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A structural blueprint for antibacterial discovery: microwave- and ultrasound-assisted synthesis of pyrrolidine-fused quinoxalines as novel inhibitors of DNA gyrase and biofilm

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In order to lessen the severity of infectious diseases, anti-infective agents—drugs that prevent, combat, or control infections brought on by microorganisms—are essential in contemporary medicine. To tackle antimicrobial resistance, this project intends to design and synthesize hybrid compounds that contain pyrrolidine, quinoxaline and a hydrazinyl bridge, and assess the antimicrobial and antifungal properties of these compounds against a variety of pathogenic strains. The bactericidal properties of hybrids **24**, **27**, and **29** against *E. coli* were verified. The MIC of 12.5 μM was shown by hybrids **24**, **25**, and **31**, which suggests bactericidal hybrids are effective against *P. aeruginosa* at greater concentrations. In comparison to Levofloxacin, treatment with all hybrids produced an 89–92% reduction in biofilm formation at 90% MIC. Eight hybrids' killing kinetics against *P. aeruginosa* were time-dependent, with an abrupt decrease in CFU number observed at higher concentrations. While 4-fold and 8-fold MICs resulted in nearly total bacterial eradication, primary bacterial elimination happened after three hours. The most effective DNA gyrase inhibitors were hybrids **25**, **28**, and **31**; their IC_{50} values were significantly less than that of ciprofloxacin (77.3, 87.6, and 65.5 μM , respectively). To determine the best drug-like qualities, the study examined the physicochemical and pharmacokinetic features of active compounds. Molecular docking simulation experiments were also conducted to comprehend the binding interactions and mechanisms of action of these hits.

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1. Introduction

Diseases caused by infectious bacteria, viruses, and fungi have appeared and are continuing to grow in tandem with daily life.^{1–4} The need for novel antimicrobial drugs with lower toxicity and resistance strain activity appears unresolved.^{5–8}

Although many antibacterial medications are already accessible, microorganism resistance to antibiotics remains an immense issue.^{9–12} Compounds that are heterocyclic are frequently included in numerous synthetic and biological fields

because of their ubiquitous occurrence in nature.^{13–16} Amongst the several scaffolds considered for the prospective pharmacological applications of them, nitrogen-containing heterocyclic frameworks have acquired a lot of focus owing to the diverse spectrum of biological processes,^{17–19} comprising antifungal,^{20,21} antimicrobial,^{22,23} anticancer,^{24,25} antiviral,^{26,27} and antidiabetic properties.^{28,29}

A useful pharmacophore in medicinal science, quinoxaline, is a bicyclic molecule constructed when a pyrazine ring and a benzene ring combine.^{30–33} Its distinct structure permits a variety of alterations, producing molecules with ameliorated biological activity.^{34–37} Due to its capacity to communicate with the essential cellular structures and microbial enzymes, quinoxaline derivatives have recently been shown to potentially be potent antimicrobial drugs.³⁸ The naturally occurring antibiotic phenazinomycin, which has anticancer properties, is another antibacterial medication containing a quinoxaline scaffold.^{39–43}

Additionally, pyrrolidine ring piqued the curiosity of scientists because of its strong antibacterial qualities.^{44–47} Numerous quinolone antibiotics, including Gemifloxacin and Moxifloxacin, include pyrrolidine.^{48–50} In addition, the newly approved drug vonoprazan, which destroys *Helicobacter-pylori*, has a scaffold

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with a pyrrole ring.^{51–53} Including an azo part to the compounds which are heterocyclic enhances the drug's biological impact.^{27,54–57}

DNA gyrase is a potential target for antibacterial therapy, a crucial bacterial enzyme associated with transcription, recombination, and DNA replication.⁵⁸ The effective prevention of bacterial growth is achieved by DNA gyrase inhibitors, which disrupt DNA supercoiling and replication.⁵⁹ Since DNA gyrase is essential for bacterial life yet is absent from human cells, it is a desirable target for targeted antimicrobial therapy.⁶⁰

The notion that bacteria must attain a particular cell density in infectious illnesses before they may establish virulence factors and interfere with the host defences is widely recognized.^{61,62} Pathogenic bacteria have evolved a special mechanism known as “biofilm formation” to withstand the lethal assaults of oxidative stress, nutritional deprivation, and the host immune system or antibiotics. Microorganisms form a biofilm while they latch on to an abiotic or biotic surface in a matrix of extracellular polymeric substance (EPS). The ability of *Candida* species forge biofilms, in addition to bacteria, increases the possibility of resistance to existing antifungal medications.^{63,64} The need for novel molecules or hybrids of already-existing chemicals that can suppress or destroy the biofilm matrix is critical because there is currently no treatment option for biofilms in clinical settings.

The purpose of this work is to use pyrrolidine-fused quinoxaline employing azo bridge to produce a variety of novel hybrid compounds, as seen in Fig. 1. The evaluation of the novel hybrids involved examining their DNA gyrase inhibitory, antibacterial, and antifungal properties. The antibiofilm activity of each bactericidal derivative was examined. Each compound was examined for pharmacokinetic and physicochemical characteristics. Lastly, an evaluation of binding mode simulations through docking for the most successful samples were carried out.

2. Findings and analysis

2.1. Chemical synthesis

The methods indicated in Schemes 1–3 were used to synthesis the target compounds.

Benzene-1,2-diamine **1** was refluxed with oxalic acid **2** using water as a solvent in presence of conc. HCl affording 1,4-dihydroquinoxaline-2,3-dione **3**. Further reaction of **3** with thionyl chloride resulted in 2,3-dichloroquinoxaline **5**. Hydrazinylquinoxaline **4** or **6** were obtained by refluxing **3** or **5** with hydrated-hydrazine, respectively, in ethyl alcohol for 24 h, accordingly as represented in Scheme 1.⁶⁵

Furthermore, the reaction of itaconic acid **7** with various aniline derivatives **8–15**, afforded the 5-oxo-pyrrolidine derivatives **16–23** (Scheme 2). The reaction was conducted using three methods implementing the principles of green chemistry (discussed in the experimental section). Using ultrasound-assisted reaction afforded the highest yield and consumed the least time. The reaction time and yield are given in Table 1. Structure elucidation using ¹H NMR confirmed the structures of oxo-pyrrolidine derivatives **16–23**. The COOH proton appeared as a singlet-signal at approximately δ 12 part per million (ppm). The hydrogens for the two methylene carbons appeared as four doublet of doublet signals each integrating one proton at approximately δ 2 and 4 ppm. The methine hydrogens from pyrrolidine showed up as a multiplet at approximately δ 3 ppm.

The targeted quinoxaline-pyrrolidine hybrids **24–31** were afforded by refluxing the respective pyrrolidine fragment with hydrazinyl quinoxaline **4** or **6** (Scheme 3). A characteristic singlet appeared in the ¹H NMR spectra of hybrids **24–27** at approximately δ 11–12 ppm representing the 2-OH proton on the quinoxaline ring. In all synthesized hybrids, –NH–NH–CO showed as doublet at δ 11 ppm, while –NH–NH–CO showed as a doublet roughly between δ 5 and 8 ppm. The protons of the pyrrolidine ring exhibited the identical signal pattern in all

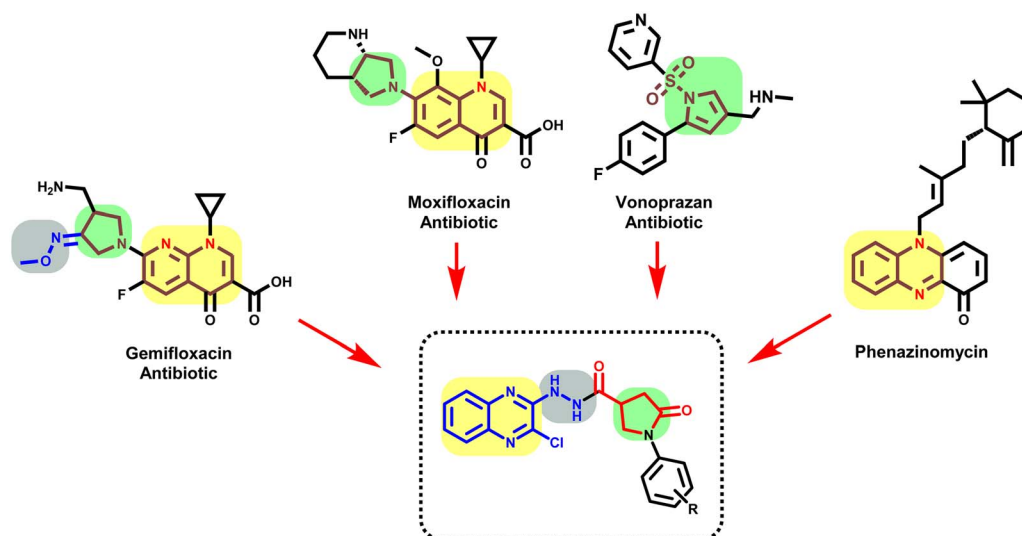
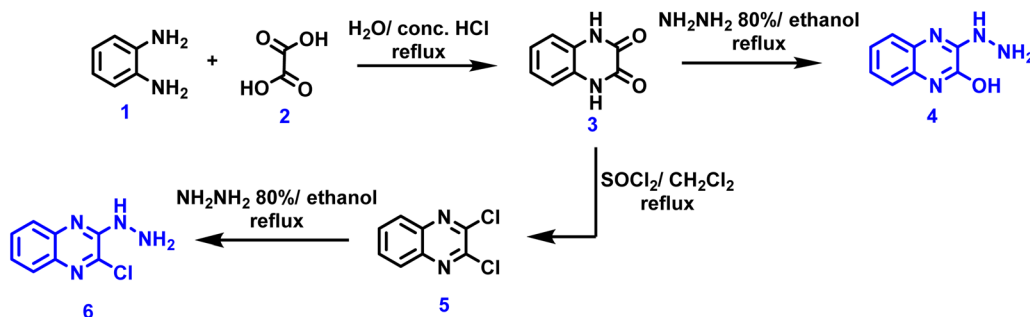
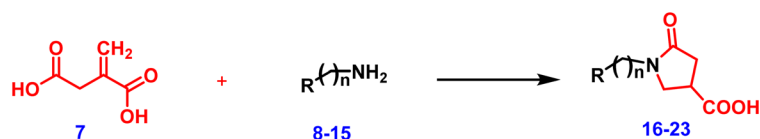


Fig. 1 Schematic representation of hybridized substances featuring quinoxaline, pyrrole/pyrrolidine, azo-bridge, and various substitutes.





Scheme 1 Creation of hydrazinylquinoxaline (4) and (6).



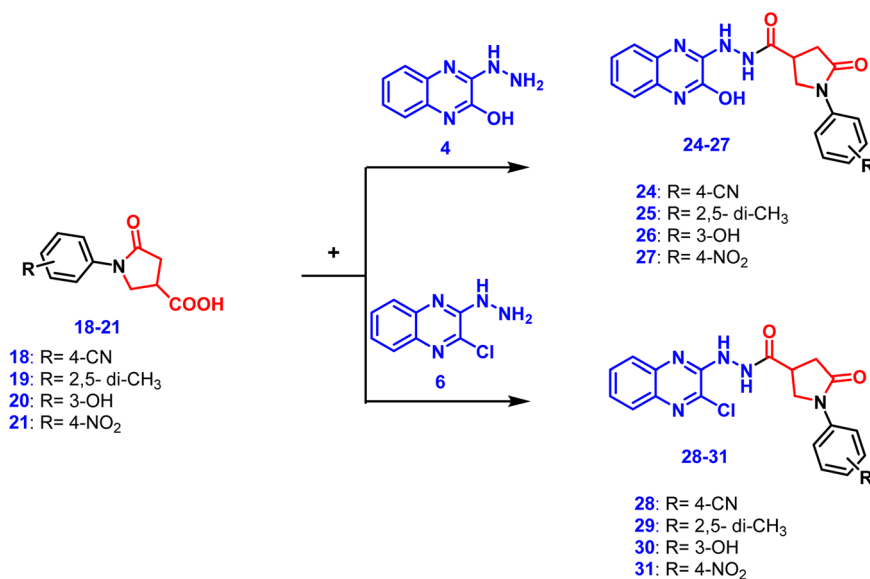
- 8,16:** $n=0$, R= 3-Cl phenyl
9,17: $n=0$, R= 4-Cl phenyl
10,18: $n=0$, R= 4-CN phenyl
11,19: $n=0$, R= 2,5- di-CH₃ phenyl
12,20: $n=0$, R= 3-OH phenyl
13,21: $n=0$, R= 4-NO₂ phenyl
14,22: $n=0$, R= 1-naphthyl
15,23: $n=1$, R= phenyl

Scheme 2 Manufacturing of oxo-pyrrolidine fragments 16–23.

synthesized hybrids. The ¹³C NMR of all synthesized hybrids revealed two characteristic signals representing the two C=O groups at δ between 170 and 175 ppm.

