bands appeared at 3310  $\text{cm}^{-1}$  (-NH<sub>2</sub>) and 3217  $\text{cm}^{-1}$ (-NH) and in the PMR spectra, peaks appeared at  $\delta$  4.12–3.26

 $(-NH_2)$  and 9.11–8.88 (-NH) which indicated the conversion of an ester into a hydrazide. The formation of the hydrazones (-N=CH) was confirmed by the presence of an absorption band at 1612–1630 cm<sup>-1</sup> and a peak at  $\delta$  8.62–8.30. The presence of all requisite peaks and absence of extraneous peaks in the PMR and CMR spectra confirm the structures. Furthermore, the mass values obtained were in good agreement with the structures assigned (spectral data are provided in the ESI<sup>†</sup>).

#### 2.2. Biology

**2.2.1. Antioxidant activity.** The *in vitro* antioxidant activity of the synthesized compounds was evaluated with DPPH,<sup>29</sup> DMPD,<sup>30,31</sup> and ABTS<sup>32</sup> radical scavenging assays. The values were determined in terms of the  $IC_{50}$ , the concentration at which 50% of the radicals were scavenged, and were calculated to evaluate the antioxidant activity. The  $IC_{50}$  values were compared with those of the standards: ascorbic acid (AA) and gallic acid (GA). The results are tabulated in Table 1.

Most of the synthesized compounds showed antioxidant properties in all three radical scavenging assays. The conjugations of the QZN moiety with Asp (1 and 2) and Glu (15 and 16) showed antioxidant properties at higher concentrations which is in accordance with our earlier observations.<sup>33</sup> Conversion of



Scheme 1 Bis-hydrazones of quinazolinone derived from anionic amino acid linkers. Reagents and conditions: (i) NMM, EDCI/HOBt, 0 °C to rt, (ii)  $NH_2NH_2 \cdot H_2O$ , ethanol, reflux, 16 h, (iii) R-CHO, EtOH, reflux, 7–8 h.

Table 1 Antioxidant activity and docking studies of the synthesized quinazolinone derivatives<sup>a</sup>

