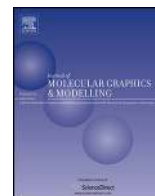




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## In-silico screening for anti-Zika virus phytochemicals



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### ABSTRACT

Zika virus (ZIKV) is an arbovirus that has infected hundreds of thousands of people and is a rapidly expanding epidemic across Central and South America. ZIKV infection has caused serious, albeit rare, complications including Guillain–Barré syndrome and congenital microcephaly. There are currently no vaccines or antiviral agents to treat or prevent ZIKV infection, but there are several ZIKV non-structural proteins that may serve as promising antiviral drug targets. In this work, we have carried out an *in-silico* search for potential anti-Zika viral agents from natural sources. We have generated ZIKV protease, methyltransferase, and RNA-dependent RNA polymerase using homology modeling techniques and we have carried out molecular docking analyses of our in-house virtual library of phytochemicals with these protein targets as well as with ZIKV helicase. Overall, 2263 plant-derived secondary metabolites have been docked. Of these, 43 compounds that have drug-like properties have exhibited remarkable docking profiles to one or more of the ZIKV protein targets, and several of these are found in relatively common herbal medicines, suggesting promise for natural and inexpensive antiviral therapy for this emerging tropical disease.

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## 1. Introduction

Zika virus (ZIKV) is an emerging arbovirus that belongs to the genus *Flavivirus* in the family *Flaviviridae*. Other members of the genus include the West Nile virus, dengue virus, yellow fever virus, and tick-borne encephalitis virus [1]. Zika virus was initially isolated from a rhesus monkey in the Zika forest of Uganda [2]. There had been sporadic human infections in sub-Saharan Africa and Southeast Asia [3]. However, in 2007 there was an outbreak in Yap Island (Micronesia) and subsequent epidemics in French Polynesia, New Caledonia, the Cook Islands, and Easter Island between 2013 and 2014 [4]. In 2015 the number of human infections with ZIKV showed a dramatic increase in the tropical Americas, particularly Brazil [5,6]. By the end of 2015, there have been an estimated 0.5–1.3 million cases of ZIKV infection [7].

The virus is primarily transmitted by *Aedes* spp. mosquitoes, including *Ae. aegypti*, *Ae. africanus*, *Ae. apicoargenteus*, *Ae. furcifer*, *Ae. hensilli*, *Ae. luteocephalus*, and *Ae. vitattus* [3]. However, there is now evidence that the disease can be sexually transmitted; replicative Zika viruses have been found in the semen of an infected man [8]. In addition, there is evidence that the virus can be transmitted

perinatally and transplacentally; Zika viral RNA has been detected in amniotic fluid samples from fetuses of infected mothers [9]. Zika virus infections are typically characterized with symptoms of maculopapular rash, fever, headache, arthralgia, myalgia, and conjunctivitis. In addition, however, ZIKV infection has been associated with additional complications such as Guillain–Barré syndrome [10], congenital microcephaly [11], and macular atrophy [12].

There are currently no vaccines or antiviral agents available to treat or prevent Zika virus infection [13]. However, several *Flavivirus* non-structural proteins have been implicated as potential targets for antiviral drug discovery.

### 1.1. Zika virus protease

ZIKV protease (NS2B–NS3pro) is a trypsin-like serine protease that cleaves the ZIKV polyprotein into individual proteins necessary for viral replication. Flaviviral proteases from dengue virus [14] and West Nile virus [15] have been identified as a potential targets for development of antiviral agents [16].

### 1.2. Zika virus helicase

Flaviviral NS3 helicases possess RNA helicase, nucleoside and RNA triphosphatase activities and are involved both in viral RNA

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replication and virus particle formation. They are essential for viral replication and have been identified as potential drug targets [17].

### 1.3. Zika virus methyltransferase

The flavivirus NS5 methyltransferase (MTase) enzyme is responsible for methylating the viral RNA cap structure [18]. Viruses deficient in cap methylation result in attenuated or non-replicative viruses, and therefore, MTase is an attractive target for discovery of antiviral agents [19,20].

### 1.4. Zika virus RNA-dependent RNA polymerase

RNA-dependent RNA polymerases are employed by flaviviruses and are responsible for initiating and catalyzing replication of viral RNA synthesis in the cytoplasm of infected cells. There are no similar enzymes found in the host since host cells do not require RNA replication. This enzyme, then, is considered to be a prime target for antiviral development [21,22].

In this work, we have carried out an *in-silico* search for potential anti-Zika viral agents from natural sources. We have generated ZIKV protease, methyltransferase, and RNA-dependent RNA polymerase using homology modeling techniques and we have carried out molecular docking analyses of our in-house virtual library of phytochemicals with these protein targets and with the X-ray crystal structure of ZIKV helicase.