The ¹H NMR of hybrids 25 and 29 demonstrated two characteristic signals between δ 2 and 2.3 ppm, each integrating for 3 protons representing the two methyl groups. Additionally, the

two methyl groups were shown as two signals in ¹³C NMR at approximately δ 16 and 19 ppm. The ¹H NMR of hybrids 26 and 30 a signal representing the phenolic-OH between delta (δ) 7 and delta (δ) 8 part per million. The ¹³C NMR of hybrids 24 and 28 confirmed the presence of the cyano groups at approximately δ 118 ppm.



Scheme 3 Preparation of quinoxaline-pyrrolidine conjugates 24–31.



Table 1 Effect of reaction green method A, B and C on reaction time and yield

Compound	Reflux in water		Ultrasound		Microwave	
	Time	Yield %	Time	Yield %	Time	Yield %
16	24 h	56%	6 min	91%	12 min	73%
17	24 h	44%	4.5 min	95%	13 min	77%
18	24 h	62%	5 min	90%	12 min	84%
19	24 h	50%	5.5 min	92%	11 min	79%
20	24 h	55%	4 min	89%	15 min	86%
21	24 h	58%	5.5 min	88%	14 min	73%
22	24 h	65%	5 min	92%	15 min	88%
23	24 h	57%	4 min	90%	12 min	84%

2.2. In vitro assessment

2.2.1. Agar well-diffusion zones of inhibition (ZOI). At a dosage of 5 mM, the eight novel conjugates were screened for antibacterial activity using agar well diffusion, which revealed varying antimicrobial action in opposition to the assessed bacterial and fungal organisms (Table 2). With the exception of compound **29**, every synthetic derivative was found to promote the growth of *E. coli*. ZOI ≥ 15 mm at a dose of 5 mM. Remarkably, hybrids **24**, **26**, **27**, **28**, **30**, and **31** displayed *E. coli* ZOI that was greater than 17.6 ± 0.33 mm, superior to that of amoxicillin. With ZOI ranging from 18 to 23 mm, *P. aeruginosa* illustrated higher sensitivity to all synthesized hybrids than to Amoxicillin (15.3 ± 0.33 mm), with the exception of the formerly described inactive hybrid **29**.

Additionally, the ZOI values of hybrids **26**, **28**, **30**, and **31** ranged from 20 to 23 mm, indicating a greater reduction in *P. aeruginosa* growth than that of Levofloxacin (19.6 ± 1.3 mm). Out of all the synthetic derivatives, hybrids **24**, **25**, **26**, and **30** showed the highest ZOI against *K. pneumoniae*, with values ranging from 17 to 21 mm, outperforming Amoxicillin (16.3 ± 0.33 mm). Only hybrid **31** showed the greatest ZOI in the case of *A. baumannii*, with a value of 17.16 ± 0.44 mm surpassing that

of Amoxicillin (15.6 ± 0.33 mm), while only hybrid **28** was on par with Amoxicillin.

Nevertheless, the hybrids under investigation showed modest antibacterial efficacy against MSSA. With ZOI varying from 13 to 15 mm, all hybrids excluding **25** showed considerable antifungal efficacy against *C. albicans*, which is comparable to the common antifungal medication Fluconazole (19 ± 0.57 mm).

2.2.2. Assessment of MIC, MBC and MFC. The investigated active compounds were further tested for their minimum inhibitory concentration (MIC), minimum bactericidal concentration (MBC) and minimum fungicidal concentration (MFC). The values of the MIC, MBC, and MFC as well as the computed MBC/MIC (or MFC/MIC) ratio of the active hybrids against the aforementioned bacterial and fungal organisms, are seen in Table 3. Every synthetic chemical demonstrated broad-spectrum anti-infective activity. The strongest effect on MIC for *E. coli*, was observed upon administration of **29** with values of $6.25 \mu\text{M}$, Verifying advantage over Levofloxacin. In MBC, the same pattern was noted, using value of $12.5 \mu\text{M}$. Additionally, **25**, **26**, **28**, and **30** still achieved excellent MIC value of $12.5 \mu\text{M}$ against *E. coli*. Based on calculated MBC/MIC ratio (*R*), values of **2**, **24**, **27**, and **29** were confirmed as bactericidal agents against *E. coli*.

Hybrids **28** and **30** displayed substantial MIC value of $6.25 \mu\text{M}$, and MBC value of $25 \mu\text{M}$ and $50 \mu\text{M}$, against *P. aeruginosa*, resulting in (*R*) value of 4 and 8, indicating bacteriostatic activity. Moreover, **24**, **25** and **31** displayed MIC of $12.5 \mu\text{M}$, whereas the calculated (*R*) indicate bactericidal hybrids at higher concentrations.

In case of *K. pneumoniae*, **24** and **26** displayed excellent MIC value of $12.5 \mu\text{M}$, and a much higher MBC value resulting in a ratio above 2 indicating bacteriostatic hybrids. Compounds **28**, **29**, and **31** demonstrated moderate MIC value of $25 \mu\text{M}$. Hybrid **28** emerged as the most potent derivative against *A. baumannii*, with MIC and MBC values of $6.25 \mu\text{M}$ and $12.5 \mu\text{M}$, respectively, thereby identifying it as a bactericidal contender.

Table 2 ZOI diameter (mm) of the investigated hybrids, Levofloxacin, Amoxicillin and Fluconazole (at 5 mM)^a

Cpd. no.	Average inhibition zone diameter (mm)					
	Bacteria (Gram negative)				Bacteria (Gram positive)	Fungi
	<i>E. coli</i>	<i>P. aeruginosa</i>	<i>K. pneumoniae</i>	<i>A. baumannii</i>	MSSA	<i>C. albicans</i>
24	20.83 ± 0.60	18 ± 0.57	20.3 ± 1.2	—	—	13.8 ± 0.16
25	16.3 ± 0.33	19.6 ± 0.33	17.8 ± 0.16	—	—	—
26	19 ± 0.01	20.16 ± 0.6	20.6 ± 0.33	—	—	15 ± 0.01
27	19.3 ± 0.33	18.3 ± 0.88	15.5 ± 0.28	13.8 ± 0.16	—	14.6 ± 0.33
28	18.3 ± 0.66	22.3 ± 0.3	16.6 ± 0.33	15.5 ± 0.28	15.5 ± 0.28	13.8 ± 0.4
29	14.6 ± 0.33	14.6 ± 0.66	14 ± 0.01	13.8 ± 0.16	14.6 ± 0.66	14 ± 0.57
30	20.6 ± 0.33	21.3 ± 0.33	20.5 ± 0.28	15.3 ± 0.33	17 ± 0.01	15 ± 0.57
31	18 ± 0.57	20.6 ± 0.88	15.16 ± 0.16	17.16 ± 0.44	15.6 ± 0.33	14.6 ± 0.6
Levofloxacin	26.3 ± 0.33	19.6 ± 1.3	27.5 ± 0.28	30.5 ± 0.28	41.1 ± 0.01	NA
Amoxicillin	17.6 ± 0.33	15.3 ± 0.33	16.3 ± 0.33	15.6 ± 0.33	25 ± 0.01	NA
Fluconazole	NA	NA	NA	NA	NA	19 ± 0.57

^a DMSO was used as a solvent with no inhibition zone.

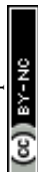


Table 3 MIC, MBC, and MFC (μM) of the tested hybrids and calculated rate (R) of (MBC or MFC) to MIC

Cpd. no.	Bacteria (Gram negative)												Bacteria (Gram positive)			Fungi		
	<i>E. coli</i>			<i>P. aeruginosa</i>			<i>K. pneumonia</i>			<i>A. baumannii</i>			MSSA			<i>C. albicans</i>		
	MIC	MBC	R	MIC	MBC	R	MIC	MBC	R	MIC	MBC	R	MIC	MBC	R	MIC	MFC	R
24	25	50	2	12.5	25	2	12.5	50	4	>200	>200	—	>200	>200	—	50	100	2
25	12.5	50	4	12.5	100	8	50	200	4	>200	>200	—	>200	>200	—	12.5	50	4
26	12.5	50	4	25	50	2	12.5	50	4	>200	>200	—	>200	>200	—	50	100	2
27	25	50	2	50	100	2	50	100	2	50	50	1	>200	>200	—	50	100	2
28	12.5	50	4	6.25	25	4	25	50	2	6.25	12.5	2	12.5	25	2	100	200	2
29	6.25	12.5	2	100	200	2	25	100	4	50	100	2	6.25	12.5	2	12.5	50	4
30	12.5	50	4	6.25	50	8	50	200	4	25	50	2	25	50	2	50	100	2
31	50	200	4	12.5	50	4	25	100	4	100	100	1	6.25	25	4	100	200	2
Levofloxacin	8	—	—	4	—	—	8	—	—	2	—	—	<0.5	—	—	NA	—	—
Amoxicillin	1	—	—	256	—	—	>1024	—	—	>1024	—	—	<0.5	—	—	NA	—	—
Fluconazole	—	—	—	—	—	—	—	—	—	—	—	—	—	—	—	64	—	—

Hybrids 29 and 31 displayed outstanding MIC value of 6.25 μM and MBC value of 12.5 μM and 25 μM , respectively, against MSSA. Hybrid 28, however, demonstrated MIC and MBC values of 12.5 and 25 μM , respectively. MIC/MBC ratio confirmed hybrids 28 and 29 as bactericidal and 31 as bacteriostatic against MSSA.

With the exception of 28 and 31, the assessed hybrids outperformed fluconazole in opposition to *C. albicans*, according to an evaluation of their fungistatic/fungicidal activities. With MBC values of 50 μM and MIC values of 12.5 μM , hybrids 25 and 29 indicated the strongest anti-fungal activity against *C. albicans*; nevertheless, the MFC to MIC Rate deemed that they were fungistatic.

2.2.3. Activity against biofilms. Since bacterial biofilm can cause a number of chronic illnesses, as was previously noted, effective therapies have become uncertain because of significant antibiotic resistance.^{66,67} Therefore, we evaluated the candidate hybrids for their antibiofilm activity at sub-inhibitory

concentrations to confirm that the antibiofilm activity was not due to bacteriostatic or bactericidal effect. Treatment of *P. aeruginosa* with each of the synthesized hybrids at 90% MIC resulted in 89 to 92% reduction in the production of biofilms in comparison to the untreated ones, outperforming Levofloxacin. Similar biofilm inhibition was noted upon administering 75% MIC of each assessed hybrid, excluding an outlier hybrid 31. Examining biofilm inhibition at 50% MIC of assessed hybrids, similarly, displayed the strong biofilm inhibition, with however two outlier compounds 28 and 31 (Fig. 2).