	Antioxidant activity			Molecular docking studies with 2HCK protein				
Entry	DPPH, $IC_{50}$ (µg mL <sup>-1</sup> )	DMPD, $IC_{50}$ (µg mL <sup>-1</sup> )	ABTS, $IC_{50}$ (µg mL <sup>-1</sup> )	Docking score	H-bond interactions with $AA^b$ residues	$\pi$ -cation interactions	π–π stacking interactions	
1	>300	$280\pm5.23$	$290\pm3.26$	-3.865	Arg 136	Arg 76	NF	
2	$260\pm3.56$	>300	>300	-4.047	NF	NF	NF	
3	$250\pm2.21$	$240 \pm 1.35$	$180\pm1.65$	-4.545	Ala 96	NF	NF	
4	$190 \pm 1.40$	>300	$240 \pm 1.32$	-4.606	Ala 96, Asn 46	NF	NF	
5	$85\pm0.95$	$100 \pm 1.56$	$70 \pm 1.45$	-6.250	Asn 46	NF	NF	
6	$50\pm0.45$	$55\pm0.56$	$45\pm1.20$	-7.770	Ala 96, Asn 46	NF	NF	
7	$35\pm0.65$	$40\pm0.72$	$25\pm0.95$	-9.375	Asp 73, Asp 73, Asn 46	Arg 76	NF	
8	$120 \pm 1.26$	$140\pm2.36$	$150\pm0.68$	-5.066	Arg 76, Arg 136	NF	NF	
9	$180\pm0.65$	$160\pm0.65$	$145 \pm 1.98$	-5.983	Arg 136	NF	NF	
10	$70\pm0.65$	$95 \pm 1.67$	$100 \pm 1.85$	-6.193	NF	NF	NF	
11	$50\pm1.01$	$55\pm0.98$	$45\pm0.89$	-7.779	Arg 136	Arg 76	NF	
12	$25\pm0.26$	$30\pm0.65$	$40\pm0.65$	-9.007	Asp 73, Asp 73, Asn 46	NF	Arg 136	
13	$150\pm3.26$	$135\pm1.69$	$160 \pm 1.98$	-5.390	Arg 136, Asn 46	NF	NF	
14	$180\pm2.10$	$170\pm0.98$	$155\pm1.69$	-5.797	Arg 136, Asn 46	NF	NF	
15	$260\pm3.65$	>300	$285\pm2.65$	-3.865	Arg 136, Gly 77	NF	NF	
16	>300	$260\pm3.45$	>300	-4.188	NF	NF	NF	
17	$240 \pm 2.98$	$225 \pm 1.65$	$250\pm3.56$	-4.582	Ala 96, Asn 46, Gly 117	NF	NF	
18	>300	$245\pm2.56$	$280\pm2.45$	-4.606	Ala 96, Gly 117	NF	NF	
19	$95\pm0.95$	$80 \pm 1.20$	$75\pm0.56$	-6.149	Arg 136	NF	NF	
20	$45\pm0.88$	$55\pm0.75$	$40\pm0.45$	-7.983	NF	NF	NF	
21	$30\pm0.23$	$35\pm0.98$	$35\pm0.69$	-9.256	Arg 136, Asp 49, Asp 49, Glu 42, Glu 42	Arg 76	Arg 76	
22	$150\pm2.23$	$120\pm2.45$	$135\pm1.90$	-5.416	Arg 136, Asn 46	NF	NF	
23	$200\pm2.48$	$220 \pm 1.65$	$205 \pm 1.98$	-5.129	NF	NF	NF	
24	$75 \pm 1.45$	$80 \pm 1.70$	$100 \pm 1.65$	-6.132	Arg 136	NF	NF	
25	$35\pm0.68$	$45\pm1.20$	$50\pm1.95$	-8.097	Glu 50	NF	NF	
26	$25\pm0.24$	$30\pm0.50$	$45\pm0.64$	-9.510	Arg 136, Asp 73, Asp73, Glu 42	Arg 76	NF	
27	$135\pm1.98$	$150\pm2.25$	$160 \pm 1.40$	-5.309	Asn 46, Glu 117	NF	NF	
28	$165\pm0.65$	$140\pm2.50$	$130\pm2.60$	-4.973	Arg 136	NF	NF	
AA	$50\pm1.33$	$55\pm2.26$	$45\pm1.65$	-6.499	Asp 73, Val 71	NF	NF	
GA	$60 \pm 1.46$	$65 \pm 1.26$	$50\pm0.36$	-6.397	Asp 73	NF	NF	

 $a^{a}$  The values are the means of three determinations, the ranges of which are <5% of the mean in all cases.  $b^{b}$  Amino acid; NF: not formed, AA = ascorbic acid, GA = gallic acid.

the esters into hydrazides (3, 4, 17 and 18) slightly improved the antioxidant potential, which may be due to the radical scavenging ability of NH-NH2 groups.34 The enhanced activity of the resultant hydrazones could be due to the presence of groups on the phenyl ring. Those compounds that lack substituents on the phenyl ring (5, 10, 19 and 24) showed moderate antioxidant activity with IC<sub>50</sub> values ranging from 70–100  $\mu g m L^{-1}$ . Compounds with electron donating substituents like -OH and -OCH<sub>3</sub> on the phenyl ring (6, 7, 11, 12, 20, 21, 25 and 26) exhibited excellent radical scavenging activity with IC50 values ranging from 25–55  $\mu$ g mL<sup>-1</sup> in all three antioxidant assays and even better results compared to the standards AA (45-55 µg  $mL^{-1}$ ) and GA (50–65  $\mu$ g  $mL^{-1}$ ). In addition, the presence of OH slightly improved the antioxidant activity compared to that with OCH<sub>3</sub>. In contrast, compounds with electron withdrawing groups like -Cl and -NO<sub>2</sub> on the phenyl ring (8, 9, 13, 14, 22, 23, 27 and 28) showed the lowest antioxidant activity with  $IC_{50}$ values ranging from 120–220  $\mu$ g mL<sup>-1</sup>. On the basis of the above observation, it may be stated that compounds containing EDGs are excellent antioxidants,35-37 whereas the presence of EWGs decreases the antioxidant potential of the synthesized

analogues.<sup>22</sup> With respect to the length of the alkyl chain in the QZNs, there was no significance for the antioxidant potential. Replacement of Asp with Glu makes the antioxidant activity even better.

