


Cite this: *RSC Adv.*, 2023, 13, 18306

Exploring the inhibitory potential of *Nigella sativa* against dengue virus NS2B/NS3 protease and NS5 polymerase using computational approaches†

Mamuna Mukhtar, ^a Haris Ahmed Khan ^{ab} and Najam us Sahar Sadaf Zaidi ^{*a}

Dengue fever, a highly infectious and rapidly spreading vector borne illness, is classified as a Neglected Tropical Disease (NTD) by WHO because they generally afflict the world's poor and historically have not received as much attention as other diseases. DENV NS2B/NS3 protease and NS5 polymerase are regarded as significant prospective therapeutic targets because of their critical involvement in the viral replication cycle. To date, no specific antiviral agents exist for dengue. The commonly used herbal plant *Nigella sativa* is known for its antibacterial, antiviral, anti-inflammatory, wound-healing, and dermatological properties. Nevertheless, not enough studies on the antiviral effects of *Nigella sativa* against DENV are reported. The current study used several prediction techniques to anticipate the oral bioavailability of substances, druglikeness, and non-toxic and non-mutagenic effects which could lead to the development of novel, safer medications. Therefore, the current study was conducted to explore the inhibitory potential of 18 phytochemicals from *Nigella sativa* against two important enzymes of dengue virus *i.e.*, NS2B/NS3 and NS5. Promising results have been observed for NS2B/NS3 with Taraxerol ($-9.1 \text{ kcal mol}^{-1}$), isoquercetin ($8.4 \text{ kcal mol}^{-1}$), apigenin, and stigmasterol ($-8.3 \text{ kcal mol}^{-1}$). Similarly, NS5 has shown favorable outcomes with apigenin ($-9.9 \text{ kcal mol}^{-1}$), rutin ($-9.3 \text{ kcal mol}^{-1}$), nigellicine ($-9.1 \text{ kcal mol}^{-1}$), and stigmasterol ($-8.8 \text{ kcal mol}^{-1}$). MD simulations validated the structural flexibility of the NS2B/NS3-taraxerol and NS5-apigenin docking complexes based on an RMSF value below 5 Å. The study concluded that among the understudied phytocomponents of *N. sativa*, apigenin, nigellicine, nigellicidine, dithymoquinone, taraxerol, campesterol, cycloeucalenol, stigmasterol and beta-sitosterol have been revealed as potential drug candidates, expected to show antiviral activity and promising drug likelihood. Phytochemicals on the short list may serve as inspiration for the creation of new drugs in the future. Further *in vitro* examination will assist in elucidating the molecular complexity of therapeutic and antiviral capabilities, opening several opportunities for researchers to identify novel medications throughout the drug development process.

Received 19th April 2023
Accepted 29th May 2023

DOI: 10.1039/d3ra02613b

rsc.li/rsc-advances

Introduction

Dengue fever, a highly infectious viral and fast spreading vector-borne disease is responsible for more than 50 million dengue infections every year, and approximately 250 000 people lose their lives to this terrible disease.^{1,2} The WHO classified dengue as a Neglected Tropical Disease (NTD), indicating that the disease is still largely uncontrolled globally, as these regions are likely to have more dengue cases.³ Dengue virus (DENV) belongs to the Flavivirus genus of family *Flaviviridae* and is mostly transmitted by *Aedes* mosquitoes, particularly *Aedes aegypti*. The

four serotypes of dengue are responsible for an extensive spectrum of clinical signs and symptoms, from mild febrile illness to severe and fatal diseases like dengue shock syndrome (DSS) and dengue haemorrhagic fever (DHF).⁴⁻⁶ There are various challenges in creating effective medications against dengue fever, despite the large number of recent research efforts. Due to the fact that DENV exhibits antibody-dependent enhancement (ADE), which allows pre-existing antibodies to attach to DENV and increase the risk of dengue infection.⁷ Numerous vaccine candidates are undergoing preliminary clinical and preclinical research. The only approved vaccination at the moment is CYD-TDV (Dengvaxia®), however WHO advises using it for those between the ages of 9 and 45.⁸ Moreover, a re-infection with the same DENV serotype can occur after immunization due to genetic and antigenic variations.^{9,10}

DENV is a positive sense single-stranded RNA (+ssRNA) virus with an overall genome size of approximately 10.7 kb (*i.e.*, 3391 amino acids) encoding a precursor polyprotein.¹¹ The precursor

^aAtta ur Rahman School of Applied Biosciences (ASAB), National University of Sciences and Technology (NUST), H-12, 44000, Islamabad, Pakistan. E-mail: sadafzaidi@asab.nust.edu.pk

^bDepartment of Biotechnology, University of Mianwali, 42200, Punjab, Pakistan

† Electronic supplementary information (ESI) available. See DOI: <https://doi.org/10.1039/d3ra02613b>



polyprotein (5'-C-prME-NS1-NS2A-NS2B-NS3-NS4A-NS4B-NS5-3') is further cleaved to produce three structural proteins (capsid protein C, membrane protein prM and envelope protein E) and seven non-structural proteins (NS1, NS2a, NS2b, NS3, NS4a, NS4b and NS5).^{3,12} The genome of flaviviruses *e.g.*, Dengue virus replicates by the RNA-dependent RNA polymerase (RdRp) domain of non-structural protein 5 (NS5). It also encodes a unique two-component NS2B/NS3 protease for its maturation and infectivity.^{13,14} Among non-structural proteins, NS3 and NS5 are known to perform a variety of enzymatic reactions. The NS5 protein is highly conserved between the four virus serotypes and is essential for virus replication and transcription. Since there is no enzyme with RdRp activity in humans, NS5 has been reported as an appropriate anti-dengue drug target.^{15,16} NS3 encodes the RNA helicase serine protease that cleaves the viral polypeptide into functional proteins, nucleocapsid triphosphatase (NTPase), and RNA triphosphatase (RTPase) activities. NS3 protease has apo behaviour and requires a small 14 kDa protein (*i.e.*, NS2B protein), as a cofactor. The NS2B protein stabilizes the NS2B/NS3 complex and NS2B and NS3 perform proteolytic cleavage in the virus.^{17,18} Thus, these essential enzymatic activities of NS2B/NS3 and NS5 make them an attractive target for anti-flaviviral drugs.

Along with the creation of several vaccine candidates, research is being done on antivirals that can stop the spread of viruses or stop illness. The first antiviral substance, acyclovir, entered clinical use in 1960, marking the field of antiviral medications' maturation. Gertrude B. Elion received the Nobel Prize in 1988 for the scientific discovery she made in 1959. Over time, many plants have been used to make rudimentary medicines that have been used to cure a variety of human ailments and illnesses.¹⁹ The antiviral activity of phytochemicals has been studied for burgeoning viral infections, and these natural antiviral compounds are the best replacements for antiviral medications or drugs due to their lower toxicity.^{20,21} *Nigella sativa* (*N. sativa*), a member of the family Ranunculaceae, is an annual flowering plant that grows in Mediterranean and Asian countries including Pakistan, India, Indonesia, Afghanistan, and Italy. It is commonly known as black seed, black cumin. In Pakistan it is called Kalonji while in China it is referred to as Hak Jung Chou.^{22,23} *N. sativa* has been used in the folk medicine systems such as Unani and Tibb, Ayurveda and Siddha for the treatment of different ailments.²⁴ It is one of the most used herbal medicines throughout the world and pharmacological studies have confirmed the potential of *N. sativa* as an antidiabetic, anticancer, antioxidant, antimicrobial, hepatoprotective, neuroprotective, gastroprotective, spasmolytic, immunomodulator, analgesic, anti-inflammatory, and bronchodilator (Ahmad *et al.* 2021 (ref. 52)). Various chemical constituents of *N. sativa* have been identified, including dithymoquinone, -pinene, -pinene, thymoquinone, thymohydroquinone, thymol, nigellidine, carvacrol, *p*-cymene, nigellidine, nigellimine, nigellidine, alpha-hederin, and limonene. (Thakur *et al.* 2021;⁵⁰ Kabir *et al.*⁷⁶ 2020).

Despite its clear severity and enormous morbidity, there is currently no direct-acting or host-directed antiviral therapy licensed to treat these illnesses, and the only approved vaccine

used in certain countries is restricted to seropositive patients.^{25,26} A lot of attention has been paid to medicinal plants as a natural source of antiviral medications due to the poor clinical results and side effects of the dengue immunization program. For the treatment of viral infections, medicinal plants have long been thought of as the primary source of pharmacologically bioactive compounds. Additionally, the difficulties brought on by frequent virus mutations and the development of viral resistance to traditional antiviral therapies are igniting interest in natural substances as antiviral alternatives.^{27,28} The current study was conducted with the purpose of detecting the inhibitory potential of eighteen phytochemicals from *Nigella sativa* against two important enzymes of the dengue virus (*i.e.*, NS2B/NS3 protease and NS5 polymerase). Molecular docking and MD simulation studies were conducted for ligand-protein complexes to determine the stability of docked structures and several pharmacokinetic analyses evaluated the drug-likeness and toxicity potential of the phytoligands for the evaluation of suitable leads to develop an anti-dengue drug.

Materials and methods

Ligand selection

For the present study, eighteen (18) bioactive components from seed oil and a seed source of black cumin (*Nigella sativa*) were selected from the literature. The selected phytochemicals have previously been studied for various having inhibitory potential against infectious diseases.^{23,29–31} 3D structures of the selected phytochemicals were downloaded from PubChem in SDF formats and Biovia Discovery studio v2021 was used to convert them into PDB format. Ligands were prepared for docking by fixing charges and adding polar hydrogen before being saved in PDBQT format. The physicochemical properties and molecular formulas and their 3D structures of the selected phytochemicals are given in Table 1.

Receptor preparation

The 3D crystal structures of two dengue virus proteins, *i.e.*, NS2B/NS3 protease (PDB ID: 2FOM) and NS5 polymerase (PDB ID: 4V0Q), available in the RCSB database, were downloaded from Protein Data Bank in PDB format. The downloaded files were opened in Discovery Studio to remove the original ligands and other small molecules attached to the protein structures. The dengue virus NS2BNS3 was originally bound with glycerol (GOL) whereas NS5 polymerase was complexed with *S*-adenosyl-L-homocysteine (SAH). The original ligand in NS5 and Curcumin, known to possess inhibitory potential against NS2BNS3 were used to serve as positive controls and their binding energies and interactions have been recorded for comparison. Both 3D protein structures (Fig. 1) were refined before docking analyses were performed. The refinement of the selected proteins was carried out by removing irrelevant ions, ligands (if any), and water molecules. Furthermore, polar hydrogen atoms and Kollman charges were added to the receptors and then saved in PDBQT format for docking.