2.2.4. Time-kill kinetics. The time-killing results of 24–31 are plotted in Fig. 3. For all eight hybrids, the killing kinetics against *P. aeruginosa* was time dependent. Higher hybrid concentrations led to a more rapid decrease in CFU number. For hybrids 24, 26, 27 and 29, at 2-fold MIC, primary bacterial elimination was noticed after 3 hours, whereas bacterial elimination almost competed after 12 hours. Administering 4-fold and 8-fold MIC resulted in significant *P. aeruginosa* elimination starting at 6 hours. At 4-fold MIC of hybrids 28 and 31, *P. aeruginosa* were almost completely killed in 24 hours, whereas primary bacterial elimination was noted at 3 hours. Administering 8-fold MIC resulted in significant *P. aeruginosa* elimination starting at 12 hours. At 8-fold MIC of hybrids 25 and 30, all *P. aeruginosa* elimination was primarily observed after 3 hours and almost completed after 24 hours.

2.2.5. Inhibition of DNA gyrase. The potential of the novel quinoxaline hybrids to inhibit *P. aeruginosa* DNA-gyrase was assessed *in vitro* in comparison to a widely used antibiotic ciprofloxacin (Table 4). The produced compounds' IC_{50} values were expressed in-relation to ciprofloxacin, which was identified to possess a value of an IC_{50} of 143.1 μM (Fig. 4). The most effective DNA gyrase inhibitors were hybrids 25, 28, and 31, with IC_{50} values that were noticeably lower than those of ciprofloxacin (77.3, 87.6, and 65.5 μM). With values of 144.7 and 147.8 μM , respectively, hybrids 26 and 30 displayed DNA gyrase inhibition analogous to that of ciprofloxacin. With IC_{50} values surpassing 150 μM , the remaining hybrids showed minimal inhibition against DNA gyrase.

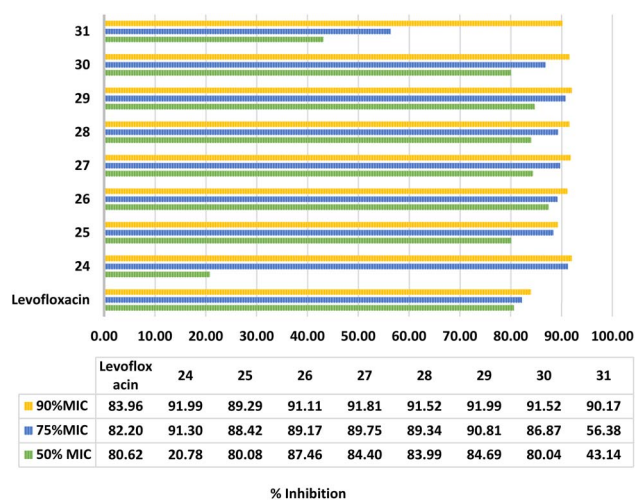


Fig. 2 Biofilm formation inhibition by evaluated hybrids versus Levofloxacin at 50%, 75% and 90% MIC.



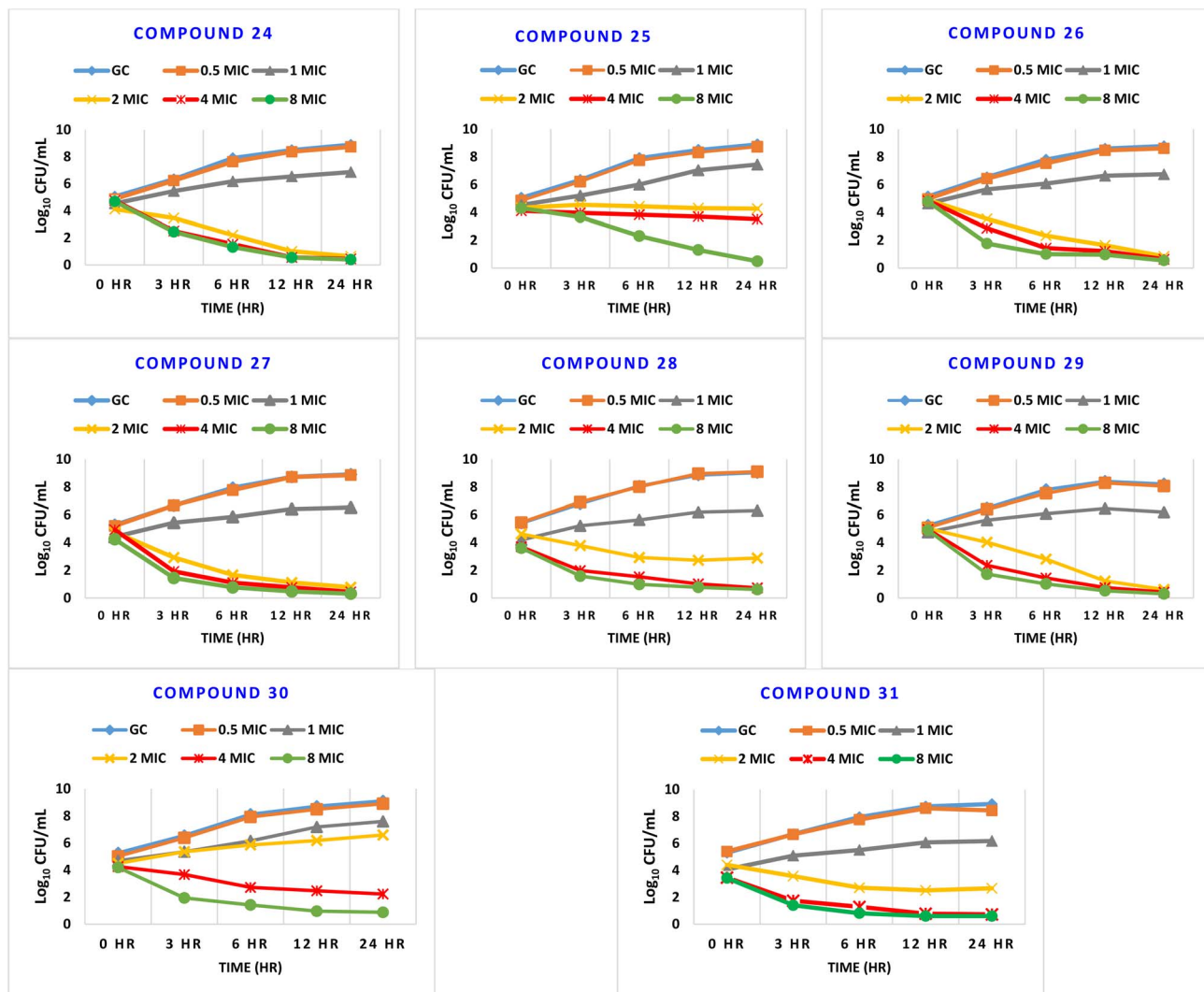


Fig. 3 Time-kill effect of 24–31 against *P. aeruginosa* versus growth control (GC).

2.3. The study of the structure–activity relationship (SAR)

This work exploited the novel compounds' biological processes to understand the connection between structure and the action (Fig. 5). The impact of the most potent chemicals on DNA Gyrase inhibitory verified that the 3-chloro-quinoxaliny bearing

hybrids are generally more active than 3-hydroxy-quinoxaliny derivatives. Substitution on the ring of the *N*-phenyl demonstrated a pattern in the inhibitory effect of DNA gyrase, where

Table 4 DNA gyrase inhibition activity of synthesized hybrids versus ciprofloxacin

Cpd. no.	DNA gyrase supercoiling IC ₅₀ (μM)
24	172.7 ± 15.1
25	77.3 ± 15.5
26	144.7 ± 15.2
27	200.8 ± 16.4
28	87.6 ± 16.0
29	182.2 ± 16.5
30	147.8 ± 15.6
31	65.5 ± 16.8
Ciprofloxacin	143.1 ± 11.2

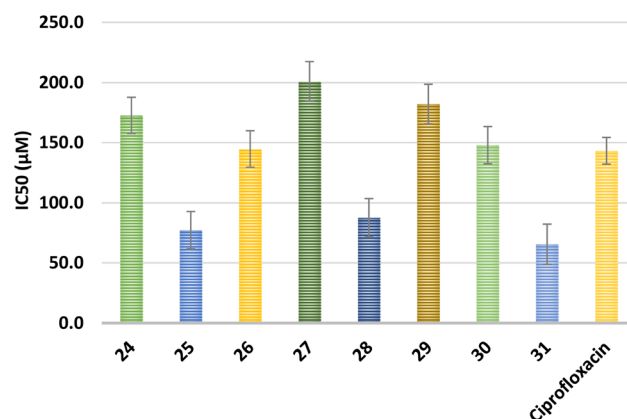


Fig. 4 The IC₅₀ values of artificial hybrids on *P. aeruginosa* DNA gyrase.



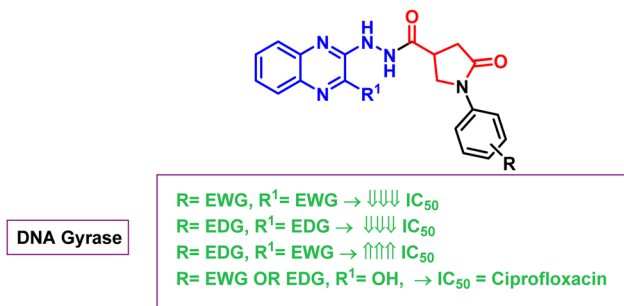


Fig. 5 SAR of pyrrolidine–quinoxaline hybrids.

electron-withdrawing groups on the *para* position demonstrated superior activity to Ciprofloxacin as observed with **28** and **31**. The remaining 3-chloro-quinoxaliny hybrids bearing electron-donating groups on the *N*-phenyl ring (**30**) and (**29**) demonstrated DNA gyrase inhibitory activity either equipotent or lower than Ciprofloxacin, respectively. This finding confirms the importance of bearing electron-withdrawing groups at both R and R¹. Most importantly, Presence of 3-hydroxy substitution on the *N*-phenyl ring generally resulted in lower DNA gyrase inhibitory activity, however equipotent to Ciprofloxacin. Such a trend was not noted in hybrid **26**, bearing two hydroxyl groups. A plausible explanation for this exception suggests the formation of an extra compensatory hydrogen bond with DNA gyrase. It was noted, however that hybrids bearing both R groups as electron-donating still exhibited equipotent or superior DNA gyrase inhibitory activity to Ciprofloxacin (hybrids **26** and **25**, respectively).

2.4. *In silico* ADME

To assess the physicochemical characteristics of the produced hybrids in comparison to levofloxacin, amoxicillin, and fluconazole, a simulated study was conducted.^{68–70} In terms of their physicochemical characteristics, all compounds and reference drugs—aside from **27**—display zero breaches of the Lipinski's requirements regarding oral medications, as stated by Table 5. With the exception of **27** and amoxicillin, all examined derivatives and references adhere to Veber's requirements.

Table 6 TPSA and %ABS of the synthesized differences

Cpd. no.	TPSA	% ABS
24	131.240	63.720
25	107.450	71.920
26	127.680	64.950
27	153.270	56.120
28	111.010	70.700
29	87.220	78.900
30	107.450	71.920
31	133.040	63.100
Levofloxacin	84.240	79.180
Amoxicillin	158.260	52.980
Fluconazole	81.650	80.090
Ciprofloxacin	74.570	82.600

Additionally, all hybrids exhibit high flexibility with no more than six rotatable bonds. This implies that oral consumption of the eight hybrids is possible. Each analogue was given a score between 3.05 and 3.76 for ease of manufacturing, suggesting that a large number of them can be produced on a large scale.

Additionally, the percentage was computed using TPSA as follows: % ABS = 109 – (0.354 × TPSA).^{71,72} All of the prospective antibacterial agents possess estimated absorption percentages between 87 and 154%, which indicates strong oral bioavailability analogous to reference medications. The percentage of ABS ranges from 56 to 79% (Table 6).

The findings of monitoring the new antibacterial hybrids for pharmacokinetic and medicinal chemistry characteristics suggest that, with the exception of **27** and amoxicillin, every other compound has demonstrated adequate GI tract absorption. They can't overcome all, however, overcome the brain-blood barrier. Every hybrid has a high probability of effluxing outside the cell and is a P-gp (P-glycoprotein) substrate.

