


In silico Molecular Docking, DFT Analysis and ADMET Studies of Carbazole Alkaloid and Coumarins from Roots of *Clausena anisata*: A Potent Inhibitor for Quorum Sensing

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Introduction: In modern drug design, in silico methods are largely used to understand drug-receptor interactions and quantum chemical properties. In the present study, a computational de novo design approach was used to confirm mode of binding for antibacterial activity, elucidating quantum chemical properties and ADMET-drug-likeness of carbazole alkaloid (1) and three coumarins (2–4) isolated from roots of *Clausena anisata*.

Methods: Docking studies were performed with DNA-Gyrase (6F86) and LasR binding domain (2UV0) employing a flexible ligand docking approach using AutoDock Vina. SwissADME prediction and toxicological predictions were performed by ADMET. The optimized structures and molecular electrostatic potential surface of the isolated compounds were predicted by DFT analysis using B3LYP/6-31G basis levels.

Results and Discussion: The docking results revealed that compound 3 showed better docking scores against both DNA gyrase B and LasR binding domain compared with ciprofloxacin with potential as an inhibitor of bacterial DNA gyrase and quorum sensing LasR binding domain. The SwissADME prediction results showed that all the isolated compounds (1–4) satisfy Lipinski's rule of five with zero violations. Toxicological prediction results suggested that all compounds and ciprofloxacin are non-hepatotoxic, non-carcinogenic, non-irritant, immunogenic, and non-cytotoxic. The DFT analysis results revealed that compound 3 has large electronegativity (χ_eV), global softness (σeV^{-1}), global electrophilicity (ω_eV), and mutagenicity value closer to ciprofloxacin (with LD_{50} value of 480 mg/kg) suggesting better bioactivity and chemical reactivity with considerable intra-molecular charge transfer between electron-donor to electron-acceptor groups.

Conclusion: Overall, compound 3 may serve as a lead molecule that could be developed into a potent *E. coli* DNA gyrase B inhibitor and efficient inhibitor for quorum sensing auto-inducer LasR binding domain of *Pseudomonas aeruginosa*.

Keywords: alkaloids, coumarins, DNA gyrase, LasR binding domain, de novo DFT and docking studies

Introduction

Clausena anisata is one of the traditional medicinal plants that belong to the family of Rutaceae used for the treatment of bacterial and fungal infections.^{1–4} Recently, our group reported isolation and antibacterial activity of a carbazole alkaloid (1) and three coumarins (2–4) from the roots of *Clausena anisata* against strains of *Staphylococcus aureus*, *Escherichia coli*, *Pseudomonas aeruginosa* and *Bacillus*

subtilis.² Additionally, the alkaloids and coumarins have been reported to be active against both Gram-positive and Gram-negative bacteria and fungi.^{2,3,5-13} However, the mechanism of action of these compounds has not been previously validated experimentally and little assessment was made on the potential “drug-likeness” of these compounds. In the discovery of new and complementary antibacterial agents, phytochemicals that show antibacterial activity can be examined for potential inhibition of bacterial target proteins such as peptide deformylase (PDF), topoisomerase IV (TopoIV), DNA gyrase B (GyrB), protein tyrosine phosphatase (Ptp), UDP-galactopyranosemutase (UGM), cytochrome P450 (CYP121), and NAD⁺-dependent DNA ligase, as well as phytochemical inhibitors of bacterial efflux pumps or quorum sensing proteins, or agents that enhance the immune system.

Pseudomonas aeruginosa is the most extensively studied strain due to the potentially fatal effects of its infections, that are responsible for many deaths each year worldwide. It is the most common Gram-negative bacterium found in hospital-acquired infections, being responsible for nosocomial pneumonia, urinary tract infections, and surgical wound or bloodstream infections. In general, many Gram-negative bacteria coordinate the activities of their populations through autoinducers.¹⁴ This is called quorum sensing and it can regulate pathogenic virulence factor production and antimicrobial resistance. Quorum sensing inhibitors can diminish the pathogenicity of *P. aeruginosa*.¹⁴⁻¹⁶ The emergence of LasR as a key regulator of *P. aeruginosa* pathogenesis has made its inhibition an interesting pharmaceutical goal. Additionally, DNA topoisomerases are crucial enzymes for maintaining the topological state of DNA.¹⁷⁻¹⁹ The ATP-dependent bacterial type II topoisomerases, DNA gyrase and DNA topoisomerase IV are responsible for the maintenance of the correct level of supercoiling in bacterial DNA. These enzymes have been well-validated as antibacterial targets as the fluoroquinolone antibiotics act at the DNA-cleavage site of these enzymes.²⁰ The aminocoumarin antibiotics, including novobiocin, which was used clinically in the 1960s, target the ATP-binding site.²¹ Therefore, in the present study we examined the binding modes of reported inhibitors (coumarins and a carbazole alkaloid from the roots of *Clausena anisata*) within bacterial DNA-GyrB and LasR, considering their promising antibacterial activity along with DFT analysis for optimized structure, drug-likeness, pharmacokinetics, toxicity profile and compared with the clinically approved drug ciprofloxacin.

Materials and Methods

Preparation of Ligands

The 2D structures (.mol) of all compounds were drawn (Figure 1) and analyzed using ChemDraw 16.0. All the compounds (1-4) are converted to 3D structure (.pdb) using Chem3D 16.0. The 3D coordinates (.pdb) of each molecule were loaded on to Chem3D for energy minimization.

Preparation of Macromolecule

The protein target, which is retrieved from the RCSB Protein Data Bank (PDB code 6F86 and 2UV0), serves as docking receptor.^{15,22} All the bound ligands and water molecules were removed from the active site of the receptor.

Molecular Docking Analysis (AutoDock Vina)

The molecular docking studies were carried out using AutoDockTools (ADT), which is a free graphic user interface (GUI) for the AutoDockVina program.²³ AutoDockVina with standard protocol was used to dock the compounds (1-4) against the active site of protein (PDB ID: 6F86 and 2UV0).^{15,22,23} The grid box was constructed using 58, 58, and 40, pointing in x, y, and z directions, respectively, with a grid point spacing of 0.375 Å. The center grid box is of 14.527 Å, 56.689 Å and -5.122 Å. Nine different conformations were generated for each ligand scored using AutoDockVina scoring functions (Supplementary materials [Tables 1-8]) and ranked according to their binding energies (Supplementary materials). AutoDock Tools and PyMOL were used for the post-docking analyses. The conformations with the most favourable (least) free binding energy were selected for analysing the interactions between the target receptor and ligands by PyMOL.