## 2. Methodology

### 2.1. Homology modeling

Currently there is only one protein structure available (helicase, PDB 5JMT [23]) for Zika virus non-structural proteins. Homology models for each of the other Zika proteins listed in Table 1 were constructed from crystal structure templates found in the Protein Data Bank using FASTA sequences downloaded from the National Center for Biotechnology Information's (NCBI) GenBank. Sequences with high sequence similarity in the PDB were identified with NCBI's BLAST utility using the BLOSUM80 scoring matrix. Sequences with high similarity to the reference sequence, as well as having good coverage for the active sites in the protein, were chosen for single reference homology modeling.

The protein sequences were first aligned to their respective template sequences using the BLOSUM62 substitution matrix and a protein backbone constructed and superposed to the reference structure using the protein alignment tool in MOE 2014.0901. The homology modeling interface in MOE was used to generate a set of putative protein structures by aligning atomic coordinates of the amino acid sequence to those of the template sequence backbone and minimizing permutations of side chain orientations using the AMBER10:EHT force field [24–26] with reaction field solvation. The candidate structure with the lowest RMSD deviation from the template backbone was selected and optimized using a constrained minimization.

The homology models for the NS3 protease was generated for the stretch of the whole Zika protein (accession number Q32ZE1) that correspond to the incomplete 257 amino acid NSP3 sequence, AHL16750.1, found in the NCBI GenBank. The closest matching template in the PDB was the combined NSP3 Protease-Helicase from Dengue Virus (2WHX) [27]. The protease binding site in this template is poorly defined in the crystal structure with the highest similarity (2WV9) [28], so a second template, from the homologous protease in West Nile Virus (2FP7; exp. value  $4 \times 10^{-7}$ ) [29] with a bound ligand (Bz-Nle-Lys-Arg-Arg-H; (*N*-benzoyl-L-norleucyl-6-ammonio-L-norleucyl-N-5-[amino(imino)methyl]-N-[(2S)-5-carbamimidamido-1-hydroxypentan-2-yl]-L-ornithinamide)) was

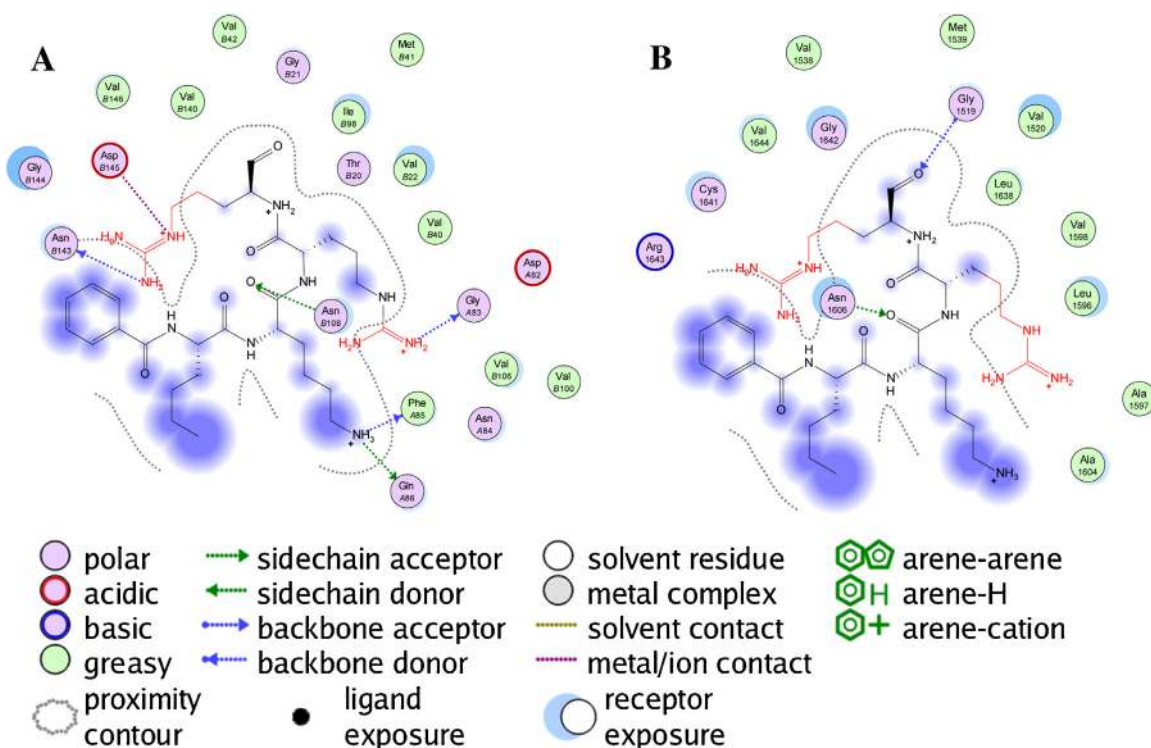
used to generate a homology model specifically for the protease. Since the sequence similarity with the NS3 protease is much lower for this template, a constrained minimization of the peptide-receptor complex was carried out using the AMBER10:EHT force field and the bound peptide removed prior to docking. The RMSD between 2FP7 and the homology model is 0.753 Å for all common atoms. The binding interactions for both are shown in Fig. 1. The GenBank protein sequence A0A0B4ZYV2 was used for the NS5 methyltransferase and RNA-dependent RNA polymerase homology models using the relevant portion of the complete NS5 protein from Japanese Encephalitis Virus (4K6M) [30]. A second homology model for NS5 RdRp was generated using the template structure 2HFZ from West Nile Virus [31]. Protein sequence alignments and Ramachandran plots of the template proteins and the homology modeled ZIKV proteins are available as Supplementary material.