Furthermore, inhibition of metabolic enzymes by investigated hybrids was computationally assessed. Predictions revealed that hybrid **26** did not inhibit any of the assessed metabolic enzymes, whereas **27** only inhibited CYP2C9.

All investigated hybrids received a bioavailability score of 0.55, with the exception of derivative **24**, which received a score of 0.56, analogous to the other reference drugs. We assessed every hit, and PAINS showed no warnings. In order to avoid

Table 5 Physicochemical characteristics of the most physiologically active drugs compared to reference medications

Cpd. no.	HBD	HBA	MW	<i>M log P</i>	No. of rotatable bonds	Lipinski's violations	Veber's violations	Synthetic accessibility
24	3	6	388.38	1.09	5	0	0	3.64
25	3	5	391.42	2.16	5	0	0	3.76
26	4	6	379.37	1.2	5	0	0	3.67
27	3	7	408.37	0.88	6	1	1	3.71
28	2	5	406.83	1.68	5	0	0	3.05
29	2	4	409.87	2.76	5	0	0	3.31
30	3	5	397.82	1.8	5	0	0	3.30
31	2	6	426.81	1.47	6	0	0	3.49
Levofloxacin	2	7	379.38	0.2	2	0	0	3.74
Amoxicillin	4	6	365.40	0.23	5	0	1	4.17
Fluconazole	1	7	306.27	1.47	5	0	0	2.45
Ciprofloxacin	2	5	331.34	1.28	3	0	0	2.51



Table 7 Medicinal chemistry parameters and pharmacokinetic features

Cpd. no.	GI absorption	BBB permeation	Pgp substrate	CYP1A2 inhibitor	CYP2C19 inhibitor	CYP2C9 inhibitor	CYP2D6 inhibitor	CYP3A4 inhibitor	Bioavailability score	PAINS alerts
24	High	No	Yes	Yes	No	Yes	No	No	0.56	0
25	High	No	Yes	Yes	No	Yes	Yes	Yes	0.55	0
26	High	No	Yes	No	No	No	No	No	0.55	0
27	Low	No	Yes	No	No	Yes	No	No	0.55	0
28	High	No	Yes	Yes	No	Yes	Yes	Yes	0.55	0
29	High	No	Yes	Yes	Yes	Yes	Yes	Yes	0.55	0
30	High	No	Yes	Yes	No	Yes	Yes	Yes	0.55	0
31	High	No	Yes	No	Yes	Yes	No	Yes	0.55	0
Levofloxacin	High	No	Yes	No	No	No	No	No	0.55	0
Amoxicillin	Low	No	No	No	No	No	No	No	0.55	0
Fluconazole	High	No	Yes	No	Yes	No	No	No	0.55	0
Ciprofloxacin	High	No	Yes	No	No	No	No	No	0.55	0

inaccurate outcomes, PAINS are important considerations while developing drugs (Table 7). The evaluated substances' bioavailability radar is illustrated within Fig. 6. The intended hybrids revealed promise as attractive agents for antibacterial treatment by exhibiting typically very good physicochemical properties.

2.5. Toxicity assessment *in silico*

The pkCSM online tools confirm the security of the most active hits since, as shown in Table 8, with the exception of **31**, none have demonstrated the potential for mutagenicity according to AMES toxicity, equivalent to that of Levofloxacin and Fluconazole. Furthermore, compared to Fluconazole (0.114 log mg per kg per day) and other hits, hybrid **26** demonstrated a maximum acceptable human dose of 0.261 log mg per kg per day. The LD₅₀ value of compound **31** has been estimated to be 2.587 mol kg⁻¹, which is considerably higher than that of Fluconazole and equivalent to that of Levofloxacin. Additionally, compound **26** had an LOAEL value of 2.798 log mg per kg bw per day, surpassing the values of all hits and the reference drugs. The potassium channels encoded by the hERG gene control the repolarization of cardiac action potentials. Pharmaceutical companies are concerned because blocking. These channels may cause cardiovascular problems that could be fatal. While every potent ingredient was expected to have an inhibitory impact on hERG-II, it was shown that they had no effect on hERG-I.

The pkCSM server was used to examine hepatotoxicity. All active hits, including reference medications, were predicted to be hepatotoxic. Skin discomfort was not evident in any of the hits. It was expected that Levofloxacin, Fluconazole, and active hybrids would be categorized as a class IV (GHS) utilizing the-ProTox-II identification webtool (Table 8). It was discovered that all active hybrids had LD₅₀ values exceeding the reference medications. None of the active hits were supposed to be immunotoxic, cytotoxic, or mutagenic. The active hybrids and reference drugs were expected to exhibit both neurotoxicity and clinical toxicity. Neither the voltage-gated sodium channel (VGSC) nor the phosphoprotein (tumor suppressor) p53 were suppressed by those active hybrids.

2.6. Molecular docking

The investigated hybrids were docked against DNA gyrase (PDB ID: 8BN6) to predict binding mode and interactions. The bound ligand was separately redocked to validate the docking procedure. Validation was confirmed by visualizing the overlay of the original and the redocked pose (Fig. 7) and calculated RMSD value of 0.79 Å. The investigated hybrids were docked implementing the same parameters. The 2D interactions and energy scores are recorded (Table 9).

The interaction of co-crystallized ligand with DNA Gyrase involved six alkyl-type van der Waals interactions with Val73, Val169, Val45 and Val122. Two H-bonds were noted between the co-crystallized ligand and key amino acids: Arg138 and Thr167. Three pi-alkyl interactions were observed with Pro81 and Ile96. Glu52 interacted with the ligand through two pi-anion interactions, whereas Arg78 interacted through one pi-cation interactions. Amide-pi stacked interaction was involved with Asn48.

Docking of ciprofloxacin as a reference against DNA Gyrase demonstrated that the binding energy was -9.70 kcal per mol. Ciprofloxacin was able to interact with key amino acids; including three H-bonds with Thr167, Arg138 and Arg78, two alkyl-type van der Waals interactions with Ile96 and Ile80, one pi-alkyl interaction with Ile80, one amide-pi stacked interaction with Asn48, and two pi-anion interactions with Glu52.

Simulation of the hybrid **25** binding showed that the binding energy was -8.12 kcal mol⁻¹. Key amino acids were involved in the interaction of **25** with DNA Gyrase. Such interactions included two amide-Pi stacked interactions with Asn48, two pi-alkyl-type van der Waals interactions with Ile96 and Val122, one H-bond with Thr167 and one alkyl-type van der Waals with Ile96. Additional interactions included two pi-alkyl interactions with Ile80, two H-bonds with Asp75 and one amide-pi stacked interaction with Gly79.

Binding simulation of hybrid **28** demonstrated a binding energy of -8.05 kcal mol⁻¹. Key amino acids were involved in the interaction of **28** with DNA Gyrase. Such interactions included two pi-alkyl-type van der Waals interactions with Ile96, one H-bond with Val122. Additional interactions involved two pi-alkyl interactions with Ile 80, and one amide-pi stacked interaction with Gly79.



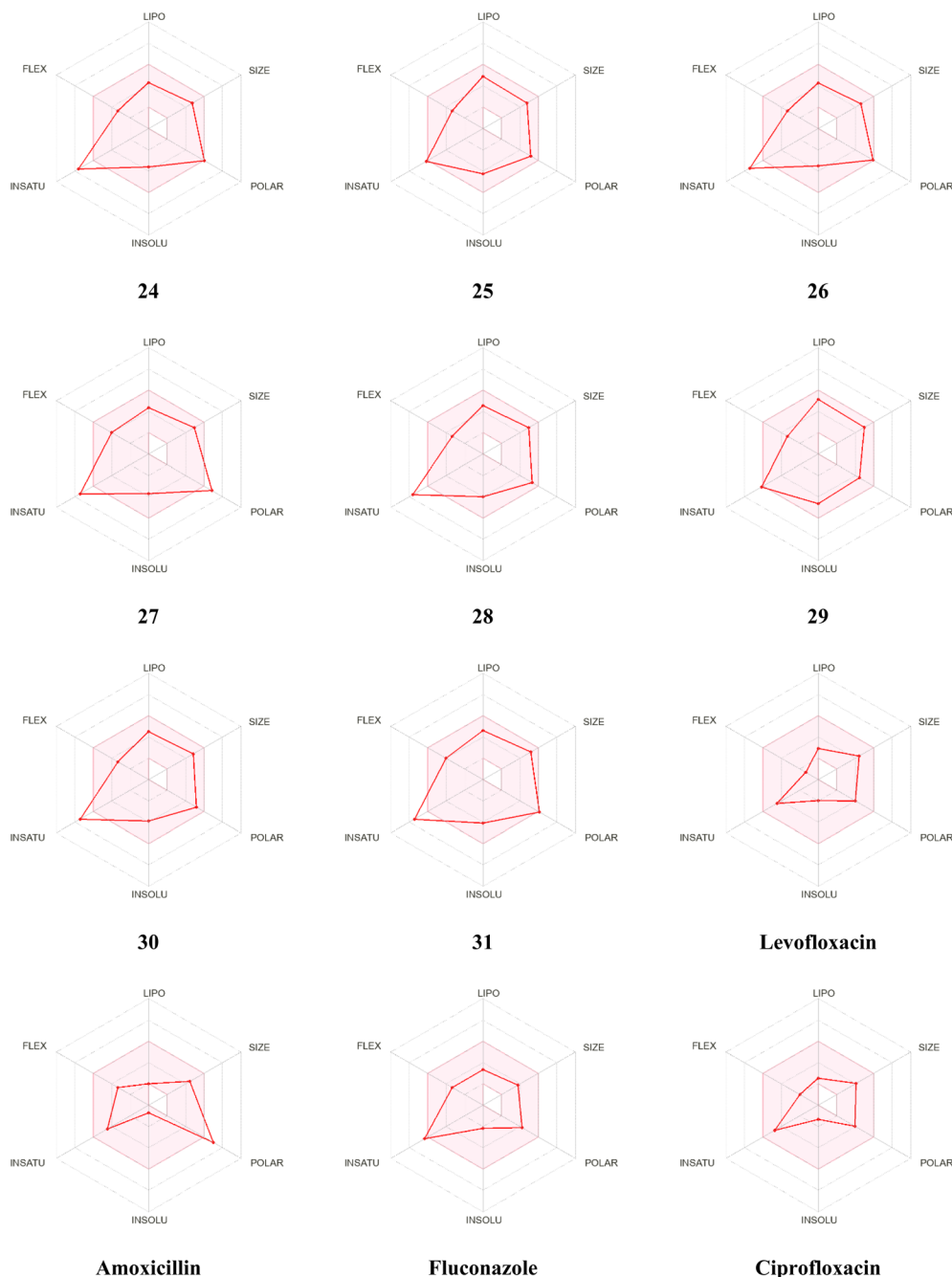


Fig. 6 The bioavailability radar of the assessed hybrids and the drugs that are referenced are correlated. The area that is colored provides ideal chemical and physical conditions for absorption through the mouth. Insolubility (INSOLU), unsaturation (INSATU), polarity (POLAR), molecular weight (SIZE), lipophilicity (LIPO), and flexibility (FLEX).

Binding simulation of hybrid **31** revealed that the binding energy score was $-10.80 \text{ kcal mol}^{-1}$. Although docking scores do not strictly follow experimental IC_{50} values, the binding energy of **31** still confirms superiority to Ciprofloxacin *in silico*. Key amino acids were involved in the interaction of **31** with DNA Gyrase. Such interactions included three pi-alkyl-type van der Waals interactions with Val169, Val122 and Ile96, one H-bond with Glu52, and two amide-pi stacked interactions with Asn48. Additional interactions involved two pi-alkyl interactions with Ile80.

The binding mode and energies of the remaining hybrids are noted in (Table 9, Fig. 8 and 9).