Table 1 Molecular weights, molecular formulas and 3D structures of the selected phytoligands

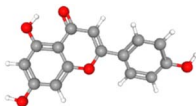
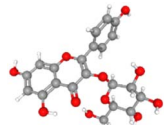

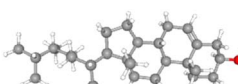
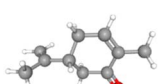

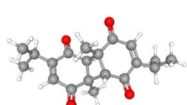
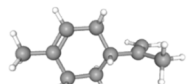
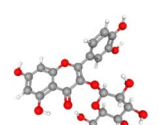
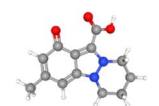
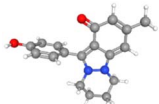
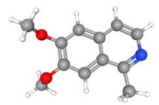
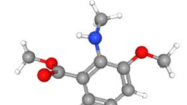
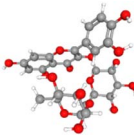
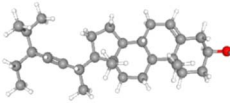
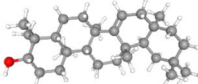
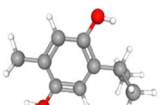
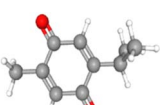
Ligands	PubChem CID	Molecular formula	Molecular weight	Structure
Apigenin	CID_5280443	C ₁₅ H ₁₀ O ₅	270.24 g mol ⁻¹	
Astragalin	CID_5282102	C ₂₁ H ₂₀ O ₁₁	448.4 g mol ⁻¹	
β-Sitosterol	CID_222284	C ₂₉ H ₅₀ O	414.7 g mol ⁻¹	
Campesterol	CID_173183	C ₂₈ H ₄₈ O	400.7 g mol ⁻¹	
Carvone	CID_7439	C ₁₀ H ₁₄ O	150.22 g mol ⁻¹	
Cycloeucalenol	CID_101690	C ₃₀ H ₅₀ O	426.7 g mol ⁻¹	
Dithymoquinone	CID_398941	C ₂₀ H ₂₄ O ₄	328.4 g mol ⁻¹	
D-Limonene	CID_440917	C ₁₀ H ₁₆	136.23 g mol ⁻¹	
Isoquercetin	CID_5280804	C ₂₁ H ₂₀ O ₁₂	464.4 g mol ⁻¹	
Nigellicine	CID_11402337	C ₁₃ H ₁₄ N ₂ O ₃	246.26 g mol ⁻¹	
Nigellidine	CID_136828302	C ₁₈ H ₁₈ N ₂ O ₂	294.3g mol ⁻¹	
Nigellimine	CID_20725	C ₁₂ H ₁₃ NO ₂	203.24g mol ⁻¹	
Nigelline	CID_21368	C ₁₀ H ₁₃ NO ₃	195.21 g mol ⁻¹	



Table 1 (Contd.)

Ligands	PubChem CID	Molecular formula	Molecular weight	Structure
Rutin	CID_5280805	C ₂₇ H ₃₀ O ₁₆	610.5 g mol ⁻¹	
Stigmasterol	CID_5280794	C ₂₉ H ₄₈ O	412.7 g mol ⁻¹	
Taraxerol	CID_92097	C ₃₀ H ₅₀ O	426.7 g mol ⁻¹	
Thymohydroquinone	CID_95779	C ₁₀ H ₁₄ O ₂	166.22 g mol ⁻¹	
Thymoquinone	CID_10281	C ₁₀ H ₁₂ O ₂	164.2 g mol ⁻¹	

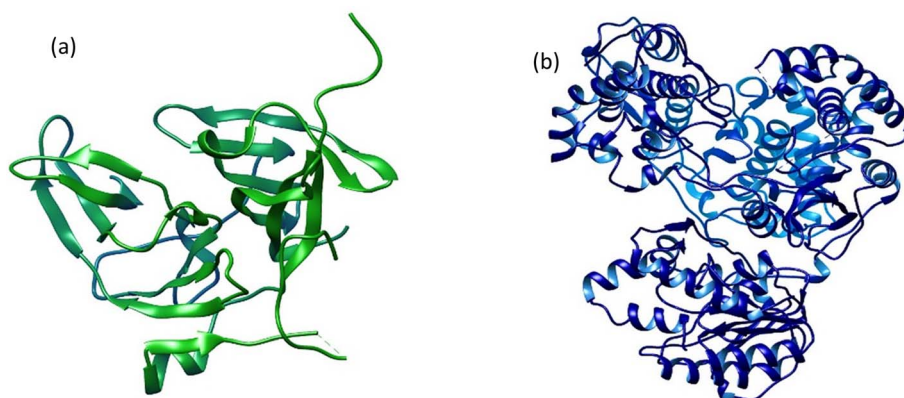


Fig. 1 The three-dimensional crystal structures of two dengue virus proteins: (a) NS2B/NS3 protease and (b) NS5 polymerase.

Prediction of activity spectra for substances (PASS) analysis

Prediction of Activity the Spectra for Substances (PASS) analysis program predicts the biological properties, pharmacological properties, drug-likeness, probable side effects, and mode of action of understudied phytoconstituents based on their structure–activity relationship with an established chemical entity. PASS analysis was carried out in this study through a variety of online and offline tools, mentioned below^{29,32}

Lipinski's rule of five

Lipinski's rule of five describes the molecular properties of drug compounds based on important pharmacokinetic parameters

such as absorption, metabolism, distribution, and excretion. The rule is helpful in the design and development of drugs.³³ The drug-likeness of the 18 phytochemicals of *N. sativa* included in the study was expected using the Lipinski rule of five. The online tool Molinspiration v2021.03 ("<https://www.molinspiration.com/cgi-bin/properties>") was used to calculate the drug-likeness of phytochemicals and compared them with standard drugs. The drug-likeness was calculated using parameters such as the logarithm of the compound partition coefficient between *n*-octanol and water ($\log P \leq 5$), molecular weight (MW ≤ 500), number of hydrogen bond donors ($\text{NOHNH} \leq 5$), hydrogen bond acceptor sites ($\text{NON} \leq 10$),



topological polar surface area (defined as a sum of surfaces of polar atoms in a molecule, $TPSA \leq 140 \text{ \AA}^2$), number of rotatable bonds (≤ 10). An orally active drug should not have more than one Lipinski violation to maintain its bioavailability.³⁴

Bioactivity score prediction

Drug score values indicate the absolute potential of a prospective complex to be a drug candidate. A web-based tool, Molinspiration version 2021.03, predicted a bioactivity score of the phytoconstituents against the human receptors like G protein-coupled receptors (GPCRs), kinases, proteases, ion channels, enzymes, and nuclear receptors. If the bioactivity score is more than 0.0, then the compound is dynamic (active); if it is in the range between -5.0 and 0.0 , then the complex is moderately active and if the bioactivity score is less than -5.0 , then it is considered to be idle (inactive).³⁵

Toxicity potential study

Toxicity risk assessment gives exploratory information on the potential side effects of phytoconstituents that can be employed in drug discovery and development. The Protox-II server and OSIRIS Data Warrior V5.2.1 software were used to observe drug-likeness and drug toxicity risk characteristics such as drug-likeness, tumorigenicity, carcinogenicity, immunotoxicity, hepatotoxicity, mutagenicity, reproductive and irritating effects.^{29,36} The understudy compounds have also been evaluated for prediction of their LD_{50} value and Drug toxicity class. The LD_{50} values for toxic dosages are frequently expressed in mg kg^{-1} body weight. The median lethal dosage, or LD_{50} , is the quantity at which 50% of test subjects pass away after being exposed to a substance. According to the Globally Harmonized System (GHS) of categorization and labeling of substances, toxicity classes are established.^{37,38}

Pharmacokinetic property prediction

The absorption, distribution, metabolism, excretion, and toxicity (ADMET) properties of all phytoconstituents in this study were predicted using two online software, *i.e.*, the Swiss-ADME software and ADMETlab. These programs examine a compound's important pharmacokinetic properties such as blood-brain barrier (BBB), distribution, GI absorption, metabolism as a P-glycoprotein (P-gp) substrate, cytochrome P450 s such as CYP1A2, CYP2C19, CYP2C9, CYP2D6, CYP3A4 inhibitor, and lipophilicity for plasma membrane absorption.^{39,40}

Docking

AutoDock Tools 1.5.6 was used to integrate phytoligands, which were considered "flexible," into the protein targets that were considered "rigid." AutoDock Vina is easy to use, has a fast processing speed, calculates the dimensions of grid boxes automatically, and conveniently estimates binding sites.^{41,42} In the present study, a grid box size of $40 \times 40 \times 40$ (x , y , and z) and a grid spacing of 0.375 were used for both receptor proteins. The center of the grid for NS2B/NS3 protease was set at positions 0.456 , -17.036 , and 13.989 for x , y , and z ,

respectively, while for NS5 polymerase the centers (x , y , and z) were set at 24.892 , 162.151 , and 24.605 , respectively. For each ligand, a configuration file containing grid box attributes was created and saved in .txt format. To begin the docking analysis, a set of codes (commands) was written in the command prompt to generate the output score.⁴³ The results displayed the values of binding energy or the Gibbs free energy (G) in kcal mol^{-1} , and the ligand with the maximum negative value was considered to have the highest binding affinity for a particular target protein. The results displayed the values of binding affinity and Gibbs free energy (G) in kcal mol^{-1} , and the ligand with the maximum negative value was considered to have the highest binding affinity for a particular target protein. From the nine different conformations generated, the best-fit model (Model 1) was selected. The interaction profiles were displayed in 2D using the Discovery Studio v2021 and then evaluated.

Molecular dynamic simulations

The investigation of the structural dynamics of the top docking complexes and the identification of motion were both carried out using iMODS. To determine deformability, B -factor, eigenvalue, variance, covariance map, and elastic network, normal mode analysis was performed on the docked complexes. The input PDB files were uploaded to iMODS with the default settings of 300 K constant temperature, 1 atm constant pressure, and 50 ns molecular dynamic simulation for each complex.^{44,45} The best docked structures were validated for their protein flexibility by performing molecular dynamic simulations using the CABS-flex 2.0 server and displayed as RMSF values (root mean square fluctuation). Simulations in CABS-flex were conducted using default parameters, *i.e.*, protein rigidity: 1.0 , number of cycles: 50 , number of cycles between trajectories: 50 , temperature range: 1.40 , and a random number of generator seed of 5711 . CABS-flex offers fast, high-resolution (10 ns) protein flexibility simulation at extremely reduced system constraints.^{46,47}

Results

Lipinski's rule of five

The physicochemical properties of eighteen phytoconstituents of *N. sativa* were assessed by PASS analysis by employing Lipinski's rule of five (Table 2). Three phytoconstituents (*i.e.*, astragaloside, isoquercetin and rutin) exhibited more than 1 Lipinski violation as they had more than five hydrogen bond donors (NOHNH) and more than ten hydrogen bond acceptors (NON) and their topological polar surface area (TPSA) was calculated to be greater than 140 \AA^2 . Five compounds (*i.e.*, β -sitosterol, campesterol, cycloeucalenol, stigmasterol and taraxerol) exhibited 1 violation as the value of $\text{clog}P$ for these compounds was calculated to be greater than the standard value (*i.e.*, ≤ 5). The remaining ten phytocompounds exhibited no violation. Since an ideal lead compound should not have more than one violation, fifteen compounds out of eighteen have met the criteria.