2.2.2. Anti-inflammatory activity. The human erythrocyte suspension was used to evaluate the in vitro anti-inflammatory activity of the synthesized molecules by employing a literature known method.38 A substantial number of compounds exhibited excellent to moderate activity compared to the standards, indomethacin (IM) and ibuprofen (IP). The results of the  $IC_{50}$ measurements were determined and are tabulated in Table 2. The compounds with electron withdrawing groups (8, 9, 13, 14, 22, 23, 27 and 28) showed excellent activity with IC<sub>50</sub> values of 45, 25, 50, 45, 50, 40, 45 and 35  $\mu$ g mL<sup>-1</sup>, respectively, which are more potent than those of indomethacin (55  $\mu g m L^{-1}$ ) and ibuprofen (50  $\mu$ g mL<sup>-1</sup>). Besides this, compounds with electron donating substituents like OH and OCH<sub>3</sub> (6, 7, 11, 12, 20, 21, 25 and 26) or without any substituents (5, 10, 19 and 24) showed moderate activity. This clearly indicates that the compounds with electron withdrawing groups (Cl, NO<sub>2</sub>) on the phenyl ring are better anti-inflammatory agents.39 Furthermore, NO2 which

#### Table 2 Anti-inflammatory activity and docking studies of the synthesized quinazolinone derivatives<sup>4</sup>

		Molecular docking studies with 1CX2 protein						
Entry	Anti-inflammatory activity, $IC_{50} (\mu g m L^{-1})$	Docking score	H-bond interactions with $AA^b$ residues	Electrostatic forces of attraction	$\pi$ -cation interactions	$\pi$ - $\pi$ stacking interactions		
1	$270 \pm 4.25$	-4.164	Lys 118	NF	NF	NF		
2	>300	-4.199	Lys 118	NF	NF	Phe 30		
3	$225\pm2.69$	-3.905	Lys 118	NF	NF	NF		
4	>300	-4.068	Lys 118	NF	NF	Phe 30		
5	$100\pm2.60$	-4.678	NF	NF	NF	Phe 30, Phe 30		
6	$75\pm1.50$	-4.879	Asp, 87, Lys 118	NF	NF	NF		
7	$80\pm2.06$	-4.976	Ala 161, Asp 90, Asp 90, Asp 120, Asp 120, Lys 118	NF	NF	Phe 30		
8	$45\pm0.25$	-8.576	NF	NF	NF	Phe 30		
9	$25\pm0.56$	-8.908	Cys 20, Lys 118, Lys 118	Asp 13, Asp 87, Lys 118, Lys 118	NF	Tyr 34		
10	$120\pm1.40$	-4.810	Lys 118, Asp 87	NF	NF	Phe 30		
11	$80 \pm 1.50$	-4.759	NF	NF	NF	NF		
12	$65\pm1.60$	-4.719	Ala 161, Asp 120, Asp 120, Glu 32, Glu 32,	NF	Lys 118	Phe 30		
13	$50\pm2.95$	-7.810	Asp 87	NF	NF	His 126, Phe 30		
14	$45 \pm 1.28$	-8.027	Asp 87, Cys 20, Lys 118, Ser 88	Asp 13, Asp 87, Asp 90, Lys 118	NF	NF		
15	$270\pm2.90$	-3.799	Lys 118, Gly 17	NF	NF	NF		
16	>300	-3.850	Lys 118	NF	NF	NF		
17	$240\pm3.40$	-4.116	Asp 120, Lys 118, Lys 162	NF	Lys 118	NF		
18	$260\pm2.75$	-3.630	Asp 87, Lys 118, Thr 127	NF	NF	Phe 30		
19	$150\pm2.35$	-4.378	NF	NF	NF	NF		
20	$80\pm0.54$	-4.646	NF	NF	Lys 118	Phe 30		
21	$95\pm0.45$	-4.506	Asp 87, Asp 87, Asp 124, Lys 118, Lys 162, Lys 162	NF	NF	NF		
22	$50\pm1.20$	-7.843	Asp 87	NF	NF	Phe 30		
23	$40\pm0.64$	-8.023	Asp 87, Asp 87, Lys 118	Asp 13, Lys 118, Lys 162	NF	Phe 30		
24	$115\pm0.75$	-4.378	Asp 87	NF	NF	His 126		
25	$75\pm0.80$	-4.482	Asp 87	NF	Lys 118	Phe 30		
26	$90\pm2.20$	-4.472	Asp 87, Asp 87, Asp 90, Ser 88	NF	NF	Phe 30		
27	$45\pm0.90$	-8.143	Asp 87, Lys 118	NF	NF	Phe 30		
28	$35\pm0.64$	-7.928	Asp 87	Tyr 34	Lys 118	Tyr 34		
IM	$55\pm2.01$	-5.074	NF	NF	Lys 118	Phe 30		
IP	$50\pm1.26$	-	-	-	-	-		