3. Experimental

3.1. Chemistry

3.1.1. Synthesis of 16–23

3.1.1.1 Method A. In a flask with a round bottom, 1 mmol (0.13 g) of itaconic acid (**1**) was mixed with 1 mmol of each of the respective amines **8–15** with water. The mixture of the reaction



Table 8 Toxicity of the most active hybrids 25, 26, 30, and 31 versus with Levofloxacin and Fluconazole

Test	Most active compounds					
	25	26	30	31	Levofloxacin	Fluconazole
pkCSM⁷³						
AMES toxicity	No	No	No	Yes	No	No
Max. tolerated dose (human)	-0.11	0.261	0.144	0.021	0.965	0.114
hERG I inhibitor	No	No	No	No	No	No
hERG II inhibitor	Yes	Yes	Yes	Yes	No	No
Oral rat acute toxicity (LD ₅₀)	2.046	2.089	2.116	2.587	2.59	2.328
Oral rat chronic toxicity (LOAEL)	1.504	2.798	2.046	2.07	1.791	1.033
Hepatotoxicity	Yes	Yes	Yes	Yes	Yes	Yes
Skin sensitization	No	No	No	No	No	No
<i>T. Pyriformis</i> toxicity	0.405	0.346	0.357	0.326	0.285	0.312
Minnow toxicity	0.812	1.777	0.187	-0.601	1.273	3.872
ProTox-II prediction⁷⁴						
LD ₅₀ (mg kg ⁻¹)	2000	1600	1600	2000	1478	1271
Toxicity class	IV	IV	IV	IV	IV	IV
Neurotoxicity	+(0.72)	+(0.73)	+(0.9)	+(0.57)	+(0.93)	+(0.91)
Nephrotoxicity	-(0.53)	-(0.53)	+(0.58)	-(0.54)	+(0.85)	-(0.61)
Immunotoxicity	-(0.99)	-(0.98)	-(0.96)	-(0.98)	-(0.99)	-(0.83)
Mutagenicity	-(0.57)	-(0.58)	-(0.54)	+(0.79)	+0.66	-(0.52)
Cytotoxicity	-(0.57)	-(0.6)	-(0.6)	-(0.53)	-(0.69)	-(0.76)
Clinical toxicity	+(0.66)	+(0.68)	+(0.73)	+(0.58)	+(0.62)	+(0.55)
Phosphoprotein (tumor suppressor)	-(0.88)	-(0.88)	-(0.87)	-(0.86)	-(0.98)	-(0.97)
p53						
Voltage gated sodium channel (VGSC)	-(0.89)	-(0.76)	-(0.7)	-(0.82)	-(0.95)	-(0.95)

underwent reflux heating for 24 hours. The resulting precipitates were filtered recrystallized from ethanol.⁷⁵⁻⁷⁸

3.1.1.2 Method B. In a beaker of 250 mL, 1 mmol (0.13 g) itaconic acid (**1**) was dissolved in the least amount of ethanol and mixed with 1 mmol of each of the respective amines **8-15**. The reaction solution was sonicated until reaction completion, monitored by TLC.

3.1.1.3 Method C. In a 250 mL beaker, 1 mmol (0.13 g) itaconic acid (**1**) was dissolved in the least amount of ethanol and mixed with 1 mmol of each of the respective amines **8-15**. The reaction solution was microwaved in domestic microwave until reaction completion, monitored by TLC.

3.1.1.4 1-(3-Chlorophenyl)-5-oxopyrrolidine-3-carboxylic acid, 8. Yellowish powder; yield: 91%, m.p. 187-189 °C; ¹H NMR (400 MHz, CDCl₃): δ = 12.38 (s, 1H, COOH), 7.28-7.24 (m, 3H, A-H), 7.20-7.15 (m, 1H, A-H), 4.19 (dd, *J* = 11.9, 4.4 Hz, 1H, pyrrolidine CH₂), 3.93 (dd, *J* = 11.9, 4.4 Hz, 1H, pyrrolidine CH₂), 3.33-3.24 (m, 1H, pyrrolidine CH), 2.78 (dd, *J* = 15.0, 5.6 Hz, 1H, pyrrolidine CH₂), 2.53 (dd, *J* = 15.0, 5.6 Hz, 1H, pyrrolidine CH₂).

3.1.1.5 1-(4-Chlorophenyl)-5-oxopyrrolidine-3-carboxylic acid, 9. Yellowish powder; yield: 95%, m.p. 184-185 °C; ¹H NMR (400 MHz, CDCl₃): δ = 12.14 (s, 1H, COOH), 7.31-7.22 (m, 2H, A-H), 7.08-7.00 (m, 2H, A-H), 4.15 (dd, *J* = 11.8, 4.5 Hz, 1H, pyrrolidine CH₂), 3.90 (dd, *J* = 11.8, 4.2 Hz, 1H, pyrrolidine CH₂), 3.37-3.27 (m, 1H, pyrrolidine CH), 2.76 (dd, *J* = 14.9, 5.5 Hz, 1H, pyrrolidine CH₂), 2.51 (dd, *J* = 14.9, 5.5 Hz, 1H, pyrrolidine CH₂).

3.1.1.6 1-(4-Cyanophenyl)-5-oxopyrrolidine-3-carboxylic acid, 10. Orange powder; yield: 90%, m.p. 181-183 °C; ¹H NMR (400 MHz, CDCl₃): δ = 12.38 (s, 1H, COOH), 7.58 (d, *J* = 7.5 Hz, 2H, A-

H), 7.30 (d, *J* = 7.1 Hz, 2H, A-H), 4.17 (dd, *J* = 11.7, 4.2 Hz, 1H, pyrrolidine CH₂), 3.92 (dd, *J* = 12.0, 4.4 Hz, 1H, pyrrolidine CH₂), 3.29 (p, *J* = 4.9 Hz, 1H, pyrrolidine CH), 2.78 (dd, *J* = 14.8, 5.6 Hz, 1H, pyrrolidine CH₂), 2.53 (dd, *J* = 14.8, 5.6 Hz, 1H, pyrrolidine CH₂).

3.1.1.7 1-(2,5-Dimethylphenyl)-5-oxopyrrolidine-3-carboxylic acid, 11. Light yellow powder; yield: 92%, m.p. 196-198 °C; ¹H NMR (400 MHz, CDCl₃): δ = 12.13 (s, 1H, COOH), 6.97 (d, *J* = 2.8 Hz, 1H, A-H), 6.87 (d, *J* = 8.5 Hz, 1H, A-H), 6.72 (dd, *J* = 8.6, 2.9 Hz, 1H, A-H), 4.32 (dd, *J* = 11.6, 4.3 Hz, 1H, pyrrolidine CH₂), 4.07 (dd, *J* = 11.6, 4.3 Hz, 1H, pyrrolidine CH₂), 3.04-2.93 (m, 1H, pyrrolidine CH), 2.70 (dd, *J* = 15.0, 5.6 Hz, 1H, pyrrolidine CH₂), 2.44 (dd, *J* = 14.9, 5.6 Hz, 1H, pyrrolidine CH₂), 2.17 (s, 3H, CH₃), 2.09 (s, 3H, CH₃).

3.1.1.8 1-(3-Hydroxyphenyl)-5-oxopyrrolidine-3-carboxylic acid, 12. Yellowish powder; yield: 89%, m.p. 179-181 °C; ¹H NMR (400 MHz, CDCl₃): δ = 12.38 (s, 1H, COOH), 7.98 (s, 1H, OH), 7.11 (t, *J* = 7.9 Hz, 1H, A-H), 7.02 (d, *J* = 7.1 Hz, 1H, A-H), 6.65 (d, *J* = 8.3 Hz, 1H, A-H), 6.59 (d, *J* = 2.6 Hz, 1H, A-H), 4.19 (dd, *J* = 11.9, 4.3 Hz, 1H, pyrrolidine CH₂), 3.93 (dd, *J* = 11.9, 4.4 Hz, 1H, pyrrolidine CH₂), 3.29 (p, *J* = 4.8 Hz, 1H, pyrrolidine CH), 2.78 (dd, *J* = 15.0, 5.4 Hz, 1H, pyrrolidine CH₂), 2.53 (dd, *J* = 15.1, 5.6 Hz, 1H, pyrrolidine CH₂).

3.1.1.9 1-(4-Nitrophenyl)-5-oxopyrrolidine-3-carboxylic acid, 13. Orange powder; yield: 88%, m.p. 198-200 °C; ¹H NMR (400 MHz, CDCl₃): δ = 13.14 (s, 1H, COOH), 8.05 (d, *J* = 7.2 Hz, 2H, A-H), 7.43 (d, *J* = 7.4 Hz, 2H, A-H), 3.98 (dd, *J* = 11.9, 4.4 Hz, 1H, pyrrolidine CH₂), 3.73 (dd, *J* = 11.8, 4.3 Hz, 1H, pyrrolidine CH₂), 3.38 (p, *J* = 4.8 Hz, 1H, pyrrolidine CH), 2.79 (dd, *J* = 14.9,



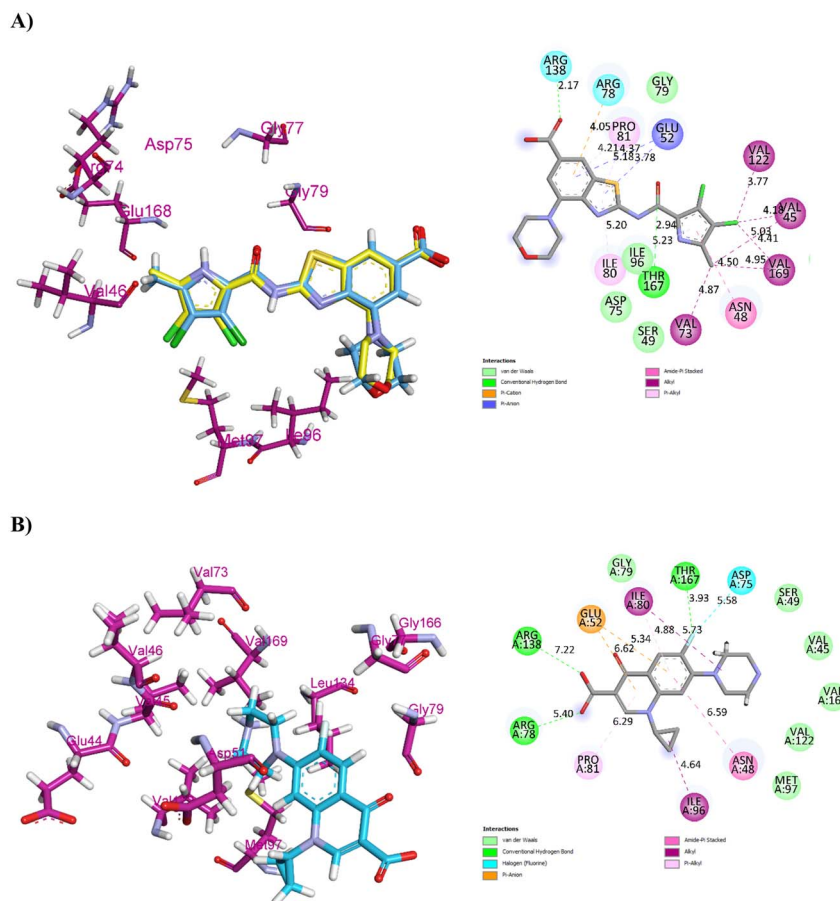


Fig. 7 (A) Overlay of bound ligand (cyan) and redocked ligand (yellow) and bound ligand interactions, (B) binding mode and ligand interactions of Ciprofloxacin.