Table 2 PASS analysis (Lipinski's rule of five) of active phytoconstituents from *N. sativa*

Phytoconstituents	clog <i>P</i> (≤ 5)	MW (≤ 500)	NOHNH (≤ 5)	NON (≤ 10)	TPSA ($\leq 140 \text{ \AA}^2$)	Number of rotatable bonds (≤ 10)	Lipinski's violations
Apigenin	2.46	270.24	3	5	90.89	1	0
Astragalin	0.125	448.38	7	11	190.275	4	2
β -Sitosterol	8.62	414.718	1	1	20.228	6	1
Campesterol	8.305	400.691	1	1	20.228	5	1
Carvone	2.513	150.221	0	1	17.071	1	0
Cycloeucalenol	7.617	426.729	1	1	20.228	5	1
Dithymoquinone	1.703	328.408	0	4	68.284	2	0
D-Limonene	3.615	136.238	0	0	0	1	0
Isoquercetin	−0.364	464.379	8	12	210.503	4	2
Nigellicine	1.652	246.266	1	5	64.238	1	0
Nigellidine	3.034	294.354	1	4	47.167	1	0
Nigellimine	1.894	203.241	0	3	31.36	2	0
Nigelline	2.107	195.218	1	4	47.566	4	0
Rutin	−1.063	610.521	10	16	269.427	6	3
Stigmasterol	7.869	412.702	1	1	20.228	5	1
Taraxerol	8.023	426.729	1	1	1	1	1
Thymohydroquinone	3.258	166.22	2	2	40.456	1	0
Thymoquinone	1.90	164.20	0	2	34.14	1	0

Prediction of bioactivity score

The prediction of the bioactivity score revealed that Apigenin was biologically active as nuclear receptor ligand, kinase and enzyme inhibitor while remaining moderately active as GPCR ligand and ion channel modulator. β -Sitosterol, campesterol, taraxerol and cycloeucalenol have demonstrated significant activity as GPCR ligand, kinase inhibitor, ion channel modulator, nuclear receptor ligand, protease inhibitors and enzyme inhibitors as all their values were above the threshold level as described in materials and methods. Nigellidine was active as GPCR ligands, ion channel modulators, nuclear receptor

ligands, and enzyme inhibitors while being moderately active as protease inhibitors. Dithymoquinone and stigmasterol were moderately active as ion channel modulator, kinase inhibitor and protease inhibitor and active as GPCR ligand, nuclear receptor ligand and enzyme inhibitor. Cycloeucalenol exhibited the highest value as protease inhibitor (*i.e.*, 0.1) and as enzyme inhibitor (*i.e.*, 0.6). Thymohydroquinone and thymoquinone were considered inactive because their maximum values were below the threshold level. The details of the bioactivity score prediction of all understudy phytoconstituents are given in Table 3.

Table 3 Bioactivity score prediction of active constituents of *N. sativa*^a

Phytoconstituents	GPCR ligand	Ion channel modulator	Kinase inhibitor	Nuclear receptor ligand	Protease inhibitor	Enzyme inhibitor
Apigenin	−0.07	−0.09	0.18	0.34	−0.25	0.26
Astragalin	0.06	−0.05	0.10	0.20	−0.05	0.41
β -Sitosterol	0.14	0.04	−0.51	0.73	0.07	0.51
Campesterol	0.11	0.01	−0.48	0.71	0.01	0.50
Carvone	−1.23	−0.30	−2.51	−0.54	−1.21	−0.45
Cycloeucalenol	0.14	0.14	−0.37	0.92	0.10	0.61
Dithymoquinone	0.18	−0.09	−0.48	0.14	−0.10	0.10
D-Limonene	−0.91	−0.27	−2.01	−0.34	−1.38	−0.21
Isoquercetin	0.06	−0.04	0.13	0.20	−0.06	0.42
Nigellicine	−0.15	0.00	−0.18	−0.29	−0.57	0.20
Nigellidine	0.07	0.01	0.29	0.10	−0.32	0.19
Nigellimine	0.47	−0.17	−0.40	−0.75	−0.64	−0.12
Nigelline	−0.77	−0.46	−0.77	−1.01	−0.98	−0.49
Rutin	0.05	−0.52	−0.14	−0.23	−0.07	0.12
Stigmasterol	0.12	−0.08	−0.48	0.74	−0.02	0.53
Taraxerol	0.21	0.02	−0.20	0.54	0.00	0.49
Thymohydroquinone	−0.92	−0.44	−1.06	−0.54	−1.17	−0.46
Thymoquinone	−1.40	−0.31	−1.27	−1.47	−1.45	−0.40

^a >0.00 active, −0.50 to 0.00 moderately active, <−0.50 inactive.

Pharmacokinetic property prediction

The ADMET analysis (Table 4) of understudied phytoconstituents suggested that apigenin, astragalín, isoquercetin, and rutin were unable to cross the blood–brain barrier, while all of the remaining selected phytoligands showed positive results, demonstrating that they can easily cross the BBB. Only β -sitosterol, stigmasterol, and nigellidine were found to be positive as permeability glycoprotein substrates (P-gp substrates), while the




rest of the phytoconstituents were found to be negative. These results suggest that being non-Pgp substrates, the phytoconstituents will persist longer in the cells, leading to their increased pharmacokinetic efficacy. For incessant plasma concentrations and enhanced bioavailability, the understudy compounds were expected to inhibit the five classes of cytochromes P450, *i.e.*, CYP1A2, CYP2C19, CYP2C9, CYP2D6, and CYP3A4. The results obtained confirmed CYP1A2 inhibition by apigenin, nigellidine,

Table 4 ADMET properties of phytoconstituents of *N. sativa*

Phytoconstituents	CYP1A2 inhibition	CYP2C19 inhibition	CYP2C9 inhibition	CYP2D6 inhibition	CYP3A4 inhibition	P-gp substrate	log <i>P</i> o/ BBB w	GI absorption
Apigenin	Yes	No	No	Yes	Yes	No	No 2.11	High
Astragalín	No	No	No	No	No	No	No −0.09	Low
β -Sitosterol	No	No	No	No	No	Yes	Yes 8.025	Low
Campesterol	No	No	No	No	No	No	Yes 2.385	Low
Carvone	No	No	No	No	No	No	Yes 2.44	High
Cycloeucalenol	No	No	No	No	No	No	Yes 7.651	Low
Dithymoquinone	No	No	No	No	No	No	Yes 2.61	High
D-Limonene	No	No	Yes	No	No	No	Yes 3.37	Low
Isoquercetin	No	No	No	No	No	No	No −0.48	Low
Nigellidine	No	No	No	No	No	No	Yes 1.45	High
Nigellimine	Yes	No	No	Yes	No	Yes	Yes 2.82	High
Nigelline	Yes	No	No	No	No	No	Yes 2.29	High
Rutin	Yes	No	No	No	No	No	Yes 1.79	High
Stigmasterol	No	No	No	No	No	No	No −0.038	Low
Taraxerol	No	No	No	No	No	Yes	Yes 7.748	Low
Thymohydroquinone	No	No	No	No	No	No	Yes 6.782	Low
Thymoquinone	Yes	No	No	No	No	Yes	Yes 2.39	High
	No	No	No	No	No	No	Yes 1.85	High

Table 5 Predicted Toxicity risk assessment of phytoconstituents of *N. sativa*.

Phytoconstituents	Tumorigenicity	Irritant	Reproductive Effective	Mutagenicity	Hepatotoxicity	Immunotoxicity	Cytotoxicity	Carcinogenicity
Apigenin	Non-toxic	Non-toxic	Non-toxic	Non-toxic	Non-toxic	Non-toxic	Non-toxic	Non-toxic
Astragalín	Non-toxic	Non-toxic	Non-toxic	Non-toxic	Non-toxic	Non-toxic	Non-toxic	Non-toxic
β -Sitosterol	Non-toxic	Non-toxic	Non-toxic	Non-toxic	Non-toxic	Highly toxic	Non-toxic	Non-toxic
Campesterol	Non-toxic	Non-toxic	Non-toxic	Non-toxic	Non-toxic	Highly toxic	Non-toxic	Non-toxic
Carvone	Non-toxic	Non-toxic	Slightly toxic	Non-toxic	Non-toxic	Non-toxic	Non-toxic	Non-toxic
Cycloeucalenol	Non-toxic	Non-toxic	Non-toxic	Non-toxic	Non-toxic	Highly toxic	Non-toxic	Non-toxic
Dithymoquinone	Non-toxic	Non-toxic	Slightly toxic	Non-toxic	Non-toxic	Non-toxic	Non-toxic	Non-toxic
D-Limonene	Non-toxic	Non-toxic	Non-toxic	Non-toxic	Non-toxic	Slightly toxic	Non-toxic	Non-toxic
Isoquercetin	Non-toxic	Non-toxic	Non-toxic	Non-toxic	Non-toxic	Slightly toxic	Non-toxic	Non-toxic
Nigellidine	Non-toxic	Non-toxic	Non-toxic	Non-toxic	Non-toxic	Non-toxic	Non-toxic	Slightly toxic
Nigellimine	Non-toxic	Non-toxic	Non-toxic	Highly toxic	Slightly toxic	Slightly toxic	Non-toxic	Non-toxic
Nigelline	Non-toxic	Non-toxic	Non-toxic	Non-toxic	Slightly toxic	Non-toxic	Non-toxic	Non-toxic
Rutin	Non-toxic	Non-toxic	Non-toxic	Non-toxic	Non-toxic	Highly toxic	Non-toxic	Non-toxic
Stigmasterol	Non-toxic	Non-toxic	Non-toxic	Non-toxic	Non-toxic	Highly toxic	Non-toxic	Non-toxic
Taraxerol	Non-toxic	Non-toxic	Non-toxic	Non-toxic	Non-toxic	Highly toxic	Non-toxic	Non-toxic
Thymohydroquinone	Slightly toxic	Non-toxic	Non-toxic	Non-toxic	Non-toxic	Non-toxic	Non-toxic	Non-toxic
Thymoquinone	Non-toxic	Non-toxic	Non-toxic	Non-toxic	Non-toxic	Non-toxic	Non-toxic	Non-toxic

 Highly toxic
 Slightly toxic
 Non-toxic



nigellimine, nigelline, and thymohydroquinone, the inhibition of CYP2D6 by apigenin, and the inhibition of nigellidine and CYP3A4 by apigenin alone. The rest of the compounds did not show inhibitory potential against cytochrome P450. The consensus lipophilicity ($\log P_{\text{o/w}}$) value was calculated to analyze the lipophilic potential of the phytoconstituents, resulting in the lipid-soluble nature of most of the compounds except astragalín,

isoquercetin, and rutin. High human intestinal absorption (GI absorption) was observed for apigenin, carvone, dithymoquinone, nigellicine, nigellidine, nigellimine, nigelline, thymohydroquinone, and thymoquinone.