In silico Drug-Likeness and Toxicity Predictions

Structures of compounds (1-4) were submitted to SwissADME tool and converted to their canonical simplified molecular input line entry system (SMILE), to estimate in silico pharmacokinetic parameters and other molecular properties based on the methodology reported by Amina et al., 2016.²⁴ SwissADME predictor provided information on the numbers of hydrogen donors, hydrogen acceptors and rotatable bonds, total polar surface area and the synthetic accessibility of the compounds. The ligands were also subjected to Lipinski et al. screenings using SwissADME predictor.

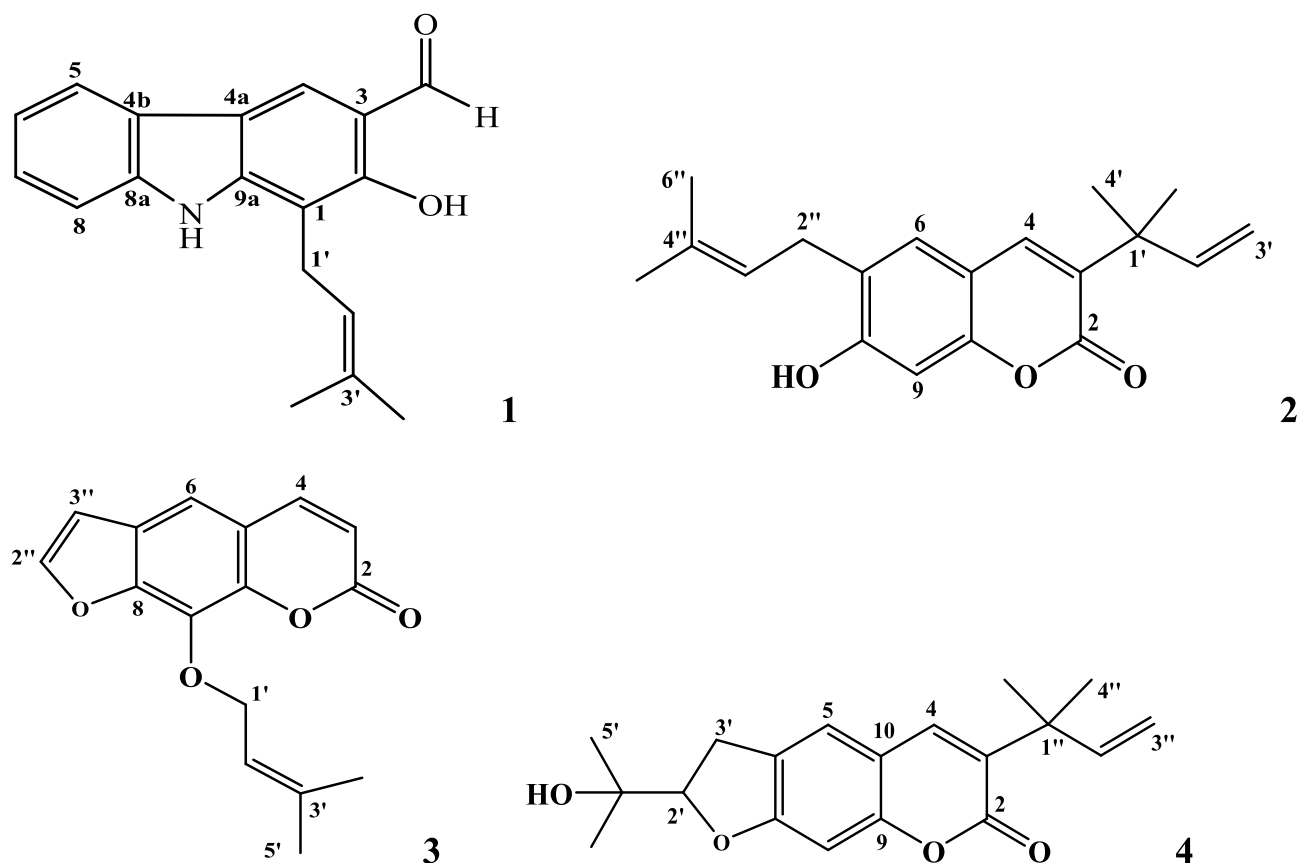


Figure 1 2D-ChemDraw structures of isolated compounds (1–4).

Drug-likeness is a prediction that determines whether a particular pharmacological agent has properties consistent with being an orally active drug. This prediction is based on an already established concept by Lipinski et al., called the Lipinski rule of five. The rule predicts that there is likely to be poor absorption or permeation when a compound possesses more than 5 H-bond donors, 10 H-bond acceptors, molecular weight greater than 500 and a calculated LogP (CLogP) greater than 5. The selection of compounds as drug candidates were determined by a parameter called drug score. The higher the drug score value, the higher the chance of the compound being considered as a drug candidate. The organ toxicities and toxicological endpoints of the ligands and their LD₅₀ were predicted using Pro Tox II.^{25–27} The analyses of the compounds were compared with that of ciprofloxacin, and only compounds without violation of any of the screenings were used for the molecular docking analysis.

Quantum Computational Studies

The DFT (density functional theory) analysis of compounds (1–4) were performed using Gaussian 09 and

visualized through Gauss view 5.0. The structural coordinates of the lead compounds were optimized using B3LYP/6-31 G (d,p) level basis set without any symmetrical constraints. The molecular electrostatic potential map and energies of the compounds were obtained from the optimized geometry. Koopman's approximation was used to estimate the HOMO-LUMO energy gap and related reactive parameters (electronegativity, chemical potential, hardness, softness, electrophilicity).²⁸

Results and Discussion

Molecular Docking Against DNA Gyrase B

Molecular docking analysis of isolated compounds showed better docking score within the active site of *E. coli* (6F86). Compounds 1–3 (–7.2, –7.7, and –7.6 kcal/mol, respectively) showed equal to better docking affinity than the preclinical drug ciprofloxacin (–7.2 kcal/mol) whereas compound 4 showed smaller docking affinity (–6.6 kcal/mol). All the molecules showed binding affinity within the binding sites of co-