### 2.2. Molecular docking

Protein-ligand docking studies were carried out based on homology-modeled structures of Zika virus (ZIKV) NSP/NS3 protease, based on the crystal structure of Murray Valley encephalitis virus protease, PDB 2WV9 [28]; ZIKV NS5 methyltransferase (MTase), based on the crystal structure of Japanese encephalitis virus methyltransferase, PDB 4K6M [30]; and ZIKV NS5 RNA-dependent RNA polymerase (RdRp), based on crystal structures of West Nile virus RdRp, PDB 2HFZ [31] and Japanese encephalitis virus RdRp, PDB 4K6M [30]. Molecular docking calculations for all compounds with each of the proteins were undertaken using Molegro Virtual Docker (version 6.0, Molegro ApS, Aarhus, Denmark) [32], with a sphere large enough to accommodate the cavity centered on the binding sites of each protein structure in order to allow each ligand to search. Standard protonation states of the proteins based on neutral pH were used in the docking studies. Each protein was used as a rigid model structure; no relaxation of the protein was performed. Assignments of charges on each protein were based on standard templates as part of the Molegro Virtual Docker program; no other charges were necessary to be set. Overall, 2263 plant-derived secondary metabolites have been docked. This molecule set was comprised of 384 alkaloids (158 indole alkaloids, 143 isoquinoline alkaloids, 5 quinoline alkaloids, 18 piperidine alkaloids, 14 steroidal alkaloids, and 46 miscellaneous alkaloids), 670 terpenoids (35 monoterpenoids, 153 sesquiterpenoids, 212 diterpenoids, 83 steroids, 51 limonoids, 42 withanolides, and 97 triterpenoids), 1043 polyphenolic compounds (22 auronos, 72 chalcones, 9 chromones, 111 coumarins, 297 flavonoids, 113 isoflavonoids, 69 lignans, 69 stilbenoids, 29 xanthones, and 172 miscellaneous phenolics), and 166 miscellaneous phytochemicals. This in-house virtual library represents a practical selection of phytochemical classes and structural types. Each ligand structure was built using Spartan '14 for Windows (version 1.1.8, Wavefunction Inc., Irvine, California). For each ligand, a conformational search and geometry optimization was carried out using the MMFF force field [33]. Flexible ligand models were used in the docking and subsequent optimization scheme. Variable orientations of each of the ligands were searched and ranked based on their re-rank score. For each docking calculation the maximum number of iterations for the docking algorithm was set to 1500, with a maximum population size of 50, and 30 runs per ligand. The RMSD threshold for multiple poses was set to 1.00 Å. The generated poses from each ligand were sorted by the calculated re-rank score. In analyzing the docking scores, we have attempted to account for the recognized bias toward high molecular weight compounds [34–37]. We have used two different schemes to re-evaluate the Molegro docking scores ( $E_{\text{dock}}$ ): (1)  $E' = 6.96 \times E_{\text{dock}} / \text{MW}^{1/3}$ , which accounts for biasing by dividing by a molecular weight function (MW), where 6.96 is a scaling con-

**Table 1**  
Homology models for Zika virus non-structural proteins.

Protein	GenBank Accession Number	Reference Structure		
		PDB Entry	Sequence Identity	E-value
NS3 Protease	AHL16750.1	2WV9 <sup>a</sup>	72%	$5 \times 10^{-135}$
NS5 MTase	A0A0B4ZYV2	4K6M	72%	$2 \times 10^{-157}$
NS5 RdRp	A0A0B4ZYV2	4K6M	72%	$2 \times 10^{-157}$
NS5 RdRp	A0A0B4ZYV2	2HFZ	73%	$3 \times 10^{-171}$