Toxicity potential study

Assessment of expected drug-likeness and toxicity potential of selected phytochemicals by data warrior and protox-II server revealed that apigenin, astragalín, and thymoquinone were non-toxic. β -sitosterol, campesterol and cycloeucalenol, rutin, stigmasterol and taraxerol were appeared to have projected immunotoxin effects. Carvone and dithymoquinone were expected to have mild toxic effects on the reproductive system, while nigellicine appeared to be slightly carcinogenic. Nigelline and nigellimine have shown potential for causing hepatotoxicity whereas nigellimine is also positive for mild immunotoxicity and high mutagenicity. None of the phytoconstituents were found to be an irritant. The expected toxicity risk assessment of phytoconstituents of *N. sativa* is presented as in Table 5. For a compound to become a good drug candidate, its druglikeness score should be 1 or close to 1. If the drug score is zero or negative, then the understudy compound is not a good drug candidate. Nigellicine, nigellidine, apigenin, rutin, β -limonene, and stigmasterol have shown positive scores for drug-likeness. Computational calculations of predicted LD_{50} and toxicity class renders apigenin, astragalín, cycloeucalenol, dithymoquinone, β -limonene, isoquercetin and rutin to class-5 while β -sitosterol, campesterol, carvone, nigellicine and nigellidine, nigelline and stigmasterol to class-4. Nigellimine (pink highlight) was the only compound in class-III (*i.e.*, toxic if swallowed) whereas taraxerol qualified to be sole potentially non-toxic lead compound having the highest LD_{50} value *i.e.*, 7000 mg kg^{-1}

Table 6 Predicted druglikeness and toxicity potential assessment of phytoconstituents of *N. sativa*^a

Phytoconstituents	Druglikeness	Predicted LD_{50}	Predicted toxicity class
Apigenin	0.28194	2500 mg kg^{-1}	Class V
Astragalín	−3.6679	5000 mg kg^{-1}	Class V
β -Sitosterol	−4.475	890 mg kg^{-1}	Class IV
Campesterol	−8.1908	890 mg kg^{-1}	Class IV
Carvone	−18.988	1640 mg kg^{-1}	Class IV
Cycloeucalenol	−7.6333	5000 mg kg^{-1}	Class V
Dithymoquinone	−21.855	2300 mg kg^{-1}	Class V
β -Limonene	0.9165	4400 mg kg^{-1}	Class V
Isoquercetin	−3.6679	5000 mg kg^{-1}	Class V
Nigellicine	2.1696	1300 mg kg^{-1}	Class IV
Nigellidine	1.1929	1000 mg kg^{-1}	Class IV
Nigellimine	−1.7979	1300 mg kg^{-1}	Class III
Nigelline	−2.6175	1800 mg kg^{-1}	Class IV
Rutin	1.9337	5000 mg kg^{-1}	Class V
Stigmasterol	1.2217	890 mg kg^{-1}	Class IV
Taraxerol	−2.422	70000 mg kg^{-1}	Class VI
Thymohydroquinone	−2.3359	1000 mg kg^{-1}	Class IV
Thymoquinone	−1.1996	2400 mg kg^{-1}	Class V

^a Class I: fatal if swallowed ($\text{LD}_{50} \leq 5$); Class II: fatal if swallowed ($5 < \text{LD}_{50} \leq 50$); Class III: toxic if swallowed ($50 < \text{LD}_{50} \leq 300$); Class IV: harmful if swallowed ($300 < \text{LD}_{50} \leq 2000$); Class V: may be harmful if swallowed ($2000 < \text{LD}_{50} \leq 5000$); Class VI: non-toxic ($\text{LD}_{50} > 5000$).

Table 7 Docking interactions of selected phytochemicals from *N. sativa* with Dengue virus NS2BNS3 protease

Ligand	Receptor proteins	Binding energies (kcal mol^{-1})	Interacting amino acid residues
Apigenin	NS2BNS3	−8.3	Val154, Ala164, Lys74, Asn152, Leu76, Leu149
Astragalín	NS2BNS3	−8.0	Asn167, Trp89, Gly87, Ala166, Val147, Glu88, Ile165, Asn152, Leu86, Lys74
β -Sitosterol	NS2BNS3	−8.3	His51, Phe130, Leu128
Campesterol	NS2BNS3	−8.1	His51, Leu128, Phe130, Pro132
Carvone	NS2BNS3	−5.6	Tyr150, Gly153, Pro132, Tyr161, Leu128
Cycloeucalenol	NS2BNS3	−8.3	His51, Pro132, Leu128
Dithymoquinone	NS2BNS3	−7.2	His51, Gly153, Pro132, Leu128
β -Limonene	NS2BNS3	−5.5	Tyr161, Leu128
Isoquercetin	NS2BNS3	−8.4	Leu149, Asn152, Ala166, Val147, Gly87, Lys74, Leu85, Leu76, Ile165
Nigellicine	NS2BNS3	−7.4	Leu128, Pro132, Gly151, Tyr161
Nigellidine	NS2BNS3	−7.3	Ala164, Leu76, Lys74, Asn167
Nigellimine	NS2BNS3	−6.2	Leu76, Trp83, Leu85, Val147, Lys74, Asn152
Nigelline	NS2BNS3	−5.5	Lys42, Val59, Val146, Lys145
Rutin	NS2BNS3	−8.2	Leu128, Pro132, Gly151, Arg54, Trp50, Lys73, Asp75
Stigmasterol	NS2BNS3	−8.3	His51, Leu128, Phe130, Pro132
Taraxerol	NS2BNS3	−9.1	Pro132, His51, Leu128, Tyr161
Thymohydroquinone	NS2BNS3	−5.6	Ala164, Val154, Ile123, Lys73
Thymoquinone	NS2BNS3	−5.6	Asn152, Ile123, Ala164, Lys74, Val154
Curcumin (positive control)	NS2BNS3	−7.2	Leu128, Pro132, Tyr161



(green highlight). The details of drug similarity and probable toxicity assessment of phytochemicals are given in Table 6.

Docking analysis

AutoDock Vina v1.5.6 was used to dock the target proteins with eighteen phytochemicals from *Nigella sativa* plants. The phytochemicals showed different interactions at the binding pockets,

displaying several binding affinities. The phytochemicals showed binding energies for NS2BNS3 ranging from $-5.5 \text{ kcal mol}^{-1}$ to $-9.1 \text{ kcal mol}^{-1}$. Taraxerol showed the highest binding affinity with lowest binding energy (*i.e.*, $-9.1 \text{ kcal mol}^{-1}$), followed by Isoquercetin (*i.e.*, $-8.4 \text{ kcal mol}^{-1}$), and the lowest binding affinity was recorded for D-limonene and nigelline (*i.e.*, $-5.5 \text{ kcal mol}^{-1}$). The highest

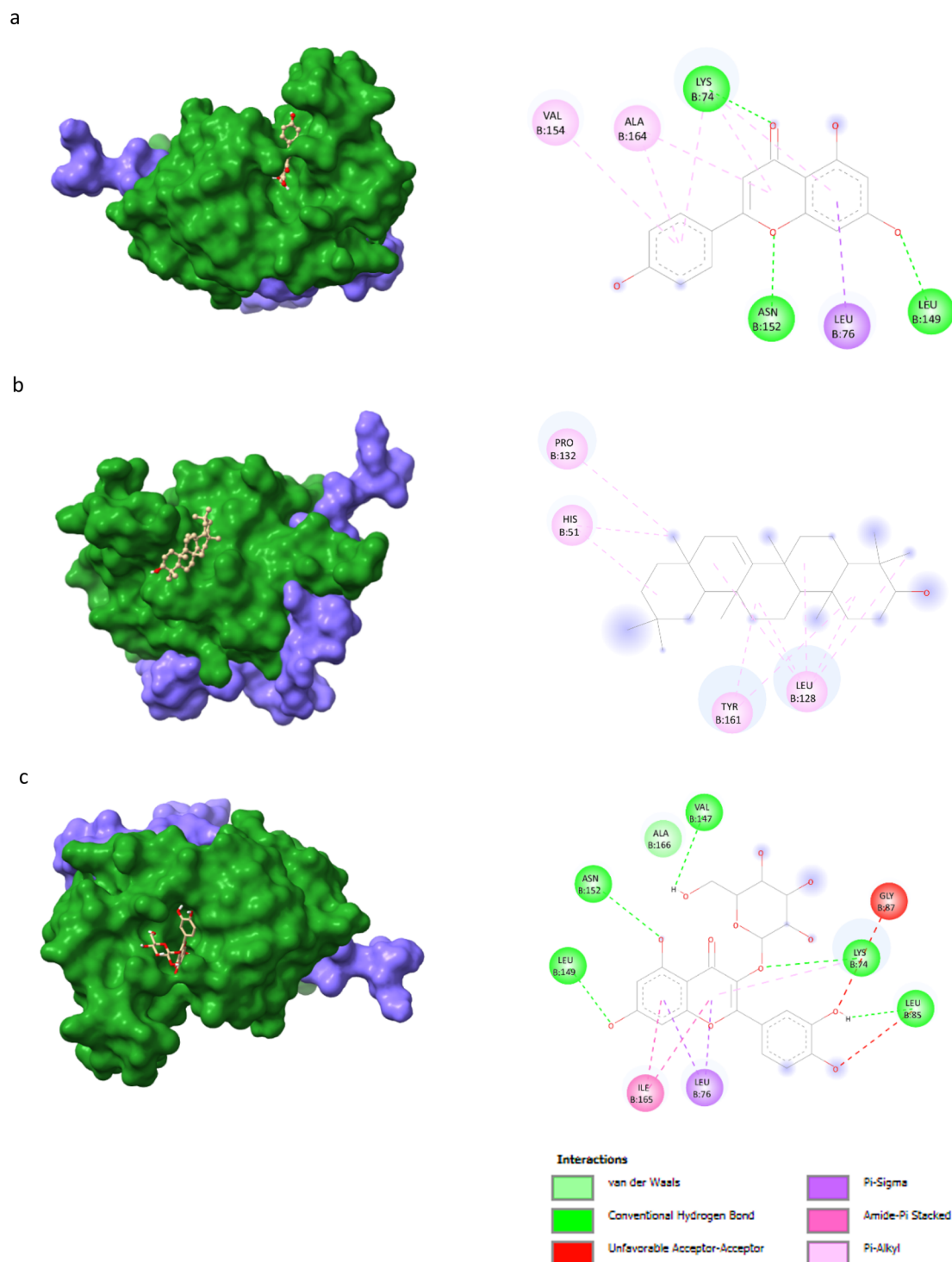


Fig. 2 Docking poses of selected phytochemicals from *N. sativa* (A) apigenin; (B) taraxerol and (C) isoquercetin with Dengue virus NS2BNS3 protease represented as 3-D models (Left side) and their respective 2-D line models (Right side).



number of interacting residues were observed in the interaction of NS2BNS3 with astragalín, they have ten interacting amino acid residues (*i.e.*, Asn167, Trp89, Gly87, Ala166, Val147, Glu88, Ile165, Asn152, Leu86, Lys74). Taraxerol, which has showed highest binding affinity containing 4 interacting amino acid residues *i.e.*, Pro132, His51, Leu128, Tyr161. α -Limonene with least binding affinity showed only two interacting amino acid residues (*i.e.*, Tyr161, Leu128). Docking interactions of selected phytochemicals from *N. sativa* with NS2BNS3 protease are shown in Table 7. The positive control (curcumin) has shown the binding affinity equals to Dithymoquinone *i.e.*, -7.2 kcal mol $^{-1}$ having two interacting residues *i.e.*, Leu128, Pro132, Tyr161. The docking poses were viewed by ChimeraX whereas interacting amino acid residues were visualized by Discovery Studio. The top 3 docking poses for NS2BNS3 protease are shown in Fig. 2. The docking results of compounds against NS2BNS3 can be compared in following order: Taraxerol > Isoquercetin > Apigenin = β -Sitosterol = Cycloeucaleanol = Stigmasterol > Rutin > Campesterol > Astragalín > Nigellícin > Nigellídine > Dithymoquinone > Nigellimine > Carvone = Thymohydroquinone = Thymoquinone > α -Limonene = Nigelline.