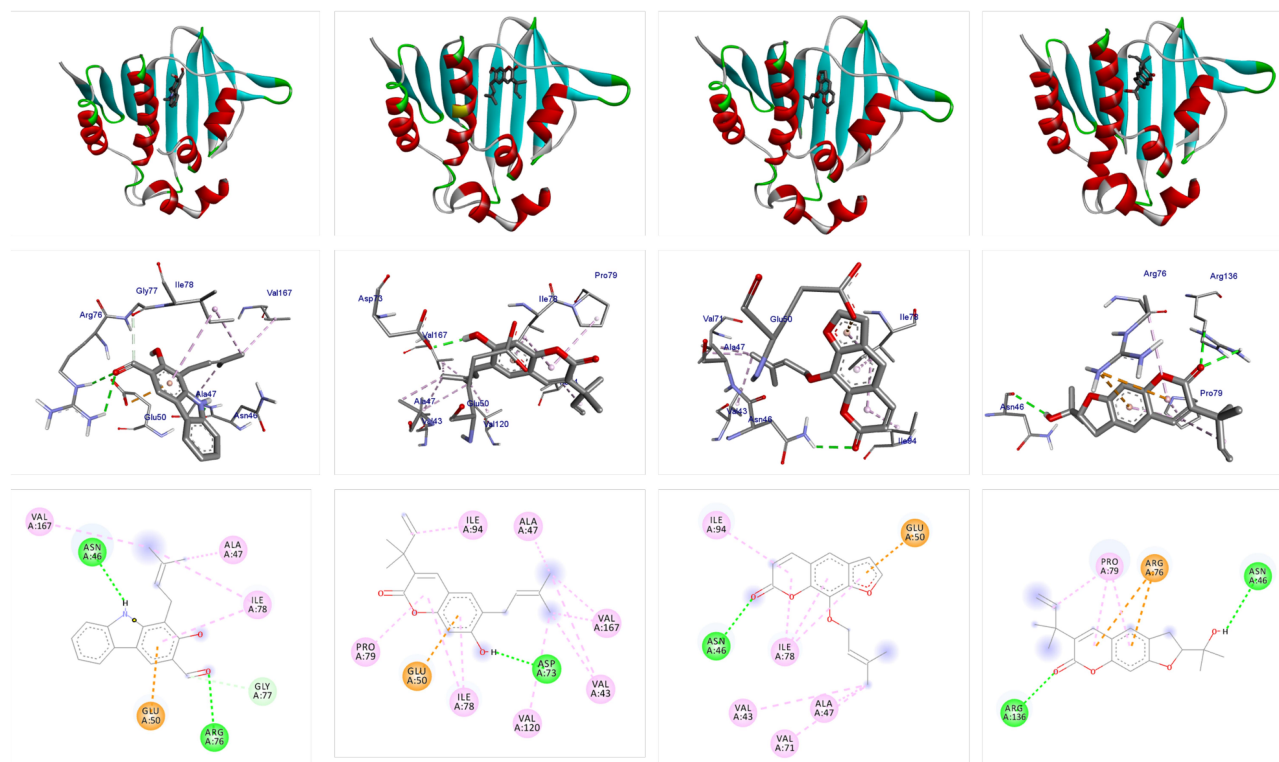


Figure 2 The 2D and 3D binding interactions of compounds (1–4) against DNA gyrase B (PDB ID: 6F86). Ribbon model shows the binding pocket structure of DNA gyrase B with compounds (1–4). Hydrogen bond between compounds and amino acids are shown as green dashed lines, hydrophobic interactions are shown as pink lines.

crystallized ligand (Figure 2). Compound 1 formed hydrogen bond interaction with Arg-76, Asn-46, Gly-77 and residual hydrophobic interaction with Ala-47, Glu-50, Ile-78, Val-167. Similarly, compound 2 showed one hydrogen bond (Asp-73) and residual hydrophobic interactions with amino acids Glu-50, Val-43, Val-47, Ile-78, Pro-79, Ile-94, Val-120, and Val-167. Compounds 3 and 4 both showed hydrogen bond interaction with Asn-46 whereas compound 4 showed an additional hydrogen bond with Arg-136 and hydrophobic residual interaction with Arg-76 and Pro-79. Compounds 1–3 demonstrated a similar binding affinity with residual amino acids compared with ciprofloxacin

(hydrogen bond with residual amino acid Asp-73, Asn-46, Arg-76 and hydrophobic or pi-cation interaction with residual amino acids Glu-50, Ala-47, Gly-77, Ile-78, Pro-79, Ile-94, Ile-78). However, compounds 1 and 4 do not show the crucial interactions between the ligand and amino acid Asp-73. In silico docking results revealed that compounds 2 and 3 have least docked score (–7.7 and –7.6 kcal/mol, respectively) compared with other compounds and comparable residual interactions with that of ciprofloxacin within the binding pocket (Table 1), suggesting that these compounds are promising antibacterial agents against *E. coli*.^{6,8,29,30}

Table 1 Molecular Docking Scores and Residual Amino Acid Interactions of Compounds 1–4 Against *E. coli* DNA gyrase B (PDB ID 6F86)

S. No.	Ligands	Affinity (kcal/mol)	H-bond	Residual Hydrophobic/Pi-Cation/Pi-Anion/Pi-Alkyl Interactions
1	C ₁₈ H ₁₇ N ₁ O ₂	–7.2	Arg-76, Asn-46, Gly-77	Ala-47, Glu-50, Ile-78, Val-167
2	C ₁₉ H ₂₂ O ₃	–7.7	Asp-73	Glu-50, Val-43, Val-47, Ile-78, Pro-79, Ile-94, Val-120, Val-167
3	C ₁₆ H ₁₄ O ₄	–7.6	Asn-46	Glu-50, Val-43, Val-47, Ile-78, Val-71, Ile-94
4	C ₁₉ H ₂₂ O ₄	–6.6	Arg-136, Asn-46	Arg-76, Pro-79
5	Ciprofloxacin	–7.2	Asp-73, Asn-46, Arg-76	Glu-50, Ala-47, Gly-77, Ile-78, Pro-79, Ile-94, Ile-78