<sup>a</sup> Homology model was constructed with superposed ligand of PDB 2FP7.**Fig. 1.** Ligand-receptor interaction map of Bz-Nle-Lys-Arg-Arg-H with the West Nile virus NS2B-NS3 protease (PDB 2FP7) (panel A) and homology-modeled Zika virus NS2B-NS3 protease (panel B).

stant to bring the average  $E'$  values comparable to  $E_{\text{dock}}$ ; and (2)  $E'' = E_{\text{dock}} - 18.3 - E_{\text{dock}} \times \text{NH}/125$ , which reduces the bias by subtracting a contribution from the number of non-hydrogen atoms (NH), where 18.3 is a factor to bring average  $E'$  values comparable to  $E_{\text{dock}}$ . Histograms of docking scores for each protein target are available as Supplementary material.

### 3. Results and discussion

A total of 2263 plant-derived natural products have been docked with Zika virus NS2B-NS3 protease, NS3 helicase, NS5 methyltransferase, and NS5 RNA-dependent RNA polymerase. Selected phytochemical ligands with notable docking profiles and that adhere to Lipinski's rule-of-five for drug likeness [38] are listed in Table 2.

#### 3.1. Zika virus NS2B-NS3 protease

The homology modeled ZIKV NS2B-NS3 protease is a small (148 residues) protein. Most low-molecular-weight ligands docked into a small hydrophobic binding site bounded by Gln1572, Thr1664, Asn1650, Lys1571, and Ile1621. Phytochemical ligands with good docking scores with ZIKV NS2B-NS3 protease are listed in Table 2. Only two alkaloids obeyed Lipinski's rule [38] and showed notable

docking to ZIKV protease, the bis-indole alkaloids flinderole A and flinderole B. The flinderoles docked with ZIKV protease such that one of the indole moieties occupied the hydrophobic binding site, while the other indole, along with the amide side chain, was placed near the active site histidine (His1549) (Fig. 2).

The chalcones angusticorin B and kuraridin showed promising docking properties with ZIKV NS2B-NS3 protease (Table 2). The lowest-energy pose of the bis-hydroxyphenylated chalcone angusticorin B placed the hydroxyphenylated B-ring into the hydrophobic cavity, with none of the ligand occupying the active site of the protease Ser1633 and His1549 (Fig. 3A). Kuraridin preferentially docked to ZIKV NS2B-NS3 protease placing the diprenyl-substituted A-ring into the hydrophobic cavity and the B-ring also not occupying the active site (Fig. 3B). Although the ligands in Table 2 did show relatively strong docking to ZIKV NS2B-NS3 protease, the docking energies are less than those for other ZIKV target proteins, likely a consequence of the small ( $42 \text{ \AA}^3$ ) binding pocket in the protease. Thus, it is unlikely that these ligands will exhibit selective protease inhibition.

#### 3.2. Zika virus NS3 helicase

The X-ray crystal structure of ZIKV helicase has recently been determined [23]. There are two binding sites for flavivirus NS3

**Table 2**  
Molegro docking scores (kJ/mol) and normalized scores for phytochemical ligands with Zika virus non-structural proteins.