For NS5 the phytochemicals showed binding energies ranging from -6.0 kcal mol $^{-1}$ to -9.9 kcal mol $^{-1}$ (Table 8). The highest binding affinity was recorded for apigenin with highest binding energy (*i.e.*, -9.9 kcal mol $^{-1}$), with rutin being the second highest (*i.e.*, -9.3 kcal mol $^{-1}$), while α -limonene bound with the lowest affinity (*i.e.*, 6.0 kcal mol $^{-1}$). Apigenin while having the highest binding affinity involved 06 interacting amino acid residues (*i.e.*, Leu94, Ile72, Pro298, Lys355, Val66, Gln351). Rutin with second highest binding affinity involved Ile691, Gly260, Arg688, Asp690, Arg540, Val687, Ala259, and

Thr360 as their 8 interacting amino acids. Highest number of interacting residues were observed in docking models of NS5-nigellícin (*i.e.*, Arg352, Lys74, Arg352, Glu151, Val66, Leu94, Asn690, Ile165, Pro298) and NS5-nigellídine (*i.e.*, Pro73, Lys96, Ile72, Leu94, Val66, Lys355, Pro298, Arg581, Glu296), both displayed 09 residues. The lowest *i.e.*, 03 number of binding residues were involved in NS5-dithymoquinone (Val687, His52, Pro692) and NS5-taraxerol (Tyr606, Ile797, Val603). The positive control *S*-adenosyl-L-homocysteine (SAH) showed a binding affinity lesser than the majority of understudy phytochemicals. The binding energy -7.2 kcal mol $^{-1}$ was higher than apigenin, astragalín, β -sitosterol, campesterol, cycloeucaleanol, dithymoquinone, isoquercetin, nigellícin nigellídine, rutin, stigmasterol and taraxerol. The docking poses and interaction of amino acid residues was visualized by chimeraX and Discovery studio, respectively (the selected 3 docking poses for NS5 polymerase are shown in Fig. 3). 3D and 2D line models of remaining compounds are given in ESI Fig. S1.† The docking results of compounds can be summed up based on their binding affinities against NS5 in following order: Apigenin > Nigellícin > Rutin > Taraxerol > Stigmasterol > Isoquercetin > Astragalín = Campesterol > Nigellídine > β -Sitosterol = Cycloeucaleanol > Dithymoquinone > Thymoquinone > Thymohydroquinone > Carvone > Nigelline > Nigellimine > α -Limonene.

MD simulations

MD simulations in normal mode analysis have been performed for NS2BNS3 and NS5-apigenin based on their highest docking score. The highest peaks, which correspond to protein areas with high deformability, are shown in Fig. 4A and 5A. The normal mode analysis and the PDB field of the complexes are

Table 8 Docking interactions of selected phytochemicals from *N. sativa* with Dengue virus NS5.

Ligand	Receptor proteins	Binding energies (kcal mol $^{-1}$)	Interacting amino acid residues
Apigenin	NS5	-9.9	Leu94, Ile72, Pro298, Lys355, Val66, Gln351
Astragalín	NS5	-8.5	Thr51, Ile691, Asp690, His52, Gln693, Val687, Ala259
β -Sitosterol	NS5	-8.0	His52, Ile691, Ala535, Val687, Ala259, Tyr119
Campesterol	NS5	-8.5	Tyr119, Ala535, His52, Val687, Ala259, Pro363, Tyr119
Carvone	NS5	-6.4	Pro298, Val66, Ile72, Arg581, Pro582
Cycloeucaleanol	NS5	-8.0	Val603, Phe485, Ile797, Tyr606, Asp663
Dithymoquinone	NS5	-7.5	Val687, His52, Pro692
α -Limonene	NS5	-6.0	Pro298, Val66, Lys95, Pro73, Leu94, Ile72
Isoquercetin	NS5	-8.6	His52, Thr51, Ile691, Asp690, Gln693, Val687, Ala259
Nigellícin	NS5	-9.1	Arg352, Lys74, Arg352, Glu151, Val66, Leu94, Asn690, Ile165, Pro298
Nigellídine	NS5	-8.4	Pro73, Lys96, Ile72, Leu94, Val66, Lys355, Pro298, Arg581, Glu296
Nigellimine	NS5	-6.1	Lys300, Lys355, Tyr119, Thr360, His52, Ala259
Nigelline	NS5	-6.2	Glu296, Pro298, Asn69, Lys355, Arg352, Val66
Rutin	NS5	-9.3	Ile691, Gly260, Arg688, Asp690, Arg540, Val687, Ala259, Thr360
Stigmasterol	NS5	-8.8	His52, Val687, Ala259, Tyr119, Pro363
Taraxerol	NS5	-9.0	Tyr606, Ile797, Val603
Thymohydroquinone	NS5	-6.5	Lys355, Lys95, Leu94, Pro298
Thymoquinone	NS5	-6.7	Lys95, Leu94, Pro298, Lys355
<i>S</i> -Adenosyl-L-homocysteine (SAH) (positive control)	NS5	-7.2	Glu356, Lys300, Val687, Ala259, Thr51, Ile691, His52, Ala535

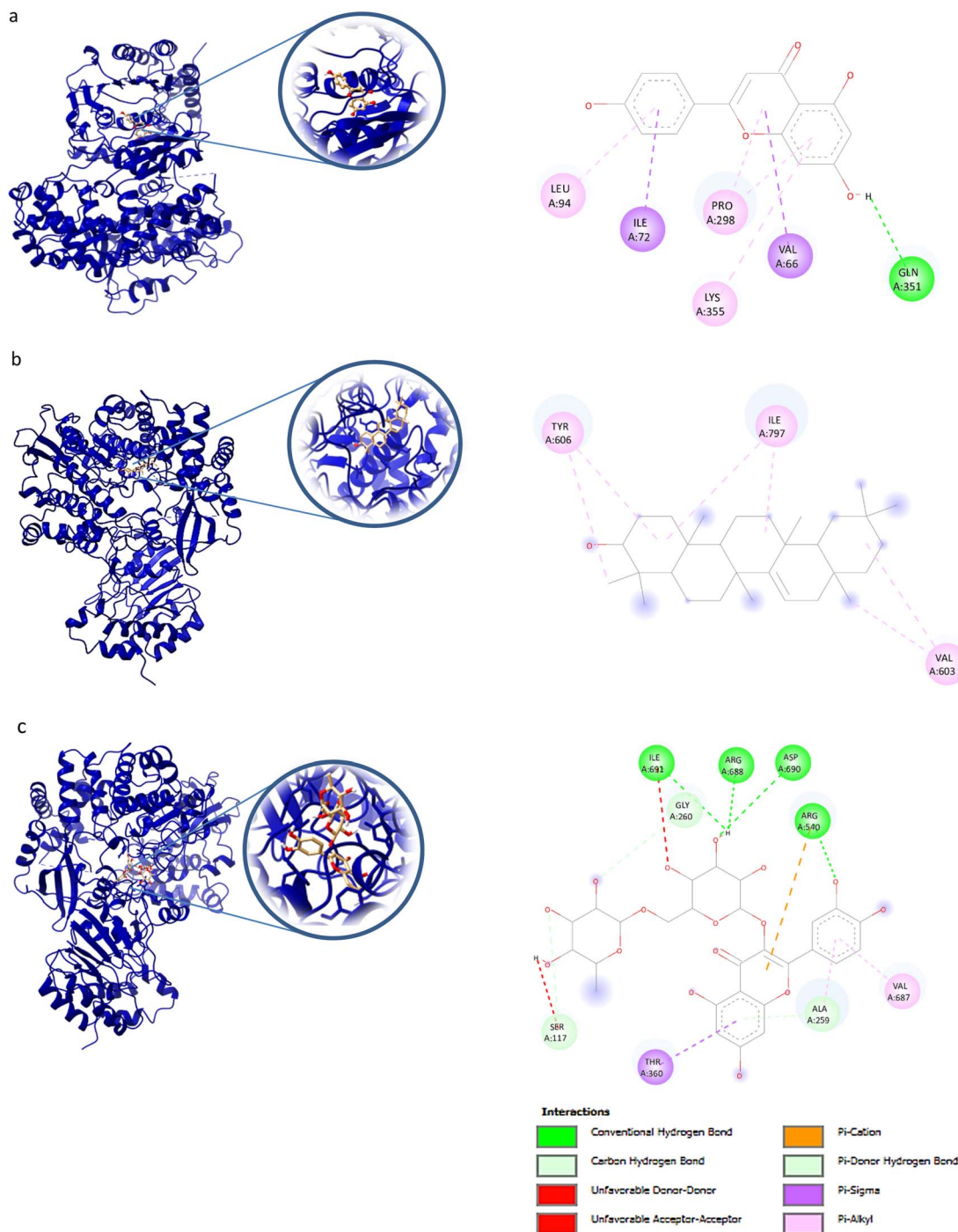


Fig. 3 Docking poses of selected phytochemicals from *N. sativa* (A) apigenin; (B) taraxerol and (C) rutin with Dengue virus NS5 polymerase represented as 3-D models (Left side) and their respective 2-D line models (Right side).

compared in the *B*-factor graphics. The *B*-factor column provides an averaged RMS since many PDB files of averaged NMR models lack *B*-factors (Fig. 4B and 5B). Every normal mode has an associated eigenvalue (Fig. 4C and 5C) and variance (Fig. 4D and 5D). The energy needed to distort the composite is correlated with the eigenvalue, and the smaller the value, the simpler the deformation. Correlations between residues in a complex are shown by the covariance matrix in Fig. 4E and 5E.