<sup>*a*</sup> The values are the means of three determinations, the ranges of which are <5% of the mean in all cases. <sup>*b*</sup> Amino acid; NF: not formed; '-': no activity/not analyzed, IM = indomethacin, IP = ibuprofen.

is a powerful EWG is better at enhancing the anti-inflammatory activity than Cl.

**2.2.3.** Antimicrobial activity. The efficacy of the synthesized compounds as antimicrobials was tested in antibacterial<sup>40</sup> studies against different strains of pathogen using both Gram negative bacteria, namely *Escherichia coli* (*E. coli*), and Gram positive bacteria, *Staphylococcus aureus* (*S. aureus*), and in antifungal<sup>41</sup> studies against *Fusarium moniliforme* (*F. moniliforme*) and *Aspergillus niger* (*A. niger*). The results obtained as the zone of inhibition (mm) are presented in Table 3. Streptomycin (SM) and bavistin (BS) served as the standard drugs for the antibacterial and antifungal studies, respectively.

Most of the synthesized compounds showed promising antimicrobial activity with few exceptions. The results of the antimicrobial studies showed that all the analogues exhibited the same trend for both antibacterial and antifungal activity. The quinazolinone-amino acid conjugates (1, 2, 15 and 16) exhibited a much smaller zone of inhibition. When C-terminal benzyl ester/methyl ester groups were converted into hydrazides (4, 5, 17 and 18), there was a slight enhancement in the activity. This may be due to the increase in the polarity of the compounds, which would help the molecule to penetrate more through the cell membrane of microbes and thereby inactivate them.<sup>20</sup> There was a drastic enhancement in the activity when the hydrazides were converted into hydrazones and we observed that the nature of the substituents present on the phenyl ring affected the biological activity of the compounds to a greater extent.

Among the bis-hydrazones, the compounds with hydroxyl groups (7, 12, 21 and 26) and nitro groups (9, 14, 23 and 28) on the phenyl ring showed potent antimicrobial activity compared to the standard drugs. Compounds bearing methoxy (6, 11, 20 and 25) and chloro (8, 13, 22 and 27) groups showed moderate antibacterial properties. Those compounds (5, 10, 19 and 24)