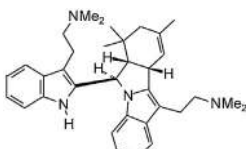
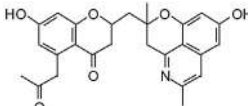
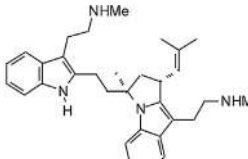
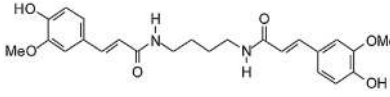
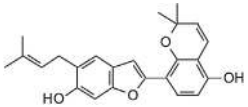
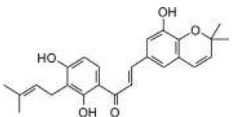
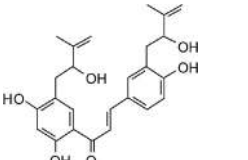
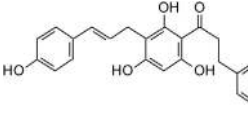
Phytochemical Ligand(MW, NH, HBD, HBA, ClogP) <sup>a</sup>	ZIKV protein	E <sub>dock</sub> <sup>b</sup>	E <sup>c</sup>	E <sup>d</sup>	Phytochemical and Bioactivity Notes and References
<b>ALKALOIDS</b>					
 4,4'-Dimethylisoborreverine (508.74, 38, 1, 4, 4.30)	Protease	-85.3	-74.3	-77.6	Isolated from <i>Flindersia fournieri</i> [39]. Isolated from <i>Flindersia amboinensis</i> ; shows antimalarial activity [40].
	Helicase (RNA site)	-144.2	-125.7	-118.6	
	Helicase (ATP site)	-87.7	-76.5	-79.4	
	MTase	-128.1	-111.7	-107.5	
	RdRp	-116.6	-101.7	-99.5	
 Cassiarin D (445.47, 33, 2, 7, 2.14)	Protease	-86.4	-78.7	-81.9	Isolated from the flowers of <i>Cassia siamea</i> ; shows antiplasmodial activity [41].
	Helicase (RNA site)	-117.2	-106.8	-104.5	
	Helicase (ATP site)	-150.7	-137.3	-106.4	
	MTase	-109.7	-100.0	-92.5	
	RdRp	-104.5	-95.2	-88.1	
 Flinderole A (480.69, 36, 3, 4, 1.67)	Protease	-103.7	-92.1	-92.1	Isolated from <i>Flindersia acuminata</i> ; shows antimalarial activity [40].
	Helicase (RNA site)	-131.5	-116.8	-111.9	
	Helicase (ATP site)	-139.1	-123.6	-117.4	
	MTase	-125.5	-111.5	-107.7	
	RdRp	-127.8	-113.5	-109.3	
 Terrestrisamide (440.49, 32, 4, 8, 2.67)	Protease	-77.6	-71.0	-76.1	From the fruit of <i>Tribulus terrestris</i> [42].
	Helicase (RNA site)	-140.3	-128.3	-122.6	
	Helicase (ATP site)	-116.7	-106.8	-105.2	
	MTase	-151.3	-138.4	-130.8	
	RdRp	-119.2	-109.0	-107.0	
<b>AURONES</b>					
 Kanzonol V (376.45, 28, 2, 4, 4.47)	Protease	-83.0	-80.0	-82.7	Isolated from licorice ( <i>Glycyrrhiza glabra</i> ) roots [43].
	Helicase (RNA site)	-110.4	-106.4	-103.9	
	Helicase (ATP site)	-133.6	-128.8	-122.0	
	MTase	-127.1	-122.5	-117.0	
	RdRp	-105.0	-101.2	-99.8	
<b>CHALCONES</b>					
 1-[2,4-Dihydroxy-3-(3-methyl-2-butenyl)phenyl]-3-(8-hydroxy-2,2-dimethyl-2H-1-benzopyran-6-yl)-2-propen-1-one (406.47, 30, 3, 5, 4.81)	Protease	-83.2	-78.2	-81.5	From licorice ( <i>Glycyrrhiza glabra</i> ) root; inhibitor of influenza virus neuraminidase [44].
	Helicase (RNA site)	-123.3	-115.9	-112.0	
	Helicase (ATP site)	-141.9	-133.3	-126.2	
	MTase	-125.6	-118.0	-113.7	
	RdRp	-116.2	-109.2	-106.6	
 Angusticornin B (424.49, 31, 5, 6, 3.70)	Protease	-99.1	-91.8	-92.8	Isolated from <i>Dorstenia angusticornis</i> ; shows antibacterial activity [45].
	Helicase (RNA site)	-124.5	-115.3	-112.0	
	Helicase (ATP site)	-149.6	-138.6	-130.8	
	MTase	-120.8	-111.9	-109.1	
	RdRp	-114.4	-106.0	-104.3	
 Balsaconin B (420.45, 31, 4, 6, 3.70)	Protease	-85.4	-79.3	-82.5	Isolated from the buds of <i>Populus balsamifera</i> ; shows antibacterial activity [46].
	Helicase (RNA site)	-123.1	-114.4	-110.9	
	Helicase (ATP site)	-143.9	-133.7	-126.5	
	MTase	-120.7	-112.2	-109.1	
	RdRp	-111.8	-103.9	-102.4	



Table 2 (Continued)