5.6 Hz, 1H, pyrrolidine CH₂), 2.54 (dd, $J = 14.8$, 5.6 Hz, 1H, pyrrolidine CH₂).

3.1.1.10 1-(Naphthalen-1-yl)-5-oxopyrrolidine-3-carboxylic acid, 14. Light yellow powder; yield: 92%, m.p. 195–196 °C; ¹H NMR (400 MHz, CDCl₃): $\delta = 12.13$ (s, 1H, COOH), 7.78–7.64 (m, 3H, A–H), 7.42–7.22 (m, 3H, A–H), 7.14 (d, $J = 7.8$ Hz, 1H, A–H), 4.32 (dd, $J = 11.5$, 4.3 Hz, 1H, pyrrolidine CH₂), 4.06 (dd, $J = 11.6$, 4.4 Hz, 1H, pyrrolidine CH₂), 2.99 (p, $J = 4.6$ Hz, 1H, pyrrolidine CH), 2.70 (dd, $J = 15.0$, 5.7 Hz, 1H, pyrrolidine CH₂), 2.45 (dd, $J = 15.0$, 5.6 Hz, 1H, pyrrolidine CH₂).

3.1.1.11 5-Oxo-1-phenylpyrrolidine-3-carboxylic acid, 15. Yellow powder; yield: 90%, m. p. 186–187 °C; ¹H NMR (400 MHz, CDCl₃): $\delta = 12.38$ (s, 1H, COOH), 7.36–7.15 (m, 5H, A–H), 4.17 (dd, $J = 11.9$, 4.2 Hz, 1H, pyrrolidine CH₂), 3.92 (dd, $J = 11.9$, 4.4 Hz, 1H, pyrrolidine CH₂), 3.33–3.24 (m, 1H, pyrrolidine CH), 2.78 (dd, $J = 15.0$, 5.6 Hz, 1H, pyrrolidine CH₂), 2.53 (dd, $J = 15.0$, 5.6 Hz, 1H, pyrrolidine CH₂).

3.1.2. Synthesis of hybrids 24–31. In a 250 mL beaker, 1 mmol of 2-hydroxy-3-hydrazinylquinoxaline (4) (0.17 g) or 2-chloro-3-hydrazinylquinoxaline (6) (0.19 g) was dissolved in the least amount of ethanol with 1 mmol of each of fragments 18–21. The reaction solution was sonicated until reaction completion monitored by TLC.^{79–83}

3.1.2.1 1-(4-Cyanophenyl)-N'-(3-hydroxyquinoxalin-2-yl)-5-oxopyrrolidine-3-carbohydrazide, 24. Orange powder; yield: (59%), m.p. 264–265 °C; IR (KBr, cm⁻¹) = 3474–3359 (2NH & OH), 3073 (CH aromatic), 2214 (CN), 1687 (br. 2C=O); ¹H NMR (400 MHz, CDCl₃): $\delta = 12.87$ (s, 1H, OH), 11.04 (d, $J = 6.6$ Hz, 1H, –NH–NH–CO), 8.18 (d, $J = 6.8$ Hz, 1H, NH–NH–CO), 8.05 (dd, $J = 7.7$, 2.0 Hz, 1H, A–H), 7.73–7.52 (m, 5H, A–H), 7.30 (d, $J = 7.8$ Hz, 2H, A–H), 4.17 (dd, $J = 12.4$, 3.6 Hz, 1H, pyrrolidine CH₂), 3.92 (dd, $J = 12.5$, 3.5 Hz, 1H, pyrrolidine CH₂), 3.12 (m, 1H, pyrrolidine CH), 2.52 (dd, $J = 15.7$, 4.9 Hz, 1H, pyrrolidine CH₂), 2.33 (dd, $J = 15.6$, 4.9 Hz, 1H, pyrrolidine CH₂); ¹³C NMR (100 MHz, CDCl₃): $\delta = 172.77$ (C=O), 171.24 (C=O), 146.52, 141.54, 140.22, 136.97, 136.51, 132.41, 128.93, 126.45, 125.38, 123.42, 118.23 (CN), 117.28, 102.21, 48.51 (pyrrolidine CH₂), 38.02 (pyrrolidine CH), 36.42 (pyrrolidine CH₂); MS(EI, 70 eV): m/z (%) = 388 [M⁺] (18%), 356.46 (100%); anal. calc. for: C₂₀H₁₆N₆O₃ (388.38): C, 61.85; H, 4.15; N, 21.64; found: C, 61.80; H, 4.17; N, 21.68.

3.1.2.2 1-(2,5-Dimethylphenyl)-N'-(3-hydroxyquinoxalin-2-yl)-5-oxopyrrolidine-3-carbohydrazide, 25. Light brown powder; yield: (77%), m.p. 271–272 °C; IR (KBr, cm⁻¹) = 3369–3280 (br. 2NH & OH), 3060 (CH aromatic), 2971, 2860 (CH aliphatic), 1698 (br. 2C=O); ¹H NMR (400 MHz, CDCl₃): $\delta = 12.86$ (s, 1H, OH), 10.79 (d, $J = 6.8$ Hz, 1H, –NH–NH–CO), 8.88 (d, $J = 6.6$ Hz, 1H,



Table 9 Binding energies and interactions of bound ligand and hybrids against DNA Gyrase (PDB: 8BN6)

Cpd. no.	S (kcal mol ⁻¹)	Type of interaction	Amino acid	Length (Å)	Cpd. no.	S (kcal mol ⁻¹)	Type of interaction	Amino acid	Length (Å)
24	-8.35	Amide-Pi stacked	Asn48	5.01	28	-8.05	Pi-alkyl	Ile80	4.44
		Amide-Pi stacked	Asn48	4.66			Pi-alkyl	Ile80	4.51
		Pi-alkyl	Ile80	4.41			Amide-Pi stacked	Gly79	4.50
		Pi-alkyl	Ile80	4.40			H-bond	Val122	4.85
		H-bond	Thr167	2.17			Pi-alkyl	Ile96	5.55
		H-bond	Asp75	1.97			Pi-alkyl	Ile96	7.25
		Amide-Pi stacked	Gly79	5.10					
		Pi-alkyl	Ile96	4.78					
		Pi-alkyl	Val122	5.35					
		H-bond	Arg138	2.96					
25	-8.12	Amide-Pi stacked	Asn48	4.64	29	-7.66	Alkyl	Ile96	5.35
		Amide-Pi stacked	Asn48	5.00			Alkyl	Pro81	4.41
		Pi-alkyl	Ile80	4.41			H-bond	Arg138	2.82
		Pi-alkyl	Ile80	4.44			Pi-alkyl	Ile80	4.25
		H-bond	Asp75	1.95			Pi-alkyl	Ile80	4.23
		H-bond	Asp75	3.12			Amide-Pi stacked	Gly79	4.80
		Amide-Pi stacked	Gly79	5.12					
		Pi-alkyl	Ile96	4.46					
		Alkyl	Ile96	4.52					
		H-bond	Thr167	2.16					
26	-7.81	Pi-alkyl	Val122	5.32	30	-10.20	H-bond	Asp51	1.96
		H-bond	Arg138	2.23			Amide-Pi stacked	Asn48	5.07
		H-bond	Asp75	2.79			Amide-Pi stacked	Asn48	4.69
		H-bond	Gly79	2.27			Pi-alkyl	Val122	5.25
		Amide-Pi-stacked	Gly79	4.36			Pi-alkyl	Val169	5.43
		Pi-alkyl	Ile80	4.10			Pi-alkyl	Ile80	5.32
		Pi-alkyl	Ile80	4.37			Pi-alkyl	Ile80	4.29
		H-bond	Glu52	2.85			Pi-alkyl	Ile80	4.29
		H-bond	ILE96	4.48			H-bond	GLU52	3.07
		Pi-alkyl	VAL122	5.34			Pi-alkyl	ILE80	4.34
27	-7.41	H-bond	ARG138	3.06	31	-10.80	Pi-alkyl	ILE80	5.42
		Amide Pi-stacked	ASN48	4.99			Pi-alkyl	VAL169	5.35
		Amide Pi-stacked	ASN48	4.65			Amide	ASN48	5.08
		Pi-alkyl	ILE80	4.43			Pi-stacked	ASN48	4.65
		Pi-alkyl	ILE80	4.41			Amide	ASN48	4.65
		H-bond	ASP75	3.12			Pi-stacked	ASN48	4.65
		H-bond	ASP75	1.95			Pi-alkyl	VAL122	5.22
		Amide Pi-stacked	GLY79	5.11			Pi-alkyl	ILE96	4.66
		H-bond	THR167	2.16					
		H-bond	ARG138	2.17					
Co-crystalized	-9.34	Pi-cation	ARG78	4.05	Ciprofloxacin	-9.70	H-bond	ARG138	7.22
		Pi-alkyl	PRO81	4.21			Pi-anion	GLU52	6.62
		Pi-alkyl	PRO81	4.37			Pi-anion	GLU52	5.34
		Pi-anion	GLU52	5.18			Pi-alkyl	ILE80	4.88
		Pi-anion	GLU52	3.78			Alkyl	ILE80	5.73
		Pi-alkyl	ILE96	5.23			H-bond	THR167	3.93
		H-bond	THR167	2.94			H-bond	ASP75	5.58
		H-bond	THR167	2.94			Amide-pi stacked	ASN48	6.59
		Alkyl	VAL73	4.87			Alkyl	ILE96	4.64
		Amide-Pi stacked	ASN48	4.50			H-bond	ARG78	5.40
		Alkyl	VAL169	4.95					
		Alkyl	VAL169	4.41					
		Alkyl	VAL45	5.03					
		Alkyl	VAL45	4.18					
		Alkyl	VAL122	3.77					



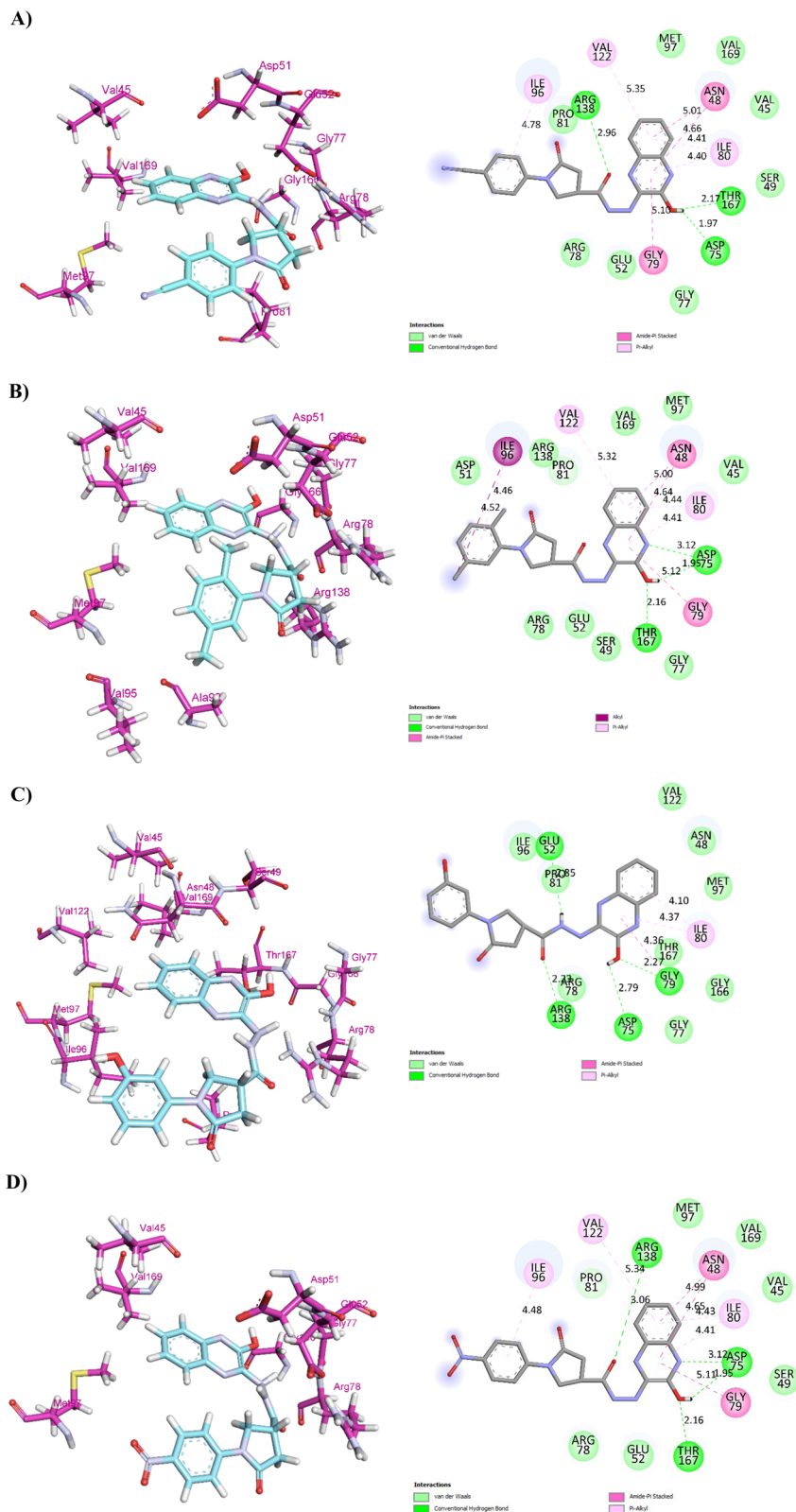


Fig. 8 Binding mode and ligand interactions of (A) 24 (B) 25 (C) 26 (D) 27.