Molecular Docking Against Protein LasR Binding Domain

LasR is a key regulator of *P. aeruginosa* pathogenesis.^{31,32} Molecular docking of compounds against LasR binding domain displayed better docking scores toward *P. aeruginosa* (2UV0) compared with the DNA gyrase B docking results. Binding mode of active compounds (1–4) demonstrated docked scores of –8.9, –9.0, –9.3, and –9.2 kcal/mol, respectively. All isolated compounds showed docking interaction within the binding sites of co-crystallized quorum sensing auto-inducer molecules N-3-oxo-dodecanoyl-L-homoserine lactone (OHN) where hydrogen bonding interactions were observed with residual amino acids similar to OHN (Ser-129, Asp-73, Tyr-64, and Trp-60) and hydrophobic interactions with amino acid residues Leu-40, Tyr-47, Ala-50, Ala-70, Val-76, and Leu-125 (Figure 3). Compound 1 showed two hydrogen bonding interactions with amino acid residues Thr-75 and Thr-115 and residual hydrophobic interaction with amino acid residues Asp-73, Tyr-64, Val-76, Leu-36, Ile-52, Ala-50, Ala-70, Tyr-47, Cys-79, Leu-125, and Ala-127. Compounds 2 and 3

showed formation of hydrogen bonding with amino acids Ser-129, Thr-115 and hydrophobic interaction with amino acid residues Asp-73, Tyr-64, Leu-36, Ile-52, Ala-50, Tyr-47, and Ala-127 within the binding pocket (Table 2). In addition, compound 3 showed two more hydrogen bonding interactions with Tyr-47 and Thr-75. Compound 4 (–9.2 kcal/mol) showed one hydrogen bond interaction with amino acid Leu-110 and hydrophobic interaction with residual amino acids Asp-73, Tyr-64, Leu-36, Val-76, Ala-50, Ala-70, Tyr-47, and Ala-127. Molecular docking results showed that all the molecules have least docking score compared with co-crystallized ligand (OHN) and clinical drug (ciprofloxacin). The docking results are in good agreement with our previous *in vitro* antibacterial analysis where compounds 3 and 4 displayed better antibacterial activity against *P. aeruginosa* (14 ± 0.1 and 12 ± 0.2 mm zone of inhibition, respectively) compared with ciprofloxacin (15 ± 0.3 mm zone of inhibition)^{2,3,14,16,31} suggesting compounds 3 and 4 may be better antibacterial agents specifically through quorum sensing activity.

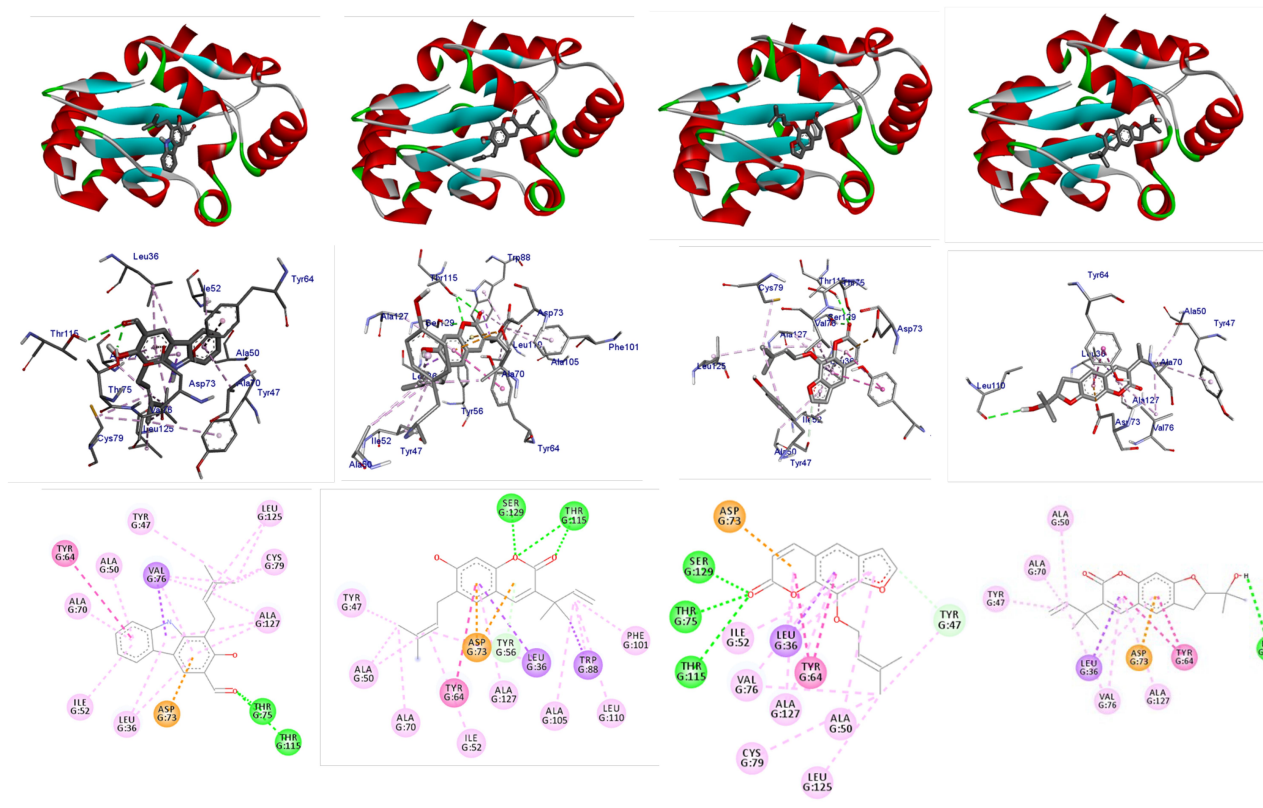


Figure 3 The 2D and 3D binding interactions of compounds (1–4) against LasR binding domain (PDB ID 2UV0). Ribbon model shows the binding pocket structure of LasR binding domain with compounds (1–4). Hydrogen bond between compounds and amino acids are shown as green dashed lines, hydrophobic interactions are shown as pink lines.