The white hue denotes uncorrelated motion, whereas the red color depicts a respectable connection between residues. Anti-correlations are further illustrated by the blue color, the better the complex, the better the correlation. The relationships between the atoms are defined by the elastic maps of the complexes, where firmer regions are shown by darker gray areas and are illustrated in Fig. 5F and 6F.



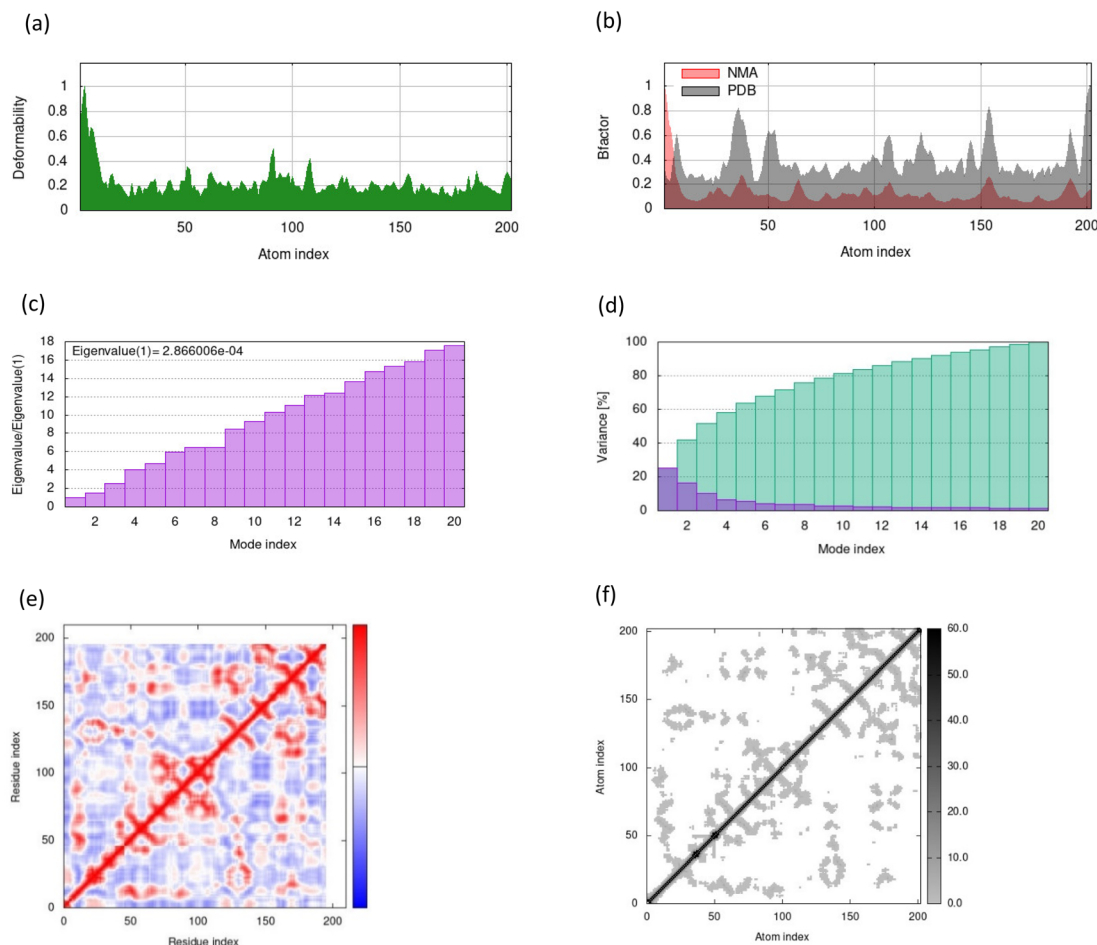


Fig. 4 Results from molecular dynamic simulations for the NS2BNS3-taraxerol complex, including (A) deformability; (B) *B*-factor plot; (C) eigenvalue plot; (D) variance plot; (E) covariance plot and (F) elastic network model.

The structural flexibility of the NS2BNS3-taraxerol docking complex and the NS5-apigenin complex was evaluated by Cabs-flex 2.0. Our results indicated that the RMSF value of NS2BNS3 when bound with naturally existing ligand glycerol was as high as 08 Å and it reduced after binding with taraxerol, and the RMSF value of the complex was recorded below 4.5 Å (Fig. 6A and B). Similarly, after binding with apigenin, the RMSF value of the NS5 protein was reduced (4.0 Å) than the value while being complexed with SAH (Fig. 6C and D). Since the RMSF value recorded for selected complexes was closer to the ideal value, *i.e.*, 3.8, the complexes were said to have stable interactions.

Discussion

Discovery and the development of novel drugs are expensive and time-consuming processes.³⁹ A planned computational approach for rational drug design has been useful in assessing the potential of selected phytochemicals before wet-lab testing by predicting their drug likeness, oral bioavailability, efficacy, toxicity risk assessment, and so on.^{29,32} *In silico* screening and prediction of phytoconstituents with good pharmacodynamic and pharmacokinetic properties is time-saving and cost-

effective. The current study attempted to investigate the antiviral potential of selected phytochemicals from *N. sativa* that have been previously documented to possess antiviral, antibacterial, and antifungal potentials. It is one of the most widely used herbal plants in the world. *N. sativa* oil was given to a 46 year-old HIV-positive patient in a documented clinical case and the findings of the investigation demonstrated full seroconversion and rescue.^{48,49} HCV-RNA was undetectable in 195 HCV patients who received *N. sativa* and chloroquine combination treatment, according to an initial trial.^{50–53} The current study attempted to investigate the antiviral potential of selected phytochemicals such as apigenin,⁵⁴ nigellicine, thymohydroquinone,⁵⁵ nigellicine, dithymoquinone,⁵⁵ nigliimine, and nigellidine, that have been tested *in vitro* to possess antiviral potentials.

To compute the drug potential of understudy compounds, all 18 active phytoconstituents were tested using Lipinski's rule of five which envisages that strong absorption/permeation is expected when the MW < 500, clog *P* ≤ 5.0, there are ≤5 H-bond donors and ≤10 H-bond acceptors with no more than 1 Lipinski's violation otherwise its bioavailability is compromised.^{56,57} Out of 18 subjected phytoconstituents from *N. sativa* three compounds *viz.* astragalin, isoquercetin and rutin

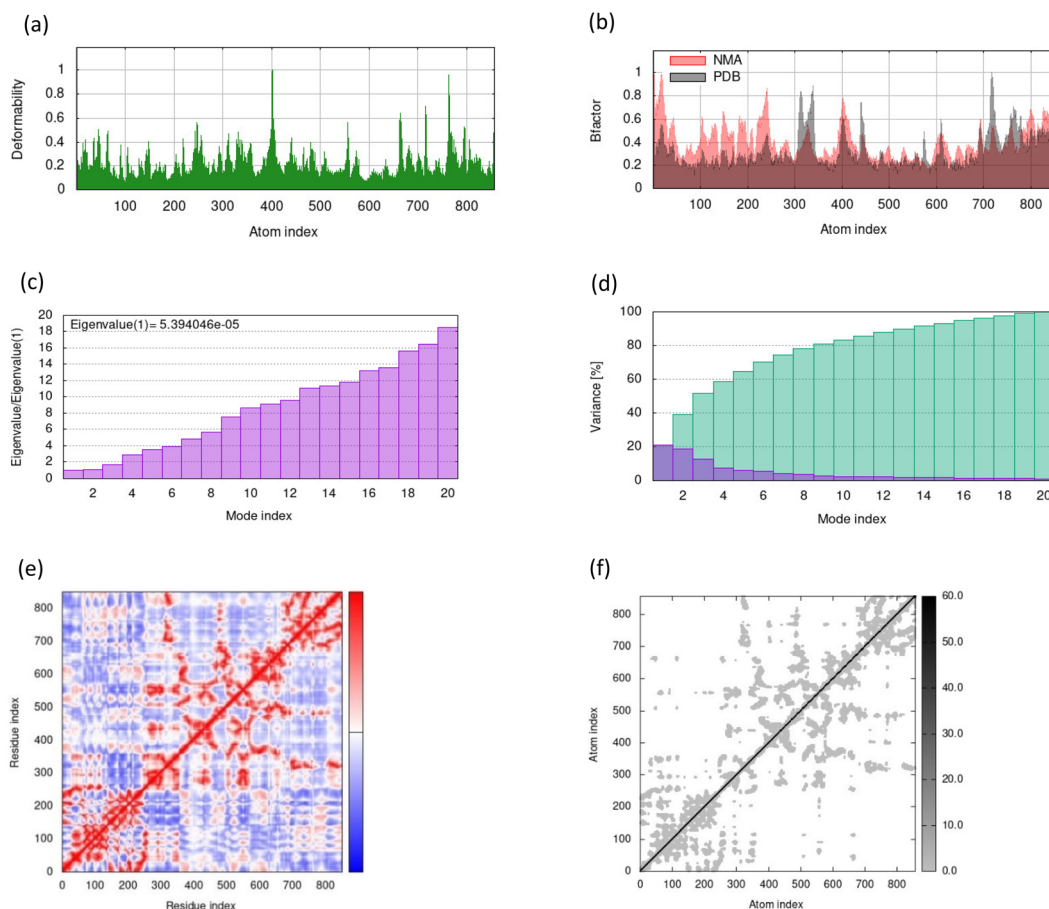


Fig. 5 Results from molecular dynamic simulations for the NS5-apigenin complex, including (A) deformability; (B) *B*-factor plot; (C) eigenvalue plot; (D) variance plot; (E) covariance plot and (F) elastic network model.

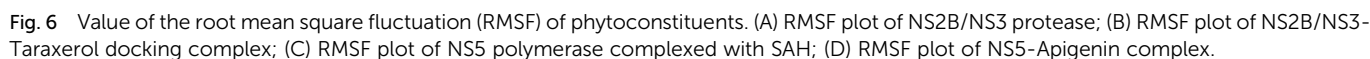
exhibited more than one Lipinski's violation and remaining fifteen phytochemicals qualified the criteria for being a good lead compound (Table 2). The phytochemicals were further evaluated for their drug-likeness capacity and to prevent inappropriate chemicals for future drug screening and to begin *in vitro* and *in vivo* evaluation, toxicity potential and bioavailability must be assessed.⁵⁸ Apigenin, campesterol, cycloeculanol, β -sitosterol, dithymoquinone, stigmasterol, taraxerol, nigellidine, and nigelline have shown significant bioavailability (Table 3) and were found to have potential as significant protease and enzyme inhibitors.^{59–61} The majority of phytochemicals, including apigenin, campesterol, carvone, nigellidine, nigellimine, and nigelline, showed high human intestinal absorption (GI absorption), lipophilicity and blood–brain permeation, indicating high absorption and transport kinetics in the stomach and the ability to easily cross the BBB (Table 4). Toxicity risk assessment BBB (Table 5) through LD₅₀ value and toxicity class BBB suggests that apigenin, astragaline, taraxerol and thymoquinone are expected to be safe as they exhibited non-mutagenic, non-tumorigenic, non-carcinogenic, non-irritant behaviour with no threats to reproductive systems. In compared to toxicity class 1–3, the LD₅₀ values for oral acute toxicity of the majority of understudy phytochemicals fell into

toxicity class 4 and 5, making them expectedly less dangerous. Only one compound was found in toxicity class 3 (*i.e.*, nigellimine) whereas taraxerol secured place in class VI and was rendered as expectedly non-toxic (Table 6).