#### Table 3 Antimicrobial activity and molecular docking studies of the synthesized analogues with 2VF5

	Antibacterial activity <sup>a</sup> (mm)		Antifungal activity <sup><i>a</i></sup> (mm)		Molecular docking studies with 2VF5			
							* $\pi$ -cation interactions/	
Entry	E. coli	S. aureus	F. moniliforme	A. niger	Docking score	Hydrogen bond interactions with amino acid residues	electrostatic forces of attraction	
1	$04\pm0.21$	$02\pm0.32$	$05\pm0.12$	$02\pm0.41$	-6.150	Ala 602, Glu 488, Ser 401	*Lys 603	
2	$03\pm0.38$	NA	$02\pm0.35$	NA	-6.196	Glu 488, Gly 301, Ser 401	NF	
3	$09\pm0.65$	$07\pm0.65$	$06 \pm 0.65$	$04\pm0.68$	-7.990	Ala 602, Lys 603, Ser 347, Gln 348, Gln 348, Ser 303, Leu 346, Cys 300	NF	
4	$07\pm0.45$	NA	$05\pm0.38$	$03\pm0.12$	-7.804	Thr 302, Ser 401, Leu 346, Cys 300, Gln 348, Thr 352, Ser 303, Ser 347	NF	
5	$14\pm0.32$	$10 \pm 0.52$	$11\pm0.42$	$09\pm0.25$	-8.203	Ala 602, Val 399, Lys 603, Gln 348, Ser 401, Ser 347	NF	
6	$16\pm0.54$	$14\pm0.16$	$15\pm0.14$	$18\pm0.54$	NF	NF	NF	
7	$28\pm0.32$	$25\pm0.25$	$26\pm0.28$	$24\pm0.32$	-12.238	Asp 354, Asp 354, Glu 488, Ala 602, Ser 401, Gln 348, Ser 303, Gly 301, Thr 352	NF	
8	$18\pm0.18$	$16\pm0.45$	$17\pm0.36$	$15\pm0.65$	-8.597	Glu 488	NF	
9	$21\pm0.23$	$19\pm0.32$	$23\pm0.23$	$24\pm0.23$	-10.017	Asp 354, Ala 602, Ala 602	<sup>#</sup> Asp 354, <sup>#</sup> Asp 354	
10	$10\pm0.56$	$09\pm0.23$	$11\pm0.65$	$12\pm0.45$	NF	NF	NF	
11	$16\pm0.36$	$15\pm0.55$	$17\pm0.11$	$18\pm0.78$	NF	NF	NF	
12	$26\pm0.65$	$24 \pm 0.85$	$25\pm0.45$	$27 \pm 0.28$	-10.553	Asp 354, Asp 354, Val 399, Cys 300, Ser 347, Ser 349, Ala 602, Asn 600	NF	
13	$18\pm0.12$	$17\pm0.35$	$19\pm0.26$	$20\pm0.45$	NF	NF	NF	
14	$22\pm0.26$	$20\pm0.38$	$21\pm0.65$	$22\pm0.22$	-9.835	Lys 603, Ser 303, Asp 354	<sup>#</sup> Asp 354, <sup>#</sup> Asp 354	
15	$02\pm0.35$	NA	$05\pm0.32$	$01 \pm 0.18$	-5.966	Ala 602, Ala 602, Val 605, Thr 352, Ser 347, Ser 349, Gln 348, Ser 303	NF	
16	NA	$01\pm0.08$	NA	$04\pm0.26$	-5.816	Ala 602, Thr 302, Ser 349, Ser 347	NF	
17	$04\pm0.21$	$03 \pm 0.21$	$05\pm0.33$	$08\pm0.21$	-7.199	Ala 602, Ala 602, Leu 346, Cys 300, Gln 348, Ser 347, Lys 603, Lys 603	NF	
18	$05\pm0.65$	NA	$07\pm0.29$	NA	-7.170	Ala 602, Ala 602, Cys 300, Ser 347, Ser 349, Gln 348, Ser 401, Ser 401, Ser 303	NF	
19	$14\pm0.35$	$11\pm0.65$	$13\pm0.26$	$14\pm0.29$	-8.083	Gln 348, Val 399, Thr 302, Ala 602, Ser 401	NF	
20	$18\pm0.16$	$16\pm0.19$	$15\pm0.32$	$16\pm0.11$	-8.317	Ala 602, Lys 603, Ser 303	NF	
21	$28\pm0.22$	$26 \pm 0.35$	$25\pm0.24$	$24 \pm 0.26$	-9.339	Thr 352, Ser 303, Asn 600, Ala 602, Asn 305, Gly 301	NF	
22	$19\pm0.32$	$20 \pm 0.17$	$18 \pm 0.35$	$17 \pm 0.56$	-8.432	Thr 352, Lys 603, Thr 302, Val 399, Ala 602	NF	
23	$22\pm0.56$	$24\pm0.29$	$21\pm0.65$	$23\pm0.36$	-8.969	Thr 302, Gln 348, Ser 303, Ser 347, Thr 352	<sup>#</sup> Glu 488	
24	$13\pm0.25$	$10\pm0.56$	$15\pm0.36$	$14\pm0.33$	NF	NF	NF	
25	$16\pm0.32$	$17\pm0.44$	$18\pm0.14$	$15\pm0.38$	NF	NF	NF	
26	$26\pm0.35$	$22\pm0.23$	$24\pm0.36$	$25\pm0.36$	-9.282	Asn 305, Thr 352, Gln 348	NF	
27	$19\pm0.26$	$16\pm0.66$	$19\pm0.56$	$20\pm0.11$	NF	NF	NF	
28	$24\pm0.12$	$23\pm0.33$	$24\pm0.45$	$22\pm0.65$	NF	NF	NF "	
SM	$15\pm0.36$	$13 \pm 0.69$	—	—	-9.517	Gly 301, Asp 354, Asp 354, Cys 300, Thr 302, Ala 602, Ala 602, Glu 488	"Glu 488	
BS	—	—	$14\pm0.65$	$13\pm0.36$	NF	NF	NF	