Table 2 Molecular Docking Scores and Residual Amino Acid Interactions of Compounds **1–4** Against LasR Binding Domain

S. No.	Ligands	Affinity (kcal/mol)	H-bond	Residual Hydrophobic/Pi-Cation/Pi-Anion/Pi-Alkyl Interactions
1	C ₁₈ H ₁₇ N ₁ O ₂	-8.9	Thr-75, Thr-115	Asp-73, Tyr-64, Val-76, Leu-36, Ile-52, Ala-50, Ala-70, Tyr-47, Cys-79, Leu-125, Ala-127
2	C ₁₉ H ₂₂ O ₃	-9.0	Ser-129, Tyr-56, Thr-115	Asp-73, Tyr-64, Trp-88, Leu-36, Ile-52, Ala-50, Ala-70, Tyr-47, Phe-101, Ala-105, Leu-110, Ala-127
3	C ₁₆ H ₁₄ O ₄	-9.3	Ser-129, Thr-75, Tyr-47, Thr-115	Asp-73, Tyr-64, Leu-36, Ile-52, Val-76, Ala-50, Cys-79, Leu-125, Ala-127
4	C ₁₉ H ₂₂ O ₄	-9.2	Leu-110	Asp-73, Tyr-64, Leu-36, Val-76, Ala-50, Ala-70, Tyr-47, Ala-127
5	Ciprofloxacin	-8.3	Asp-73, Tyr-47, Trp-60	Leu-36, Tyr-64, Val-76, Cys-79, Ala-127

In silico Pharmacokinetics (Drug-Likeness) and Toxicity Analysis

The SwissADME computed results showed that all the isolated compounds (**1–4**) satisfy Lipinski's rule of five with zero violations (Table 3). The K_p values of all molecules are within the range of -4.26 to -5.47 cm/s with ciprofloxacin (-8.98 cm/s) inferring low skin permeability. The predicted logP values revealed that they have optimal lipophilicity (ranging from 2.43 to 4.37). The SwissADME prediction parameters showed that all the compounds **1–4** have high gastrointestinal (GI) absorption, blood-brain barrier (BBB) permeation and no compounds are substrates of permeability glycoprotein (P-gp). The CYP's interaction result showed that all the compounds are inhibitors for CYP1A2, CYP2C19, and CYP2C9. Compounds

1 and **4** were found to be potential inhibitors for CYP2D6 whereas compounds **2**, **3**, and ciprofloxacin were found to be non-inhibitors. For CYP3A4, compound **1** and ciprofloxacin were found to be non-inhibitors whereas compounds **2–4** were found to be potential inhibitors (Table 4).

Acute toxicity prediction results such as LD₅₀ values and toxicity class classification [1 (toxic) to 6 (non-toxic)] revealed that none of the ligands demonstrated acute toxicity similar to ciprofloxacin with toxicity class classification **4** (harmful if swallowed) for compounds **3**, **4** and ciprofloxacin. Toxicological prediction results suggested that all compounds and ciprofloxacin are non-hepatotoxic, non-carcinogenic, non-irritant, immunogenic and non-cytotoxic with the exception of compound **3** that displayed a mutagenicity value closer to ciprofloxacin (with LD₅₀ value of 480 mg/kg) suggesting this compound can be the

Table 3 Drug-Likeness Predictions of Compounds **1–4**, Computed by SwissADME

S. No.	Formula	Mol.Wt. (g/mol)	NHD	NHA	NRB	TPSA (Å ²)	LogP (cLogP)	Lipinski's Rule of Five Violation
1	C ₁₈ H ₁₇ N ₁ O ₂	279.33	2	2	0	53.09	2.57	0
2	C ₁₉ H ₂₂ O ₃	298.38	1	3	1	50.44	2.43	0
3	C ₁₆ H ₁₄ O ₄	270.28	0	4	2	52.58	3.05	0
4	C ₁₉ H ₂₂ O ₄	314.38	1	4	2	59.67	3.47	0
5	Ciprofloxacin	401.45	2	9	5	136.13	0.963	0

Abbreviations: NHD, number of hydrogen donor; NHA, number of hydrogen acceptor; NRB, number of rotatable bonds; TPSA, total polar surface area.

Table 4 ADME Predictions of Compounds **1–4**, Computed by SwissADME and PreADMET

S. No.	Chemical Formula	log K _p cm/s	GI Absorption	BBB Permeability	Inhibitor Interaction (SwissADME/PreADMET)					
					P-gp Substrate	CYP1A2 Inhibitor	CYP2C19 Inhibitor	CYP2C9 Inhibitor	CYP2D6 Inhibitor	CYP3A4 Inhibitor
1	C ₁₈ H ₁₇ N ₁ O ₂	-4.48	High	Yes	No	Yes	Yes	Yes	Yes	No
2	C ₁₉ H ₂₂ O ₃	-4.26	High	Yes	No	Yes	Yes	Yes	No	Yes
3	C ₁₆ H ₁₄ O ₄	-5.46	High	Yes	No	Yes	Yes	Yes	No	Yes
4	C ₁₉ H ₂₂ O ₄	-5.47	High	Yes	No	Yes	Yes	Yes	Yes	Yes
5	Ciprofloxacin	-8.98	High	No	Yes	Yes	No	No	No	No

Abbreviations: Log K_p, skin permeation value; GI, gastro-intestinal; BBB, blood-brain barrier; P-gp, P-glycoprotein; CYP, cytochrome-P.

Table 5 Prediction of Toxicity of Compounds (1–4), Computed by Pro-Tox II and OSIRIS Property Explorer

S. No.	Formula	LD ₅₀ (mg/kg)	Toxicity Class	Organ Toxicity					
				Hepatotoxicity	Carcinogenicity	Immunotoxicity	Mutagenicity	Cytotoxicity	Irritant
1	C ₁₈ H ₁₇ N ₁ O ₂	2000	3	No	No	No	Yes	No	No
2	C ₁₉ H ₂₂ O ₃	2905	5	No	No	No	Yes	No	No
3	C ₁₆ H ₁₄ O ₄	480	4	No	No	Yes	Yes	No	No
4	C ₁₉ H ₂₂ O ₄	1500	4	No	No	No	Yes	No	No
5	Ciprofloxacin	500	4	No	No	No	Yes	No	No

lead compound compared with other isolated compounds in the investigation (Table 5).

DFT Analysis for Optimized Structure and Quantum Chemical Parameters

Optimized geometries (Figure 4) and molecular electrostatic potential (MEP) of isolated compounds were computed using density functional theory (DFT) with the basis sets B3LYP/6-31 G (d,p). Bond lengths, bond

angles, and dihedral angles were obtained from the optimized structure (Table 6). The determination of the MEP region is the best fit for identifying sites for intra- and inter-molecular interactions (Figure 7).³³ Red/yellow regions indicate negative electrostatic potentials and the blue region shows positive, and green color designates neutral potential region.