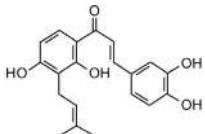
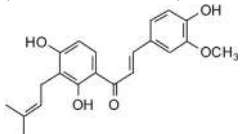
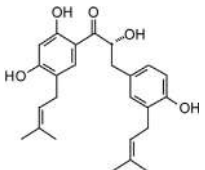
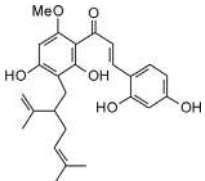
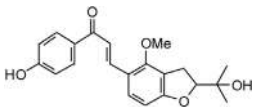
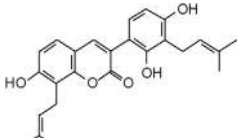
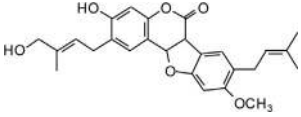
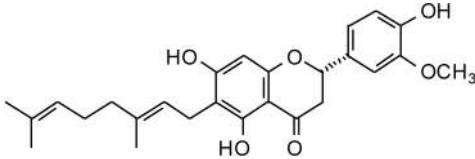
Phytochemical Ligand(MW, NH, HBD, HBA, ClogP) <sup>a</sup>	ZIKV protein	E <sub>dock</sub> <sup>b</sup>	E <sup>c</sup>	E <sup>rd</sup>	Phytochemical and Bioactivity Notes and References
 Corylifol B (340.37, 25, 4, 5, 3.80)	Protease	-83.6	-83.3	-85.2	Isolated from <i>Psoralea corylifolia</i> ; shows antibacterial activity [47].
	Helicase (RNA site)	-109.9	-109.5	-106.2	
	Helicase (ATP site)	-132.1	-131.7	-124.0	
	MTase	-109.9	-109.6	-106.2	
	RdRp	-110.5	-110.2	-106.7	
 Crotaorixin (354.40, 26, 3, 5, 4.06)	Protease	-90.2	-88.7	-89.7	Isolated from the aerial parts of <i>Crotalaria orixensis</i> ; shows antiplasmodial activity [48].
	Helicase (RNA site)	-120.4	-118.5	-113.7	
	Helicase (ATP site)	-135.9	-133.6	-125.9	
	MTase	-120.2	-118.2	-113.5	
	RdRp	-100.4	-98.7	-97.8	
 Kanzonol Y (410.50, 30, 4, 5, 4.48)	Protease	-95.6	-89.5	-90.9	Isolated from licorice ( <i>Glycyrrhiza glabra</i> ) roots [43].
	Helicase (RNA site)	-120.4	-112.8	-109.8	
	Helicase (ATP site)	-143.3	-134.2	-127.2	
	MTase	-128.8	-120.6	-116.2	
	RdRp	-111.9	-104.8	-103.3	
 Kuraridin (438.51, 32, 4, 6, 5.16)	Protease	-100.6	-92.2	-93.2	Isolated from <i>Sophora angustifolia</i> [49] and <i>Sophora flavescens</i> ; shows antibacterial [50] and tyrosinase inhibitory [51], and anti-reoviral [52] activities.
	Helicase (RNA site)	-127.0	-116.4	-112.8	
	Helicase (ATP site)	-141.1	-129.2	-123.2	
	MTase	-134.4	-123.1	-118.3	
	RdRp	-115.1	-105.5	-103.9	
 Licoagrochalcone D (354.40, 26, 2, 5, 3.09)	Protease	-74.5	-73.3	-77.3	Constituent of hairy root cultures of <i>Glycyrrhiza glabra</i> [53].
	Helicase (RNA site)	-111.5	-109.7	-106.6	
	Helicase (ATP site)	-125.6	-123.5	-117.8	
	MTase	-103.2	-101.5	-100.0	
	RdRp	-111.7	-109.8	-106.8	
COUMARINS  Licocoumarin A (406.47, 30, 3, 5, 5.39)	Protease	-74.5	-70.0	-74.9	Isolated from <i>Glycyrrhiza glabra</i> roots [54].
	Helicase (RNA site)	-108.9	-102.4	-101.1	
	Helicase (ATP site)	-140.1	-131.6	-124.8	
	MTase	-124.4	-116.9	-112.9	
	RdRp	-108.5	-101.9	-100.8	
 3-Hydroxy-9-methoxy-2-[2'-(E)-4'-hydroxy-3'-methylbutenyl]-8-isoprenylcoumestan (434.48, 32, 2, 6, 4.41)	Protease	-87.5	-80.4	-83.4	Isolated from the roots of <i>Picalima nitida</i> ; shows antibacterial activity [55].
	Helicase (RNA site)	-121.7	-111.8	-108.8	
	Helicase (ATP site)	-131.2	-120.5	-115.9	
	MTase	-129.3	-118.9	-114.5	
	RdRp	-109.4	-100.5	-99.7	
FLAVONOIDS  3'-O-Methyldiplacone (438.51, 32, 3, 6, 4.62)	Protease	-85.5	-78.3	-81.9	Isolated from the fruits of <i>Poulownia tomentosa</i> ; shows antibacterial [56] and antiprotozoal [57] activities.
	Helicase (RNA site)	-127.4	-116.7	-113.1	
	Helicase (ATP site)	-155.9	-142.8	-134.3	
	MTase	-118.8	-108.9	-106.7	
	RdRp	-122.2	-112.0	-109.2	

Table 2 (Continued)