NH-NH-CO), 7.91 (dd, $J = 8.0, 1.6$ Hz, 1H, A-H), 7.68 (dd, $J = 8.0, 1.8$ Hz, 1H, A-H), 7.65–7.52 (m, 2H, A-H), 7.12 (d, $J = 8.2$ Hz, 1H, A-H), 6.98 (d, $J = 8.2$ Hz, 2H, A-H), 4.21 (dd, $J = 12.2, 3.6$ Hz,

1H, pyrrolidine CH₂), 3.96 (dd, $J = 12.3, 3.7$ Hz, 1H, pyrrolidine CH₂), 3.34–3.25 (m, 1H, pyrrolidine CH), 2.75 (dd, $J = 15.7, 4.9$ Hz, 1H, pyrrolidine CH₂), 2.53 (dd, $J = 15.7, 4.9$ Hz, 1H,



3.1.2.3 *N'*-(3-Hydroxyphenyl)-*N'*-(3-hydroxyquinoxalin-2-yl)-5-oxopyrrolidine-3-carbohydrazide, **26**. Dark yellow powder; yield: (62%), m.p. 243–244 °C; IR (KBr, cm^{-1}) = 3422–3269 (br. 2NH & 2OH), 3070 (CH aromatic), 1656–1703 (br. 2C=O); ^1H NMR (400 MHz, CDCl_3): δ = 11.03 (s, 1H, OH), 8.80 (d, J = 6.6 Hz, 1H, –NH–NH–CO), 8.60 (s, 1H, OH), 8.17 (d, J = 8.1 Hz, 1H, A–H), 7.73 (dd, J = 8.1, 1.6 Hz, 1H, 1H, A–H), 7.70–7.54 (m, 2H, 1H, A–H), 7.09 (t, J = 7.9 Hz, 1H, 1H, A–H), 6.83 (d, J = 6.9 Hz, 1H, A–H), 6.68 (d, J = 8.6 Hz, 1H, A–H), 6.54 (t, J = 2.3 Hz, 1H), 5.90 (d, J = 6.8 Hz, 1H, –NH–NH–CO), 3.90 (dd, J = 12.5, 3.5 Hz, 1H, pyrrolidine CH_2), 3.65 (dd, J = 12.5, 3.7 Hz, 1H, pyrrolidine CH_2), 3.08–3.00 (m, 1H, pyrrolidine CH), 2.54 (dd, J = 15.6, 4.9 Hz, 1H, pyrrolidine CH_2), 2.40 (dd, J = 15.8, 4.8 Hz, 1H, pyrrolidine CH_2); ^{13}C NMR (100 MHz, CDCl_3): δ = 173.48 (C=O), 172.48 (C=O), 158.81, 147.23, 147.07, 140.85, 136.93, 135.59, 131.70, 131.16, 129.32, 125.08, 122.44, 117.35, 109.00, 106.54, 49.02 (pyrrolidine CH_2), 40.83 (pyrrolidine CH), 38.00 (pyrrolidine CH_2); MS (EI, 70 eV): m/z (%) = 379.40 [M^+] (19.76%), 190.16 (100%); anal. calc. for: $\text{C}_{19}\text{H}_{17}\text{N}_5\text{O}_4$ (379.37): C, 60.15; H, 4.52; N, 18.46; found: C, 60.19; H, 4.50; N, 18.48.

3.1.2.4 *N'*-(3-Hydroxyquinoxalin-2-yl)-1-(4-nitrophenyl)-5-oxopyrrolidine-3-carbohydrazide, **27**. Light brown powder; yield: (73%), m.p. 267–268 °C; IR (KBr, cm^{-1}) = 3480, 3359, 3219 (br. 2NH & OH), 3078 (CH aromatic), 1698 (br. 2C=O); ^1H NMR (400 MHz, CDCl_3): δ = 12.86 (s, 1H, OH), 10.79 (d, J = 6.8 Hz, 1H, –NH–NH–CO), 8.88 (d, J = 6.6 Hz, 1H, –NH–NH–CO), 8.09 (d, J = 7.8 Hz, 2H, A–H), 7.91 (dd, J = 7.7, 1.8 Hz, 1H, A–H), 7.73–7.42 (m, 5H, A–H), 4.20 (dd, J = 12.5, 3.7 Hz, 1H, pyrrolidine CH_2), 3.94 (dd, J = 12.5, 3.6 Hz, 1H, pyrrolidine CH_2), 3.28–3.16 (m, 1H, pyrrolidine CH), 2.71 (dd, J = 15.7, 4.9 Hz, 1H, pyrrolidine CH_2), 2.49 (dd, J = 15.6, 4.9 Hz, 1H, pyrrolidine CH_2); ^{13}C NMR (100 MHz, CDCl_3): δ = 172.73 (C=O), 171.24 (C=O), 146.52, 143.90, 141.54, 139.45, 136.97, 136.51, 128.93, 126.45, 125.38, 124.48, 123.42, 119.11, 48.60 (pyrrolidine CH_2), 38.03 (pyrrolidine CH), 36.42 (pyrrolidine CH_2); MS (EI, 70 eV): m/z (%) = 408.60 [M^+] (13.86%), 147.50 (100%); anal. calc. for: $\text{C}_{19}\text{H}_{16}\text{N}_6\text{O}_5$ (408.37): C, 55.88; H, 3.95; N, 20.58; found: C, 55.86; H, 3.96; N, 20.55.

Conjugates **28–31** were afforded by reacting 2-chloro-3-hydrazinylquinoxaline (**6**) with fragments **18–21**.

3.1.2.5 *N'*-(3-Chloroquinoxalin-2-yl)-1-(4-cyanophenyl)-5-oxopyrrolidine-3-carbohydrazide, **28**. Brown powder; yield: (66%), m.p. 258–259 °C; IR (KBr, cm^{-1}) = 3367–3260 (br. 2NH), 3070 (CH aromatic), 2213 (CN), 1696 (br. 2C=O); ^1H NMR (400 MHz, CDCl_3): δ = 10.33 (d, J = 7.0 Hz, 1H, –NH–NH–CO), 8.97 (d, J = 7.0 Hz, 1H, NH–NH–CO), 8.01–7.86 (m, 2H, A–H), 7.61–7.51 (m, 4H, A–H), 7.30 (d, J = 7.7 Hz, 2H, A–H), 4.19 (dd, J = 12.5, 3.7 Hz, 1H, pyrrolidine CH_2), 3.94 (dd, J = 12.5, 3.6 Hz, 1H, pyrrolidine CH_2), 3.26–3.17 (m, 1H, pyrrolidine CH), 2.71 (dd, J = 15.7, 5.0 Hz, 1H, pyrrolidine CH_2), 2.49 (dd, J = 15.7, 4.9 Hz, 1H, pyrrolidine CH_2); ^{13}C NMR (100 MHz, CDCl_3): δ = 172.24 (C=O), 171.54 (C=O), 146.11, 141.32, 140.42, 137.80, 132.33, 131.31, 129.74, 127.06, 126.42, 125.81, 117.89, 117.18 (CN), 102.13, 48.78 (pyrrolidine CH_2), 38.09 (pyrrolidine CH), 36.37 (pyrrolidine CH_2); anal. calc. for: $\text{C}_{20}\text{H}_{15}\text{ClN}_6\text{O}_2$ (406.83): C, 59.05; H, 3.72; N, 20.66; found: C, 59.08; H, 3.70; N, 20.65.

3.1.2.6 *N'*-(3-Chloroquinoxalin-2-yl)-1-(2,5-dimethylphenyl)-5-oxopyrrolidine-3-carbohydrazide, **29**. Light brown powder; yield: (75%), m.p. 274–276 °C; IR (KBr, cm^{-1}) = 3349 (br. 2NH), 3044 (CH aromatic), 2904, 2840 (CH aliphatic), 1656 (br. 2C=O); ^1H NMR (400 MHz, CDCl_3): δ = 10.50 (d, J = 7.0 Hz, 1H, –NH–NH–CO), 8.16–8.01 (m, 2H, A–H), 7.94 (d, J = 7.0 Hz, 1H, NH–NH–CO), 7.61 (dd, J = 6.0, 3.6 Hz, 2H, A–H), 7.14 (d, J = 8.7 Hz, 1H, A–H), 7.00 (d, J = 7.3 Hz, 2H, A–H), 4.18 (dd, J = 12.2, 3.6 Hz, 1H, pyrrolidine CH_2), 3.93 (dd, J = 12.3, 3.6 Hz, 1H, pyrrolidine CH_2), 3.18–3.09 (m, 1H, pyrrolidine CH), 2.53 (dd, J = 15.6, 4.9 Hz, 1H, pyrrolidine CH_2), 2.34 (dd, J = 15.6, 4.9 Hz, 1H, pyrrolidine CH_2), 2.26 (s, 3H, CH_3), 2.09 (s, 3H, CH_3); ^{13}C NMR (100 MHz, CDCl_3): δ = 172.15 (C=O), 171.54 (C=O), 146.11, 140.42, 137.80, 137.35, 132.42, 131.31, 129.74, 129.63, 129.36, 127.06, 126.42, 125.81, 123.84, 121.82, 49.31 (pyrrolidine CH_2), 38.12 (pyrrolidine CH), 36.32 (pyrrolidine CH_2), 19.52 (CH_3), 16.53 (CH_3); MS (EI, 70 eV): m/z (%) = 411.38 [$\text{M} + 2$] (24.70%), 409.44 [M^+] (33.19%), 259.30 (100%); anal. calc. for: $\text{C}_{21}\text{H}_{20}\text{ClN}_5\text{O}_2$ (409.87): C, 61.54; H, 4.92; N, 17.09; found: C, 61.58; H, 4.90; N, 17.14.