 $^{a}$  The values are the means of three determinations, the ranges of which are <5% of the mean in all cases, NF = not formed, SM = streptomycin, BS = bavistin.

which did not have any substituents on the phenyl ring showed the lowest activity. Thus the order of activity based on the groups attached to the phenyl ring of hydrazone derivatives was found to be  $OH > NO_2 > Cl > OCH_3 > H$ . An increase in the alkyl chain length of quinazolinone caused a slight decrease in the antimicrobial activity which is in good agreement with our earlier report.<sup>20</sup> Replacement of Asp with Glu resulted in a slight decrease in the antimicrobial properties which may be attributed to the increase in hydrophobicity caused by the longer carbon chain in Glu.

#### 2.3. Molecular docking studies

The results of the molecular docking studies of the title compounds with respect to the antioxidant, anti-inflammatory and antimicrobial properties are presented below.

Molecular docking was performed on the active site of tyrosine kinase (PDB ID: 2HCK) with the synthesized ligands (1–28) in order to determine the possible binding interactions of highly potent molecules. Most of the compounds showed good docking scores and potent interactions with different amino acid residues and the results are tabulated in Table 1. Among

#### Paper

the series of compounds, those possessing EDGs, especially hydroxyl groups (7, 12, 21 and 26), gave the highest docking scores. The binding interactions (2D and 3D) of 7, 12, 21 and 26 are displayed in Fig. 2. Compound 7 showed hydrogen bond interactions with Asp 73, Asp 73 and Asn 46 and  $\pi$ -cation interactions with Arg 76 and compound **12** showed hydrogen



Fig. 2 2D and 3D images of compounds 7, 12, 21 and 26 with 2HCK.

bond interactions with Asp 73, Asp 73, and Asn 46 and  $\pi$ - $\pi$  stacking interactions with Arg 136. Compound **21** showed hydrogen bond interactions with Arg 136, Asp 49, Asp 49, Glu 42

and Glu 42 and  $\pi$ -cation and  $\pi$ - $\pi$  stacking interactions with Arg 76 and compound **26** exhibited hydrogen bond interactions with Arg 136, Asp 73, Asp 73 and Glu 42 and  $\pi$ -cation



Fig. 3 2D and 3D images of compounds 9, 14, 23 and 28 with 1CX2.

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interactions with Arg 76. These molecules showed the highest docking scores because of the involvement of hydroxyl groups on the phenyl ring in hydrogen bond interactions. To rationalize the anti-inflammatory potential of the synthesized compounds (1–28), we conducted docking studies on the crystal structure of cyclooxygenase-2 (PDB ID: 1CX2).



Fig. 4 2D and 3D images of compounds 7, 9, 21 and 23 with 2VF5 protein.