The binding affinities of 18 selected phytochemicals from *N. sativa* have been studied against two target proteins of Dengue virus, *i.e.*, NS2BNS3 protease and NS5 polymerase, using molecular, pharmacokinetic, and *in silico* approaches. A member of the S7 family of serine proteases, the DENV NS3 protease domain (NS3pro), which becomes fully active by interacting with the cofactor NS2B, facilitates polyprotein processing at certain sites. Due to its critical involvement in the viral replication cycle, NS2BNS3 protease is regarded as a significant prospective therapeutic target.^{62,63} NS5 is the most conserved and largest non-structural protein encoded by flaviviruses. Increased structural understanding and multifunctionality of the dengue NS5 protein will help accelerate drug development efforts targeting NS5 as an antiviral target.^{64,65} Improved *in silico* approaches, such as molecular docking, pharmacology networks, molecular dynamic simulations, and so on, now significantly contribute to the separation of prospective drugs.³⁰

Molecular docking is a popular approach in computational drug discovery and development to anticipate the orientation





of small molecules coupled to an enzyme or receptor binding pocket.⁶⁶ Based on antiviral phytochemicals, the active site residues of the dengue NS2B/NS3 protease (Table 7) Lys73, Lys74, Leu76, Trp83, His51, Phe130, Leu128, Leu85, Glu88, Gly87, Ile165, Ala166, Ala164, Asn152, and Asn167 were observed to be involved in hydrophobic and hydrogen bond formation.^{12,67} Taraxerol ($-9.1 \text{ kcal mol}^{-1}$) interacted with Pro132, His51, Leu128, Tyr161, while astragalín showed binding affinity of $-8.0 \text{ kcal mol}^{-1}$ and interacted with residues Asn167, Trp89, Gly87, Ala166, Val147, Glu88, Ile165, Asn152, Leu86, Lys74. β -Sitosterol ($-8.3 \text{ kcal mol}^{-1}$), campesterol ($8.1 \text{ kcal mol}^{-1}$), and nigellidine ($-7.3 \text{ kcal mol}^{-1}$), each interacted actively with four amino acid residues of NS2B/NS3. For NS5 polymerase, apigenin ($9.9 \text{ kcal mol}^{-1}$) formed binding interactions with Leu94, Ile72, Pro298, Lys355, Val66, Gln351. Nigellidine (Arg352, Lys74, Arg352, Glu151, Val66, Leu94, Asn690, Ile165, Pro298) and nigellidine (Pro73, Lys96, Ile72, Leu94, Val66, Lys355, Pro298, Arg581, Glu296) interacted with nine residues in an active pocket with the binding affinity of $-9.1 \text{ kcal mol}^{-1}$ and $-8.4 \text{ kcal mol}^{-1}$, respectively (Table 8). Cycloecalenol ($-8.0 \text{ kcal mol}^{-1}$) interacted with residues of Val603, Phe485, Ile797, Tyr606, Asp663 in the hydrophobic pocket. The NS5 residues Ser56, Gly81, Cys82, Arg84, Gly85, Thr104, Lys105, His110, Asp131, Val132, Asp146, and Gly148 are conserved in the binding

pocket whereas the hydrophobic pocket contains Trp302, Phe354, Val358, Val577, Val579, Val603, and Gly599, Ala406, Ala407, Asn492, Glu507, Tyr606, and Ile797.^{68–70} An analysis of the interaction between dengue protease and ligands indicated significant contributions from van der Waals forces, conventional hydrogen bonding, carbon hydrogen bonding, alkyl interactions, and pi-alkyl interactions.⁷¹ The primary component ensuring the stability of the protein-ligand complex structure is hydrophobic interaction. However, hydrogen bonding has a less significant impact in the structural stability of the protein-ligand complexes. It is expected that existence of the hydrogen bond will matter to provide a better docking score because it has the highest negative weighted factor value among the other variables.⁴² MD simulations confirmed the structural flexibility of the NS2B/NS3-taraxerol docking complex and the NS5-apigenin complex, whose root mean square value fluctuated below the 5 Å value. RMSF values are utilized to calculate the atomic positional variation of each residue based on its C-alpha (Ca) atom.^{72,73} The complexes NS2BNS3-taraxerol and NS5-apigenin under iMODS investigation have a significant amount of deformability, according to the normal mode analysis. The low eigenvalue, covariance map, and elastic network of the complex under study are indications of the molecule's potential deformability and allowable dynamics.

The current study used several prediction techniques to anticipate the oral bioavailability of substances, which could lead to the development of novel, safer medications. *In silico* investigations can save a significant amount of cash and time before proceeding to the experimental stage. Dengue virus replication depends on two proteins, NS5 polymerase and NS2BNS3 main protease complexes,⁷⁴ which are also the primary focus of antiviral drug research. A previous study⁶⁶ has used the same target receptors to identify the potential inhibitors from *Carica papaya* and their results from molecular docking and ligand chemistry have been compared with our 18 understudied phytochemicals of *N. sativa* which suggests that apigenin, nigellicine, nigellidine, dithymoquinone, taraxerol, campesterol, cycloeucalenol, and beta-sitosterol have been revealed as the most potential lead compounds. According to another research, *Nigella sativa* demonstrated potent virucidal characteristics that inactivated encapsulated viruses from 2log10 (or 99%) to >4log10 (or 99.99%). The compound Apigenin was also shown to have antiviral, antibacterial, and antifungal effects.⁷⁵

Conclusion

In conclusion, our research has identified a small number of possible phytochemicals with potential inhibitory effects against Dengue NS2BNS3 and NS5, which may be used for future *in vitro* and animal testing, formulation, and drug development. These phytoconstituents can be further investigated *in vitro* and *in vivo* as possible antiviral drugs for the treatment of dengue fever infections due to their expected bioavailability, druglikeness, and non-toxic and non-mutagenic effects.

Author contributions

M. M. and H. A. K. conceptualized the study and performed the formal analyses. Validation was done by M. M., N. S. S. Z. and H. A. K. M. M. wrote the original draft. Original draft was reviewed and edited by N. S. S. Z. and H. A. K. All authors have read and agreed to the published version of the manuscript.

Conflicts of interest

The authors declare that they have no conflict of interest.

Acknowledgements

We acknowledge the support of the Department of Industrial Biotechnology at Atta-ur-Rahman School of Applied Biosciences, NUST, for providing the research facilities and funding.

References

- M. J. Sabir, N. B. S. Al-Saud and S. M. Hassan, *Saudi J. Biol. Sci.*, 2021, **28**, 5074–5080.
- M. T. Qamar, U. A. Ashfaq, K. Tusleem, A. Mumtaz, Q. Tariq, A. Goheer and B. Ahmed, *Pak. J. Pharm. Sci.*, 2017, **30**, 2119–2137.
- S. J. Thomas and T. P. Endy, *Curr. Opin. Infect. Dis.*, 2011, **24**, 442–450.
- A. N. Anoopkumar and E. M. Aneesh, *Environ. Dev. Sustain.*, 2021, **23**, 11217–11239.
- S. Swaminathan and N. Khanna, *Int. J. Infect. Dis.*, 2019, **84**, S80–S86.
- C. P. Simmons and J. E. Farrar, *N. Engl. J. Med.*, 2012, **366**, 1423–1432.
- A. Wilder-Smith, *Bundesgesundheitsblatt - Gesundheitsforsch. - Gesundheitsschutz*, 2020, **63**, 40–44.
- M. Mukhtar, A. W. Wajeeha and N. Bibi, *Int. J. Mol. Sci.*, 2022, **23**, 13911.
- J. Park, J. Kim and Y. S. Jang, *J. Microbiol.*, 2022, **60**, 247–254.
- C. J. Wollner, M. Richner, M. A. Hassert, A. K. Pinto, J. D. Brien and J. M. Richner, *J. Virol.*, 2021, 95.
- T. N. Adikari, N. Riaz, C. Sigera, P. Leung, B. M. Valencia, K. Barton, M. A. Smith, R. A. Bull, H. Li, F. Luciani, P. Weeratunga, T. L. Thein, V. W. X. Lim, Y. S. Leo, S. Rajapakse, K. Fink, A. R. Lloyd, D. Fernando and C. Rodrigo, *Sci. Rep.*, 2020, **10**, 18196.
- R. A. D. Costa, J. Rocha, A. S. Pinheiro, A. Costa, E. Rocha, R. C. Silva, A. D. S. Goncalves, C. B. R. Santos and D. Brasil, *Molecules*, 2022, 27.
- L. Lim, M. Dang, A. Roy, J. Kang and J. Song, *ACS Omega*, 2020, **5**, 25677–25686.
- S. P. Lim, C. G. Noble, C. C. Seh, T. S. Soh, A. El Sahili, G. K. Chan, J. Lescar, R. Arora, T. Benson, S. Nilar, U. Manjunatha, K. F. Wan, H. Dong, X. Xie, P. Y. Shi and F. Yokokawa, *PLoS Pathog.*, 2016, **12**, e1005737.
- H. Shimizu, A. Saito, J. Mikuni, E. E. Nakayama, H. Koyama, T. Honma, M. Shirouzu, S. I. Sekine and T. Shioda, *PLoS Negl. Trop. Dis.*, 2019, **13**, e0007894.
- P. Coulerie, M. Nour, A. Maciuk, C. Eydoux, J. C. Guillemot, N. Lebouvier, E. Hnawia, K. Leblanc, G. Lewin, B. Canard and B. Figadere, *Planta Med.*, 2013, **79**, 1313–1318.
- A. Wahaab, B. E. Mustafa, M. Hameed, N. J. Stevenson, M. N. Anwar, K. Liu, J. Wei, Y. Qiu and Z. J. V. Ma, *Potential Role of Flavivirus NS2B-NS3 Proteases in Viral Pathogenesis and Anti-flavivirus Drug Discovery Employing Animal Cells and Models: A Review*, 2021, vol. 14, p. 44.
- N. Rasool, A. Ashraf, M. Waseem, W. Hussain and S. Mahmood, *Turk. J. Biochem.*, 2019, **44**, 261–277.
- K. Mishra, N. Sharma, D. Diwaker, L. Ganju and S. Singh, *J. Virol. Antivir. Res.*, 2013, **2**, 2.
- R. Ghildiyal, V. Prakash, V. Chaudhary, V. Gupta and R. Gabrani, in *Plant-derived bioactives*, Springer, 2020, pp. 279–295.
- T. Rathinasabapathy, L. P. Sakthivel and S. Komarnytsky, *J. Agric. Food Chem.*, 2022, **70**, 2064–2076.
- S. Desai, S. H. Saheb, K. K. Das and S. Haseena, *Phytochemical Analysis of Nigella sativa and its Antidiabetic Effect*, 2015.
- S. L. Khan, F. A. Siddiqui, S. P. Jain and G. M. Sonwane, *Coronaviruses*, 2021, **2**, 384–402.
- A. Ahmad, A. Husain, M. Mujeeb, S. A. Khan, A. K. Najmi, N. A. Siddique, Z. A. Damanhouri and F. Anwar, *Asian Pac. J. Trop. Biomed.*, 2013, **3**, 337–352.