3.1.2.7 *N'*-(3-Chloroquinoxalin-2-yl)-1-(3-hydroxyphenyl)-5-oxopyrrolidine-3-carbohydrazide, **30**. Brown powder; yield: (63%), m.p. 278–279 °C; IR (KBr, cm^{-1}) = 3418–3397 (br. 2NH & 2OH), 3076 (CH aromatic), 1651 (br. 2C=O); ^1H NMR (400 MHz, CDCl_3): δ = 10.27 (d, J = 7.0 Hz, 1H, –NH–NH–CO), 9.38 (d, J = 7.0 Hz, 1H, NH–NH–CO), 7.81–7.72 (m, 1H, A–H), 7.71–7.61 (m, 1H, A–H), 7.49–7.39 (m, 3H, A–H & OH), 7.06–6.97 (m, 2H, A–H), 6.55–6.49 (m, 1H, A–H), 6.43–6.34 (m, 1H, A–H), 4.24 (dd, J = 12.5, 3.7 Hz, 1H, pyrrolidine CH_2), 3.98 (dd, J = 12.5, 3.7 Hz, 1H, pyrrolidine CH_2), 3.17–3.09 (m, 1H, pyrrolidine CH), 2.70 (dd, J = 15.6, 4.9 Hz, 1H, pyrrolidine CH_2), 2.45 (dd, J = 15.6, 4.9 Hz, 1H, pyrrolidine CH_2); ^{13}C NMR (100 MHz, CDCl_3): δ = 173.48 (C=O), 172.44 (C=O), 158.81, 149.69, 141.26, 140.85, 139.36, 131.53, 131.16, 130.66, 129.32, 126.83, 124.68, 117.35, 109.00, 106.54, 49.02 (pyrrolidine CH_2), 40.79 (pyrrolidine CH), 38.00 (pyrrolidine CH_2); MS (EI, 70 eV): m/z (%) = 397.20 [$\text{M} + 2$] (33.98%), 399.22 [M^+] (25.48%), 379.93 (100%); anal. calc. for: $\text{C}_{19}\text{H}_{16}\text{ClN}_5\text{O}_3$ (397.82): C, 57.37; H, 4.05; N, 17.60; found: C, 57.33; H, 4.09; N, 17.57.

3.1.2.8 *N'*-(3-Chloroquinoxalin-2-yl)-1-(4-nitrophenyl)-5-oxopyrrolidine-3-carbohydrazide, **31**. Dark yellow powder; yield: (51%), m.p. 266–267 °C; IR (KBr, cm^{-1}) = 3420–3342 (br. 2NH & OH), 3067 (CH aromatic), 1686 (br. 2C=O); ^1H NMR (400 MHz, CDCl_3): δ = 10.50 (d, J = 7.0 Hz, 1H, –NH–NH–CO), 8.15–8.01 (m, 4H, A–H), 7.94 (d, J = 7.0 Hz, 1H, NH–NH–CO), 7.61 (dd, J = 6.0, 3.5 Hz, 2H, A–H), 7.40 (d, J = 7.8 Hz, 2H, A–H), 4.17 (dd, J = 12.5, 3.7 Hz, 1H, pyrrolidine CH_2), 3.92 (dd, J = 12.5, 3.6 Hz, 1H, pyrrolidine CH_2), 3.12 (p, J = 4.2 Hz, 1H, pyrrolidine CH), 2.52 (dd, J = 15.7, 4.9 Hz, 1H, pyrrolidine CH_2), 2.33 (dd, J = 15.6, 4.9 Hz, 1H, pyrrolidine CH_2); ^{13}C NMR (100 MHz, CDCl_3): δ = 172.65 (C=O), 171.12 (C=O), 146.45, 143.82, 139.83, 139.37, 138.67, 130.84, 129.01, 126.36, 126.02, 125.18, 124.41, 119.04, 48.53 (pyrrolidine CH_2), 37.94 (pyrrolidine CH), 36.35 (pyrrolidine CH_2); MS (EI, 70 eV): m/z (%) = 428.41 [$\text{M} + 2$] (10.83%), 426.69 [M^+] (19.33%), 263.76 (100%); anal. calc. for:



C₁₉H₁₅ClN₆O₄ (426.82): C, 53.47; H, 3.54; N, 19.69; Found: C, 53.44; H, 3.50; N, 19.67.

3.2. Biology

3.2.1. Agar diffusion approach. The action of the antibacterial was assessed utilizing the well-diffusion of Agar test (Sabaraud Dextrose Agar plates (fungi) and Tryptic Soya Agar plates (bacteria)) (Oxoid Ltd, UK). Eight different hybrids were examined for antimicrobial action taken in opposition to test organisms, namely, Gram-positive bacteria (Methicillin sensitive *S. aureus* (MSSA) ATCC 25923), Bacteria that are Gram-negative (*P. aeruginosa* ATCC 27853, *A. baumannii* ATCC 19606, *K. pneumonia* ATCC 700603, *E. coli* ATCC 25922), and fungal strain (*C. albicans* ATCC 10231). Fresh cultures were utilized to create the bacterial suspensions, which were then adjusted with saline to 0.5 McFarland-standard (pertaining to 1×10^8 CFU mL⁻¹). To attain an ultimate accumulation of 1×10^6 CFU mL⁻¹, the fungal suspensions were likewise modified. A target concentration of 5 mM was obtained by dissolving the investigated hybrids in DMSO. 200 μ L of each hybrid were applied. Measurements of surrounding clear zones of inhibition were taken and compared to the negative command. Dimethylsulphoxide (DMSO) functioned as a negative control. Fluconazole as well as Levofloxacin were utilized as standard. Incubation of the plates lasted 24 hours for bacteria and 48 hours for fungus at 35 ± 2 °C. The data were asserted using the well-diffusion technique in triplicate.^{84–86}

3.2.2. Evaluation of MIC. 96-well microtiter plates with serial dilutions of the hybrids in Tryptic Soy Broth (TSB-Sigma®) medium were employed to measure the compounds' MIC implementing the broth microdilution method (BMM). The hybrids were prepared as stock solutions in DMSO/TSB 1 : 10 v/v. The concentrations varied between 1.5 and 200 μ g mL⁻¹, with fluconazole, levofloxacin, and amoxicillin serving as drug controls. Briefly, in a microtiter plate, 100 μ L of the chemicals to be tested were combined with 100 μ L of TSB and 5 μ L of 1×10^8 CFU mL⁻¹ of the strain under investigation. For bacteria or fungi, incubation of the plates lasted at 35 ± 2 °C for 24 or 48 hours, respectively. MIC was the least amount of concentration of substance which showed no signs of growth.^{87–90}

3.2.3. Evaluation of MBC and MFC. After transferring 10 μ L of MIC-tests culture either plates of Tryptic Soya-Agar or Sabaraud Dextrose-Agar and incubating them overnight at 35 ± 2 °C for 24 hours or 48 hours for bacteria or fungi, MBC and MFC of the substances under test were identified. MBC/MFC is clearly the lowest concentration, resulting in a 99.9% decrease in the growth of bacteria. If the MBC to MIC or MFC to MIC rate falls between 1 : 1 and 2 : 1, the investigated conjugate was deemed bactericidal or fungicidal; if the ratio is more than 2 : 1, it was deemed bacteriostatic or fungistatic.^{91–95}

3.2.4. Anti-biofilm activity. Using TSB, the inhibitory impact of studied hybrids on biofilm development was evaluated in 96-well polystyrene flat-bottom plates. Briefly, 200 μ L of freshly inoculated TSB with 1% glucose was aliquoted into plate wells, resulting in a last focus of 10^6 CFU per mL of the tested strain. The culture was conducted in the presence of sub-

inhibitory doses (50 and 90% of MIC). To remove all of the planktonic bacterial cells, the cups had next cultured for 24 hours at 37 °C, after that washed three times with PBS. Following the washing stage, the cups had opened inside a hot air at 60 °C for 60 minutes, then 150 μ L of crystal-violet (CV) at a 0.1% concentration was put into every well for 15 minutes for fixing and staining. To get rid of any remaining CV stains, the washing process was reapplied. Addition of 150 μ L of 100% ethanol to each well followed the washing process and the well OD at 630 nm was measured in a 96-well plate spectrophotometer (800 TS Absorbance Reader, Biotek, USA). Every plate included sterility and growth controls. Ultimately, The formula below was utilized to ascertain the inhibitory percentages of biofilm: biofilm inhibitory (%) = $[1 - \text{OD nm isolate}/\text{OD nm of positive control}] \times 100$.^{96–103}

3.2.5. Time kill kinetics. The bactericidal efficacy of the conjugates under evaluation was confirmed by the time-kill experiment. *P. aeruginosa* ATCC 27853 bacterial cultures in the exponential growth phase (1×10^6 CFU mL⁻¹) were administered treatment in drug-containing tubes under examination at 8-fold, 4-fold, 2-fold, 1-fold and 0.5-fold MIC, and incubated at 37 °C. Viable count measurements were made using samples collected at 0, 3, 6, 12, and 24 hours of incubation. After being serially diluted, each sample was spread out across two to four agar plates. CFU were established following 18 to 24 hours of the incubated at 35 °C in 5% CO₂. 10 CFU mL⁻¹ was the limit of detection (LOD). Visual examination of the colony dispersion on the plates was used to gauge drug carryover. Growth control was included, involving an experiment without addition of hybrids. A plot of log 10 CFU mL⁻¹ vs. time (h) was used to depict the bactericidal kinetics of conjugates.^{104–109}

3.2.6. DNA gyrase inhibition assay. See SI.^{110–115}

3.3. In silico ADMET

Pharmacokinetics prediction was performed as reported.^{116–123}

3.4. Molecular docking

The 3D layout for DNA gyrase (PDB.: 8BN6) had been extracted through data bank of the protein (<https://www.rcsb.org/>) and prepared using UCSF Chimera.^{124–127} Briefly, the mol2 file of protein was set by deleting water molecules, adding and charges and energy minimization. The generated compounds and respective co-crystallized ligands were sketched in Avogadro software, and their geometries were optimized.^{128,129} The prepared protein was imported in AutoDock. Hydrogens were added and Kollman charges and AD4 atom parameters were assigned. The grid calculations were set up; $-5.5 \times 23.5 \times 1.1$ Å grid with dimensions of 35, 35, 35 respectively, at a resolution of 0.375 Å, was created around the compound. Docking commenced using genetic algorithm search parameters. During docking, partial Gasteiger charges were applied. Docked conformations were generated using the Lamarckian evolutionary method and arranged according to their docking energy in descending order.^{130–134} Results visualizations were performed using Biovia Discovery Studio. The same steps were



repeated for the co-crystallized ligand in order to perform validation. The RMSD value of the redocked pose was calculated using DockRMSD online tool.^{135,136}

4. Conclusion

This study successfully synthesized and studied several novel hybrid chemicals, such as quinoxaline, pyrrolidine, and a hydrazinyl bridge, indicating their potential as strong antibacterial and antifungal medicines. Several compounds had strong efficacy against different pathogenic strains, according to the *in vitro* tests, with eight derivatives demonstrating notable bactericidal effects. Eight novel conjugates were tested for antibacterial activity at a dosage of 5 mM. All synthesized derivatives caused *E. coli* growth ZOI \geq 15 mm, with hybrids **26**, **28**, **30**, and **31** showing greater reduction than Amoxicillin and Levofloxacin. Hybrids **24**, **27**, and **29** were confirmed as bactericidal agents against *E. coli*. Hybrids **24**, **25**, and **31** displayed MIC of 12.5 μ M, whereas they indicate bactericidal hybrids at higher concentrations against *P. aeruginosa*. Treatment with involving all hybrids resulted in 89–92% decrease in biofilm formation at 90% MIC, outperforming Levofloxacin. The killing kinetics of eight hybrids against *P. aeruginosa* were time-dependent, with higher concentrations leading to a rapid decrease in CFU number. Primary bacterial elimination occurred after 3 hours, while 4-fold and 8-fold MICs led to almost complete elimination. Hybrids **25**, **28**, and **31** were the most efficient DNA gyrase inhibitors; their IC₅₀ values were significantly lower than those of ciprofloxacin (77.3, 87.6, and 65.5 μ M). The study reveals that molecular docking studies reveal effective compounds' binding interactions, confirming their biological activities. Compounds having drug-like properties are identified by examining their physicochemical and pharmacokinetic properties, indicating the possibility of further development. With future research concentrating on optimizing lead compounds for clinical applications, the findings support the continuing attempts to develop novel treatment medicines for infectious disorders and resistant microbial strains.

Conflicts of interest

The author(s) report no conflicts of interest in this work.

Data availability

The data supporting this article have been included as part of the supplementary information (SI). Supplementary information is available. See DOI: <https://doi.org/10.1039/d6ra00675b>.

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