The DFT calculated Mulliken's atomic charges (Figure 5) revealed charge distribution in individual atoms (Supplementary materials [Tables 9–16]). The

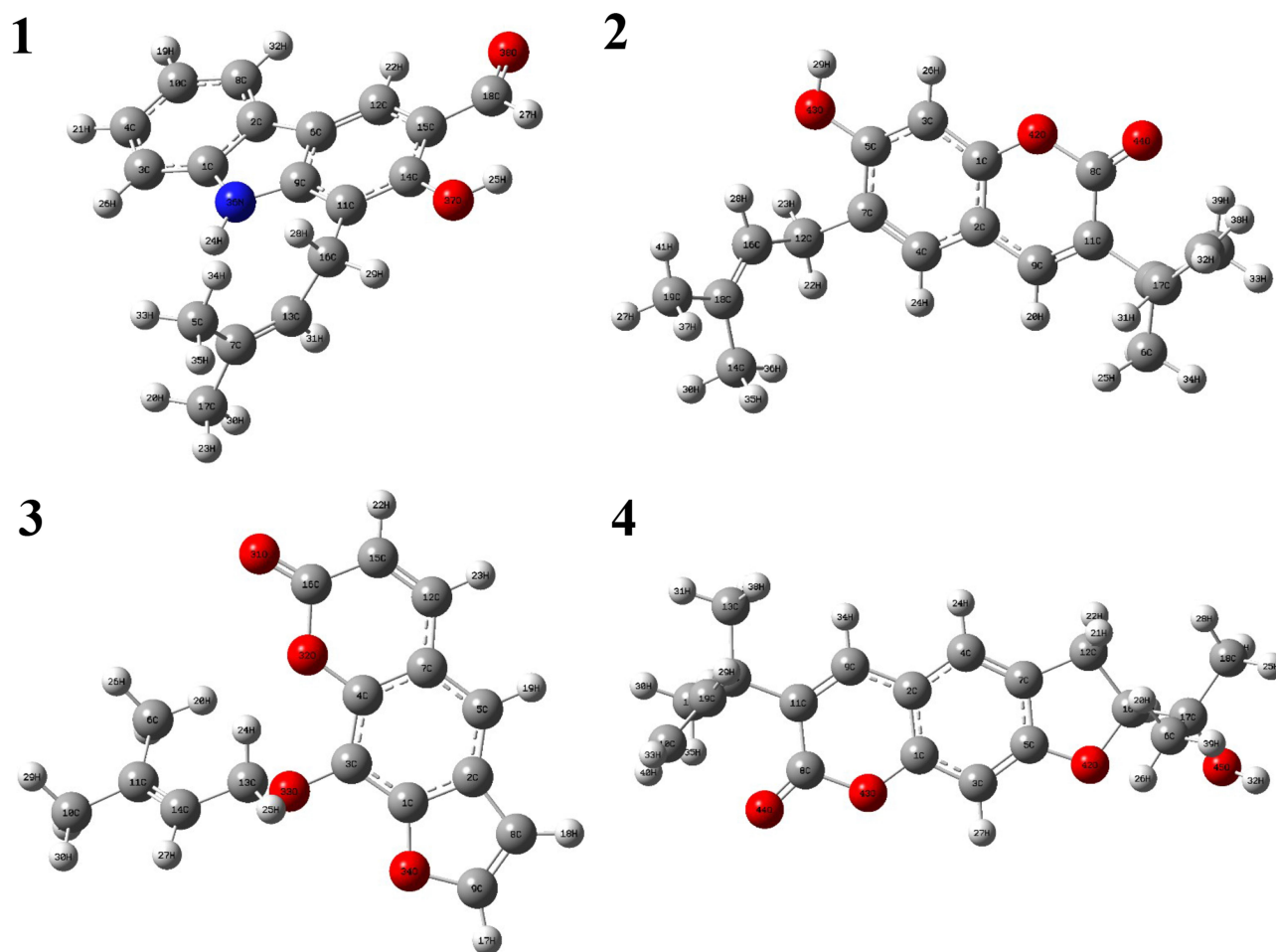
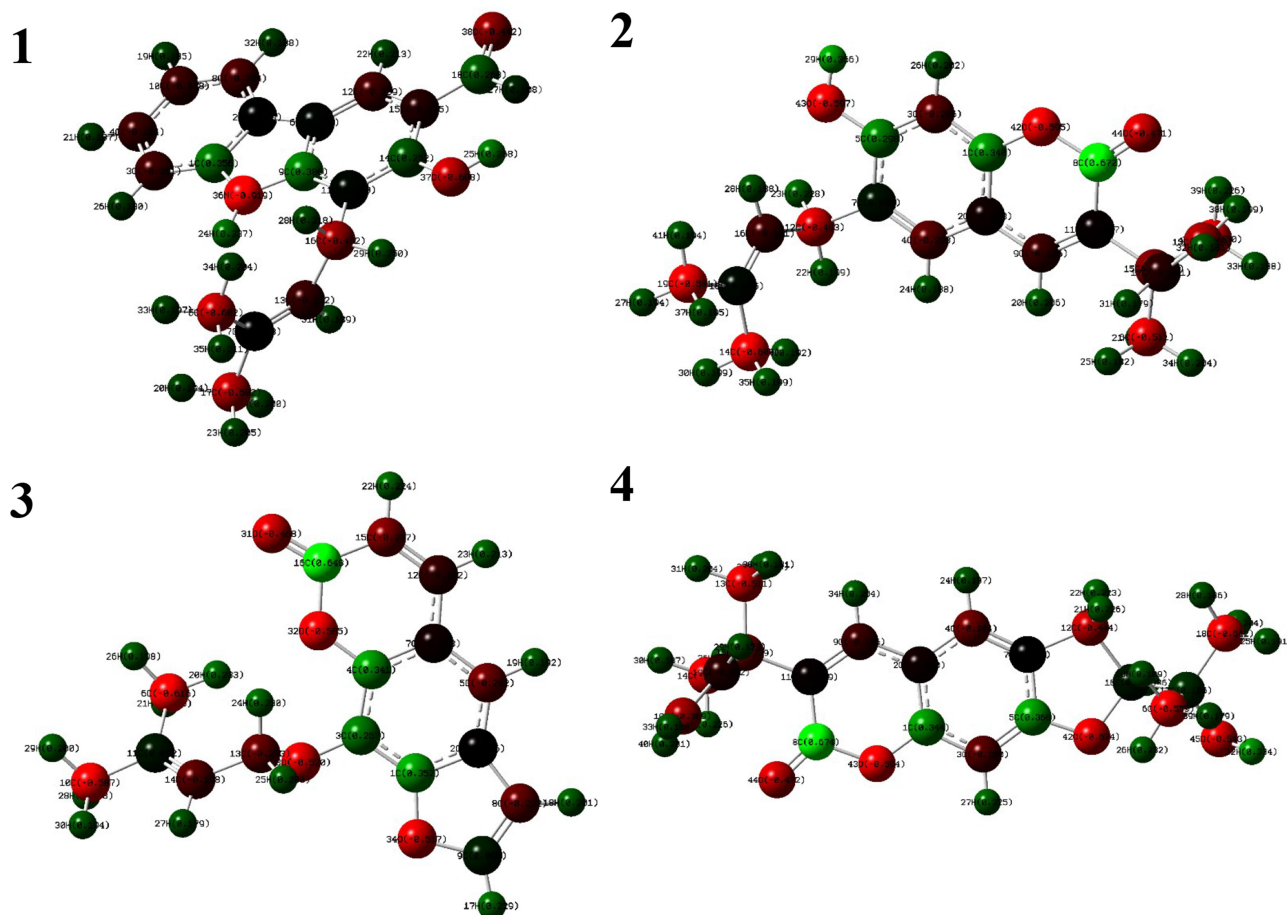
**Figure 4** The optimized structures of compounds (1–4).