Most of the compounds exhibited a good docking score and different kinds of interaction with amino acid residues and these are tabulated in Table 2. Analogues with EWGs, particularly with NO<sub>2</sub> (9, 14, 23 and 28) on the phenyl ring, displayed good interactions with amino acid residues and had the highest docking scores. Molecular docking studies provided an interaction map of cyclooxygenase-2 with 9, 14, 23 and 28 and the results are presented in Fig. 3. Compound 9 showed hydrogen bond interactions with Cys 20, Lys 118 and Lys 118, electrostatic forces of attraction with Asp 13, Asp 87, Lys 118 and Lys 188 and  $\pi$ - $\pi$  stacking interactions with Tyr 34; and compound 14 showed hydrogen bond interactions with Asp 87, Cys 20, Lys 118 and Ser 88 and electrostatic interactions with Asp 13, Asp 87, Asp 90, Lys 118 and Lys 118. Meanwhile, compound 23 displayed hydrogen bond interactions with Asp 87, Asp 87 and Lys 118, electrostatic interactions with Asp 13, Lys 118 and Lys 162 and  $\pi$ - $\pi$  stacking interactions with Phe 30, whereas compound 28 showed hydrogen bond interactions with Asp 87,  $\pi$ -cation interactions with Lys 118 and Tyr 34 and  $\pi$ - $\pi$  stacking interactions with Tyr 34. The NO<sub>2</sub> groups present on the phenyl ring involved in electrostatic forces of attraction and  $\pi$ -cation interactions led to an enhanced docking score.

In order to gain insight into the exact binding location of the ligands with the protein, all the synthesized molecules (1-28) including the standards were subjected to molecular docking with the active site of GlcN-6-P synthase (PDB ID: 2VF5). They showed good binding interactions with the surrounding amino acid residues. The docking score and the interacting amino acids are tabulated in Table 3. Most of the synthesized compounds exhibited well established bonds with one or more amino acids in the receptor active pocket of 2VF5 protein. The potential of the compounds as antimicrobial agents was determined based on the docking scores. The docking scores with 2VF5 protein ranged from -12.238 to -5.816 and the highest negative value indicates the best docked ligand to the targeted site. The 2D and 3D images of compounds 7, 9, 21 and 23 are shown in Fig. 4. Aspartic acid linked hydrazones (7) showed hydrogen bond interactions with Asp 354, Asp 354, Glu 488, Ala 602, Ser 401, Gln 348, Ser 303, Gly 301 and Thr 352 and compound 9 showed hydrogen bond interactions with Asp 354, Ala 602 and Ala 602 and electrostatic forces of interaction with Asp 354 and Asp 354. On the other hand, glutamic acid linked hydrazones (21) displayed hydrogen bond interactions with Thr 352, Ser 303, Asn 600, Ala 602, Asn 305 and Gly 301 and compound 23 showed hydrogen bond interactions with Thr 302, Gln 348, Ser 303, Ser 347 and Thr 352 and electrostatic forces of interaction with Glu 488. The involvement of the hydroxyl groups in hydrogen bond interactions and the nitro groups in electrostatic forces of attraction enhanced the docking scores of these analogues.

## 3. Conclusions

In summary, we have developed a novel class of antioxidant, anti-inflammatory and antimicrobial agent with quinazolinonehydrazones linked *via* Asp and Glu as connectors. In addition, molecular docking studies were performed on all the synthesized compounds. The correlation between molecular docking studies and biological assays suggested that compounds 6, 7, 11, 12, 20, 21, 25 and 26 with electron donating groups (OH, OCH<sub>3</sub>) exhibited stronger radical scavenging activity than ascorbic acid and gallic acid. Compounds 8, 9, 13, 14, 22, 23, 27 and 28 with electron withdrawing groups (Cl, NO<sub>2</sub>) demonstrated better *in vitro* anti-inflammatory activity than indomethacin and ibuprofen. The compounds with electron donating hydroxyl groups (7, 12, 21 and 26) and electron withdrawing nitro groups (9, 14, 23 and 28) exhibited potent antimicrobial properties among this series of compounds.

## Conflicts of interest

There are no conflicts to declare.

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