- 25 J. Anjum, M. A. Sheikh, A. Tiwari, S. Sharma and B. Kumari, in *Microbial and Biotechnological Interventions in Bioremediation and Phytoremediation*, Springer, 2022, pp. 425–444.
- 26 A. McBride, P. Mehta, L. Rivino, A. V. Ramanan and S. J. Yacoub, *Lancet Microbe*, 2021, **2**, e277.
- 27 Z. Hu, J. Lin, J. Chen, T. Cai, L. Xia, Y. Liu, X. Song and Z. J. He, *Front. Pharmacol.*, 2021, **12**, 1380.
- 28 R. Chakravarti, R. Singh, A. Ghosh, D. Dey, P. Sharma, R. Velayutham, S. Roy and D. J. Ghosh, *RSC Adv.*, 2021, **11**, 16711–16735.
- 29 S. Siddiqui, S. Upadhyay, R. Ahmad, A. Gupta, A. Srivastava, A. Trivedi, I. Husain, B. Ahmad, M. Ahamed and M. A. Khan, *J. Biomol. Struct. Dyn.*, 2022, **40**, 3928–3948.
- 30 P. R. Ferdian, R. R. Elfirta, Q. Emilia and A. Z. N. Ikhwan, *Ann. Bogor.*, 2021, **24**.
- 31 R. S. Basurra, S. M. Wang and M. A. Alhoot, *J. Pure Appl. Microbiol.*, 2021, **15**, 29–41.
- 32 R. Ahmad, *PeerJ*, 2019, **7**, e6012.
- 33 C. A. Lipinski, F. Lombardo, B. W. Dominy and P. J. Feeney, *Adv. Drug Delivery Rev.*, 2012, **64**, 4–17.
- 34 P. Kulkarni, Y. Walunj and N. J. T. C. Dongare, *Theoretical Validation of Medicinal Properties of Ocimum sanctum*, 2021, p. 53.
- 35 D. Jindal and B. Rani, *In Silico Studies of Phytoconstituents from Piper longum and Ocimum sanctum as ACE2 and TMRSS2 Inhibitors: Strategies to Combat COVID-19*, 2022, pp. 1–18.
- 36 A. Agrwal, R. Saini, S. Bhandri, S. Verma, P. Srivastava and O. J. Prakash, *Mater. Today: Proc.*, 2022, **67**, 598–604.
- 37 P. Banerjee, A. O. Eckert, A. K. Schrey and R. J. Preissner, *Nucleic Acids Res.*, 2018, **46**, W257–W263.
- 38 M. N. Drwal, P. Banerjee, M. Dunkel, M. R. Wettig and R. J. Preissner, *Nucleic Acids Res.*, 2014, **42**, W53–W58.
- 39 T. Khan, R. Ahmad, I. Azad, S. Raza, S. Joshi and A. R. Khan, *Comput. Biol. Chem.*, 2018, **75**, 178–195.
- 40 E. Berinyuy, J. Ibrahim and B. Alozieuwa, *AROC in Pharmaceutical and Biotechnology*, 2021, vol. 1, pp. 1–8.
- 41 D. Flamandita, K. Lischer, D. K. Pratami, R. Aditama and M. Sahlan, *presented in part at the International Conference on Emerging Applications in Material Science and Technology: Iceamst 2020*, 2020.
- 42 R. Novianty, S. Ananta and M. A. Karim, *J. Phys. Conf.*, 2021, **1788**.
- 43 V. Adrianne Dien-Yu, H. Siaw-San, C. Xavier Wezen and S. Edmund Ui-Hang, *J. Appl. Biol. Biotechnol.*, 2022, **10**(1), 38–44.
- 44 M. Ebrahimi and M. Alijanianzadeh, *J. Mol. Divers.*, 2023, **1**–24.
- 45 A. Ibrahim and E. U. Haq, *Nat. Product Chem.*, 2023, **12**, 97–108.
- 46 Y. Yuniwati, M. F. R. Syaban, S. G. Anoraga and F. L. Sabila, *Acta. Inform. Med.*, 2022, **30**, 91–95.
- 47 R. Srivastava, *Front. Chem.*, 2022, **10**, 843642.
- 48 A. A. Onifade, A. P. Jewell and W. A. Adedeji, *Afr. J. Tradit. Complement Altern. Med.*, 2013, **10**, 332–335.
- 49 M. Riaz, M. Khan, R. Ahmad, L. H. AlLehaibi, N. Rahman and D. Deqiang, *Bol. Latinoam. Caribe Plantas Med. Aromat.*, 2022, **21**, 176–206.
- 50 S. Thakur, H. Kaurav and G. Chaudhary, *Int. J. Res. Rev.*, 2021, **8**, 342–357.
- 51 M. Akram Khan and M. Afzal, *Inflammopharmacology*, 2016, **24**, 67–79.
- 52 M. F. Ahmad, F. A. Ahmad, S. A. Ashraf, H. H. Saad, S. Wahab, M. I. Khan, M. Ali, S. Mohan, K. R. Hakeem and M. T. Athar, *J. Herb. Med.*, 2021, **25**, 100404.
- 53 Z. Sheir, G. Badra and O. Salama, *J. Liver*, 2013, **2**.
- 54 A. Hakobyan, E. Arabyan, A. Avetisyan, L. Abroyan, L. Hakobyan and H. Zakaryan, *J. Arch. Virol.*, 2016, **161**, 3445–3453.
- 55 E. R. Esharkawy, F. Almalki and T. B. Hadda, *Bioorg. Chem.*, 2022, **120**, 105587.
- 56 H. Tabassum and I. Z. Ahmad, *Med. Chem.*, 2020, **16**, 350–357.
- 57 Y. I. Shaikh, V. S. Shaikh, K. Ahmed, G. M. Nazeruddin and H. M. Pathan, *The revelation of various compounds found in Nigella sativa L.(Black Cumin) and their possibility to inhibit COVID-19 infection based on the molecular docking and physical properties*, 2020, vol. 11, pp. 31–35.
- 58 S. Riaz, M. A. Riaz, A. Zhanzhaksina, S. A. Raja and T. Javed, *Silico Therapeutic and Pharmacological Evaluation of novel apolipoprotein E4 inhibitors against Alzheimer's Disease*, 2022, vol. 3, pp. 22–31.
- 59 I. Ghosh and P. J. Talukdar, *World Sci. News*, 2019, **124**, 264–278.
- 60 S. Bandyopadhyay, O. A. Abiodun, B. C. Ogboo, A. T. Kola-Mustapha, E. I. Attah, L. Edemhanria, A. Kumari, R. Jaganathan and N. S. Adelakun, *J. Biomol. Struct.*, 2021, **1**–17.
- 61 S. Hemaiswarya, P. K. Prabhakar and M. Doble, in *Herb-Drug Combinations*, Springer, 2022, pp. 235–244.
- 62 H. N. Saleem, F. Batool, H. J. Mansoor, S. Shahzad-ul-Hussan and M. Saeed, *ACS Omega*, 2019, **4**, 1525–1533.
- 63 S. Bharadwaj, K. E. Lee, V. D. Dwivedi, U. Yadava, A. Panwar, S. J. Lucas, A. Pandey and S. G. Kang, *Sci. Rep.*, 2019, **9**, 19059.
- 64 J. O. Obi, H. Gutierrez-Barbosa, J. V. Chua and D. J. Deredge, *Trop. Med. Infect. Dis.*, 2021, **6**.
- 65 S. P. Lim, C. G. Noble and P. Y. Shi, *Antiviral Res.*, 2015, **119**, 57–67.
- 66 M. U. Farooq, B. Munir, S. Naeem, M. Yameen, S. Z. Iqbal, A. Ahmad, M. A. Mustaan, M. W. Noor, M. A. Nadeem and A. J. Ghaffar, *Pak. J. Pharm. Sci.*, 2020, **33**.
- 67 M. S. S. Shimu, S. Mahmud, T. E. Tallei, S. A. Sami, A. A. Adam, U. K. Acharjee, G. K. Paul, T. B. Emran, S. Zaman, M. S. Uddin, M. A. Saleh, S. Alshehri, M. M. Ghoneim, M. Alruwali, A. J. Obaidullah, N. R. Jui, J. Kim and B. Kim, *Molecules*, 2022, **27**(3), 653.
- 68 C. N. Powers and W. N. Setzer, *Comb. Chem. High Throughput Screening*, 2016, **19**, 516–536.
- 69 M. Tahir ul Qamar, A. Maryam, I. Muneer, F. Xing, U. A. Ashfaq, F. A. Khan, F. Anwar, M. H. Geesi, R. R. Khalid and S. A. Rauf, *Sci. Rep.*, 2019, **9**, 1–16.



- 70 K. Niranjana, M. Kumar and S. J. Bhatnagar, *Indian J. Appl. Phys.*, 2017, **6**, 114–118.
- 71 A. A. Poola, P. S. Prabhu, T. K. Murthy, M. Murahari, S. Krishna, M. Samantaray and A. J. Ramaswamy, *Front. Mol. Biosci.*, 2023, **10**.
- 72 S. A. Mir, A. Firoz, M. Alaidarous, B. Alshehri, A. A. B. Dukhyil, S. Banawas, S. A. Alsagaby, W. Alturaiki, G. A. Bhat and F. J. Kashoo, *Saudi J. Biol. Sci.*, 2022, **29**, 394–401.
- 73 A. Nag, P. Verma, S. Paul and R. Kundu, *Appl. Biochem. Biotechnol.*, 2022, **194**, 4867–4891.
- 74 M. Hasan, M. M. Mia, S. U. Munna and M. M. H. Talha, *Inform. Med. Unlocked*, 2022, **30**, 100932.
- 75 S. Sharad and S. J. P. Kapur, *Indian herb-derived phytoconstituent-based antiviral, antimicrobial and antifungal formulation: an oral rinse candidate for oral hygiene and the potential prevention of COVID-19 outbreaks*, 2021, vol. 10, p. 1130.
- 76 Y. Kabir, Y. Akasaka-Hashimoto, K. Kubota and M. Komai, *Heliyon*, 2020, **6**(10), e05343.