Table 6 The Various Quantum Chemical Parameters of Isolated Compounds (1–4)

S.No.	Compounds	E_{HOMO} (eV)	E_{LUMO} (eV)	Energy Gap ΔE (eV)	Electronegativity χ (eV)	Global Hardness η (eV)	Global Softness σ (eV^{-1})	Global Electrophilicity ω (eV)
1	$C_{18}H_{17}N_1O_2$	-0.2064	-0.0490	0.15746	0.12773	0.07873	12.70164	0.103613
2	$C_{19}H_{22}O_3$	-0.2154	-0.0564	0.15907	0.13592	0.07953	12.57308	0.116148
3	$C_{16}H_{14}O_4$	-0.2209	-0.0692	0.15168	0.14506	0.07584	13.18565	0.138729
4	$C_{19}H_{22}O_4$	-0.2110	-0.0546	0.15644	0.13283	0.07822	12.78445	0.112783

HOMO, LUMO, and energy gap (ΔE) of the isolated compounds are calculated as -0.2064 , -0.0490 , and 0.15746 eV for **1**, -0.2154 , -0.0564 and 0.15907 eV for **2**, -0.2209 , -0.0692 , and 0.15168 eV for **3**, and -0.2110 , -0.0546 , and 0.15644 eV for **4** respectively (Figure 6). The results revealed that compound **3** showed the least energy gap (ΔE) compared with the other isolated compounds suggesting high chemical reactivity and

considerable intramolecular charge transfer from electron donor (HOMO) to electron acceptor (LUMO) groups.³⁴ Based on this results compound **3** has better bioactivity compared with other reported compounds herein. Additionally, compound **3** has large electronegativity (χ eV), global softness (σ eV^{-1}), and global electrophilicity (ω eV) compared with other compounds (Table 6).²⁸

**Figure 5** The DFT calculated Mulliken's atomic charges of compounds (1–4).

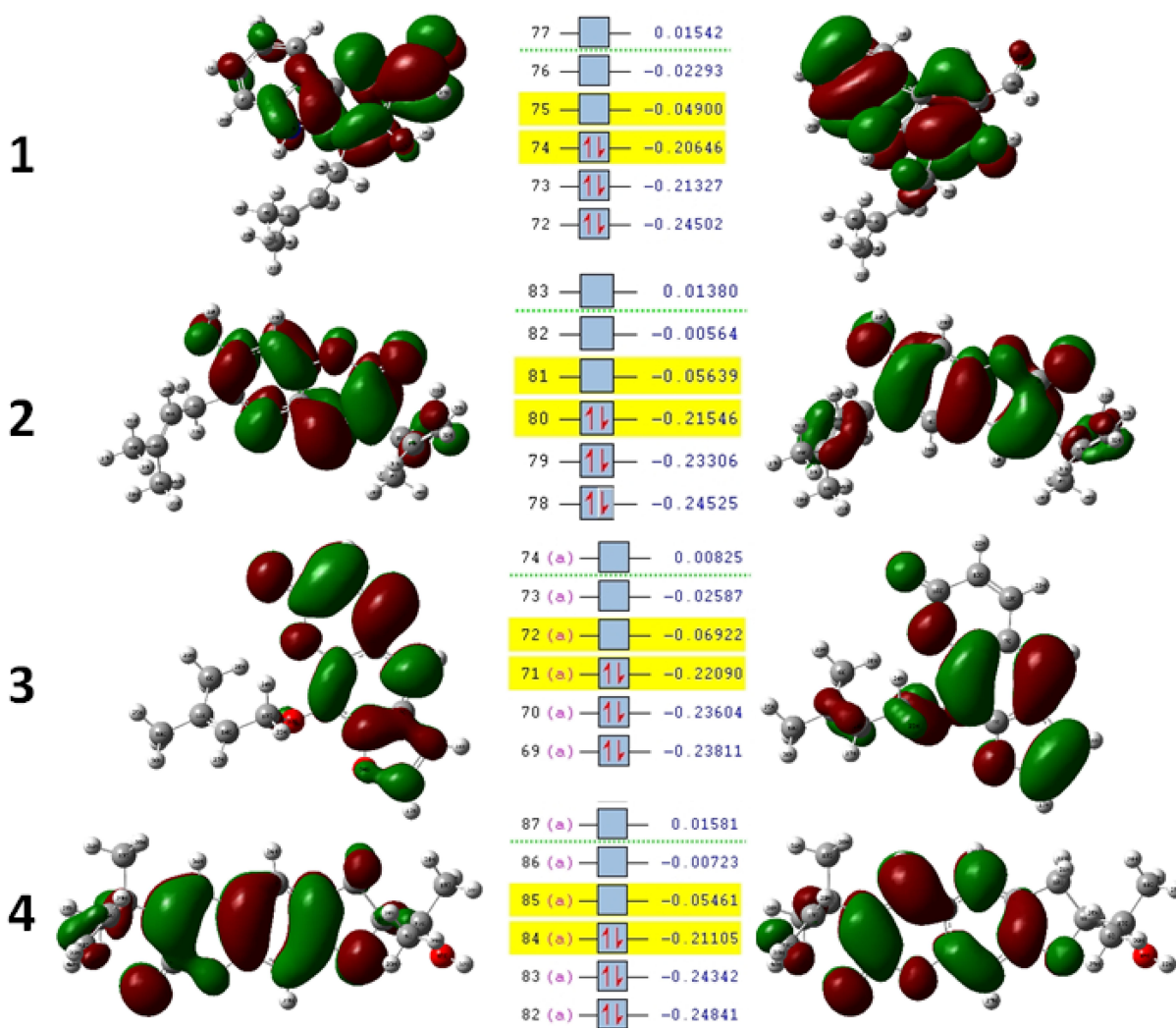


Figure 6 Molecular orbitals and energies for the HOMO and LUMO of isolated compounds (1–4).