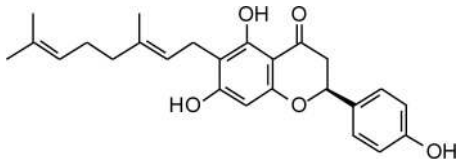
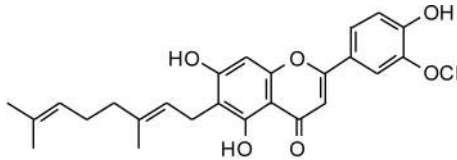
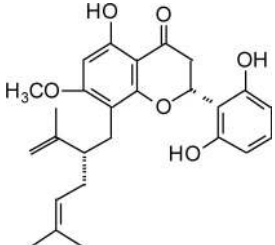
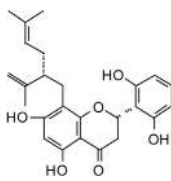
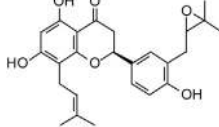
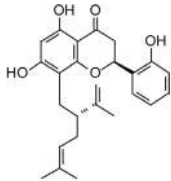
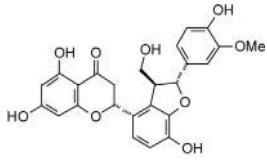
Phytochemical Ligand(MW, NH, HBD, HBA, ClogP) <sup>a</sup>	ZIKV protein	E <sub>dock</sub> <sup>b</sup>	E <sup>c</sup>	E <sup>rd</sup>	Phytochemical and Bioactivity Notes and References
 <p>Bonannione A (408.49, 30, 3, 5, 4.75)</p>	Protease Helicase (RNA site) Helicase (ATP site) MTase RdRp	-80.8 -121.3 -134.6 -119.0 -115.4	-75.8 -113.8 -126.2 -111.7 -108.2	-79.7 -110.5 -120.6 -108.8 -106.0	Isolated from the fruits of <i>Paulownia tomentosa</i> ; shows cytotoxic activity [58].
 <p>Cannflavin A (436.50, 32, 3, 6, 4.39)</p>	Protease Helicase (RNA site) Helicase (ATP site) MTase RdRp	-92.1 -131.7 -139.8 -126.9 -120.3	-84.5 -120.8 -128.1 -116.5 -110.4	-86.8 -116.3 -122.2 -112.7 -107.8	Isolated from <i>Cannabis sativa</i> [59]; shows antiprotozoal activity [57].
 <p>Exiguaflavanone A (438.51, 32, 3, 6, 4.59)</p>	Protease Helicase (RNA site) Helicase (ATP site) MTase RdRp	-91.7 -115.0 -156.1 -130.4 -103.9	-84.0 -105.3 -143.0 -119.5 -95.2	-86.5 -103.8 -134.4 -115.3 -95.6	Isolated from <i>Artemisia indica</i> ; shows antimalarial activity [60].
 <p>Exiguaflavanone B (424.49, 31, 4, 6, 4.32)</p>	Protease Helicase (RNA site) Helicase (ATP site) MTase RdRp	-82.6 -107.9 -134.5 -125.2 -102.4	-76.5 -99.9 -124.5 -116.0 -94.8	-80.4 -99.5 -119.4 -112.5 -95.3	Isolated from <i>Artemisia indica</i> ; shows antimalarial activity [60].
 <p>Flemiflavanone D (424.49, 31, 3, 6, 3.56)</p>	Protease Helicase (RNA site) Helicase (ATP site) MTase RdRp	-82.9 -123.4 -140.5 -121.3 -117.3	-76.8 -114.2 -130.1 -112.3 -108.6	-80.6 -111.1 -124.0 -109.5 -106.5	Isolated from <i>Flemingia stricta</i> ; shows antibacterial activity [61].
 <p>Kushenol A (408.49, 30, 3, 5, 4.71)</p>	Protease Helicase (RNA site) Helicase (ATP site) MTase RdRp	-79.1 -125.4 -137.0 -128.9 -104.2	-74.2 -117.6 -128.6 -120.9 -97.7	-78.4 -113.6 -122.5 -116.2 -97.5	Isolated from the roots of <i>Sophora flavescens</i> ; shows antibacterial activity [62].
 <p>Neosilyhermin A (466.44, 34, 5, 9, 2.02)</p>	Protease Helicase (RNA site) Helicase (ATP site) MTase RdRp	-82.1 -137.9 -136.6 -133.1 -109.2	-73.7 -123.7 -122.6 -119.4 -98.0	-78.1 -118.7 -117.7 -115.2 -97.8	A constituent of milk thistle ( <i>Silybum marianum</i> ) [63].