Conclusion

In the present work, a computational de novo approach was used to confirm mode of binding for antibacterial activity, elucidating quantum chemical properties and the ADMET-drug-likeness of a carbazole alkaloid (1) and three coumarins (2–4) isolated from roots of *Clausena anisata*. Compared with alkaloid (1), coumarins (2–4) displayed better docking score against both proteins and better docking scores than ciprofloxacin suggesting further work for functional group inclusion and modification to develop novel antibacterial agents with therapeutic activity. SwissADME results revealed that all the compounds are inhibitors for CYP1A2, CYP2C19, and CYP2C9. Compounds 1 and 4 were found to be potential inhibitors for CYP2D6 whereas compounds 2, 3, and ciprofloxacin were found to be non-inhibitors. For

CYP3A4, compound 1 and ciprofloxacin were found to be non-inhibitors whereas compounds 2–4 were found to be potential inhibitors. Toxicological prediction results suggested that all compounds and ciprofloxacin are non-hepatotoxic, non-carcinogenic, non-irritant, immunogenic, and non-cytotoxic with the exception of compound 3 that displayed a mutagenicity value closer to ciprofloxacin (with LD₅₀ value of 480 mg/kg). The DFT results revealed that compound 3 has better bioactivity and chemical reactivity with considerable intramolecular charge transfer between electron-donor and electron-acceptor groups. Based on the results of the present investigation, compound 3 may serve as a lead molecule and further work is recommended for functional group inclusion, modification, and SAR study to develop novel antibacterial agents with therapeutic activity

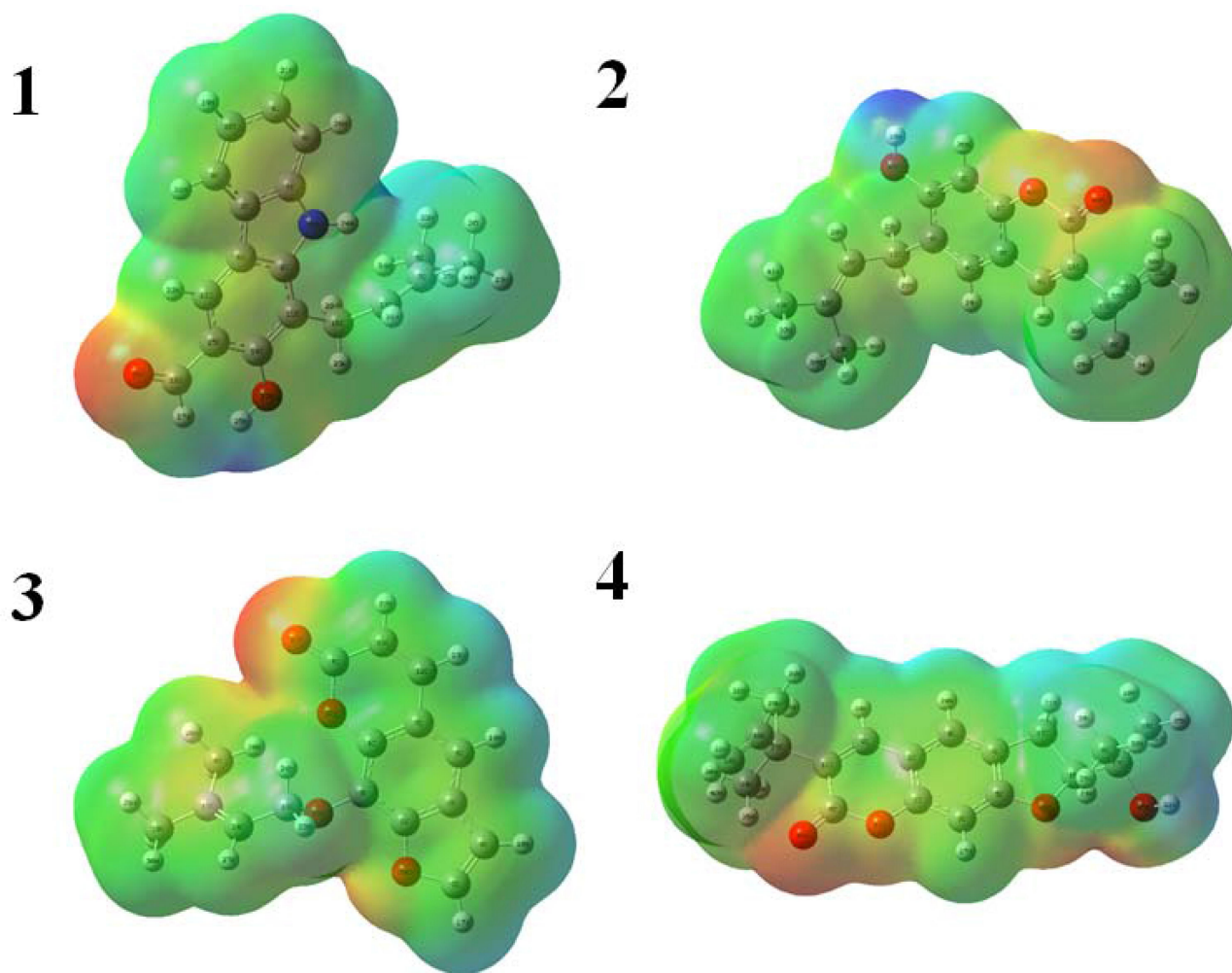


Figure 7 Molecular electrostatic potential surface of compounds (1–4).

against *E. coli* DNA gyrase B inhibitor and efficient inhibitors for quorum sensing auto-inducer LasR binding domain of *Pseudomonas aeruginosa*.

Supporting Information

Full data for Autodock Vina conformations of compounds 1–4 against LasR and DNA gyrase binding domain are included within [supplementary materials](#).

Data Sharing Statement

AutoDock Vina conformations for compounds 1–4 against LasR and DNA gyrase B binding domain can be freely accessed from [supplementary materials](#).

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Disclosure

The authors declare no conflicts of interest.

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