Table 2 (Continued)

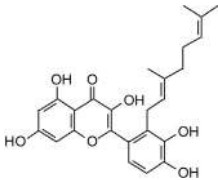
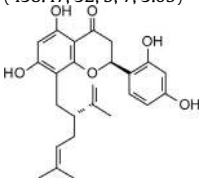
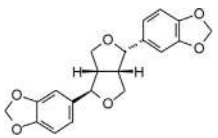
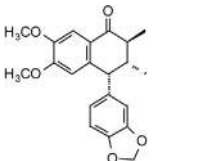
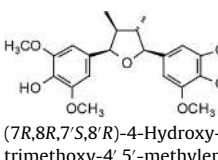
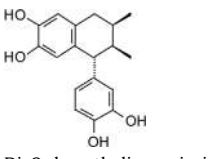
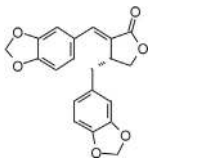
Phytochemical Ligand(MW, NH, HBD, HBA, ClogP) <sup>a</sup>	ZIKV protein	E <sub>dock</sub> <sup>b</sup>	E <sup>c</sup>	E <sup>d</sup>	Phytochemical and Bioactivity Notes and References
 Solophenol D (438.47, 32, 5, 7, 3.05)	Protease Helicase (RNA site) Helicase (ATP site) MTase RdRp	-110.5 -122.4 -139.9 -134.0 -104.4	-101.2 -112.2 -128.2 -122.8 -95.6	-100.5 -109.4 -122.4 -118.0 -96.0	Isolated from propolis collected from the Solomon Islands; shows antibacterial activity [64].
 Sophoraflavanone G (424.49, 31, 4, 6, 5.10)	Protease Helicase (RNA site) Helicase (ATP site) MTase RdRp	-80.8 -109.8 -113.7 -127.9 -104.9	-74.8 -101.6 -108.1 -118.5 -97.2	-79.0 -100.8 -105.6 -114.5 -97.2	Isolated from the roots of <i>Sophora flavescens</i> ; shows antibacterial activity [65].
LIGNANS  (-)-Asarinin (354.35, 26, 0, 6, 2.70)	Protease Helicase (RNA site) Helicase (ATP site) MTase RdRp	-75.1 -109.2 -123.0 -126.4 -103.8	-73.9 -107.4 -121.0 -124.3 -102.1	-77.8 -104.8 -115.7 -118.4 -100.5	Found in several species of <i>Asarum</i> , <i>Boronia</i> , and <i>Zanthoxylum</i> [54].
 (7 <i>R</i> , 8 <i>S</i> , 8' <i>R</i> )-4,5-Dimethoxy-3',4'-methyleneedioxy-2,7'-cyclo lignan-7-one (= Aristotetralone) (354.40, 26, 0, 5, 3.93)	Protease Helicase (RNA site) Helicase (ATP site) MTase RdRp	-66.0 -100.5 -118.5 -93.4 -115.5	-64.9 -98.9 -116.5 -91.8 -113.6	-70.6 -97.9 -112.1 -92.3 -109.8	Isolated from <i>Aristolochia chilensis</i> ; shows antiparasitodal activity [66].
 (7 <i>R</i> , 8 <i>R</i> , 7' <i>S</i> , 8' <i>R</i> )-4-Hydroxy-3,3',5'-trimethoxy-4',5'-methyleneedioxy-7,7'-epoxylignan (402.44, 29, 1, 7, 3.64)	Protease Helicase (RNA site) Helicase (ATP site) MTase RdRp	-95.9 -103.6 -126.4 -109.1 -114.5	-90.4 -97.7 -119.1 -102.8 -108.0	-91.9 -97.9 -115.4 -102.1 -106.3	Isolated from <i>Peperomia blanda</i> ; shows antitrypanosomal activity [67].
 Di- <i>O</i> -demethylisoguaiacin (300.35, 22, 4, 4, 3.96)	Protease Helicase (RNA site) Helicase (ATP site) MTase RdRp	-65.2 -77.0 -103.8 -89.6 -114.4	-67.8 -80/0 -107.9 -93.2 -118.9	-72.0 -81.7 -103.8 -92.2 -112.6	Isolated from <i>Larrea tridentata</i> ; shows antiprotozoal activity [68].
 Hibalactone (= Savinin) (352.34, 26, 0, 6, 3.60)	Protease Helicase (RNA site) Helicase (ATP site) MTase RdRp	-76.5 -116.6 -148.3 -114.7 -103.1	-75.4 -114.9 -146.2 -113.1 -101.6	-78.9 -110.7 -135.8 -109.2 -99.9	A constituent of the needles of <i>Juniperus sabina</i> and <i>Juniperus converta</i> , the bark of <i>Chloroxylon swietenia</i> and <i>Zanthoxylum nitidum</i> [54]. Shows antiviral activity against severe acute respiratory syndrome associated coronavirus (SARS-CoV) [69].



Table 2 (Continued)

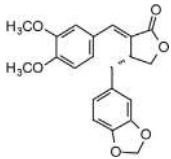
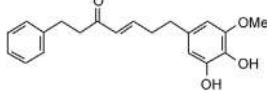
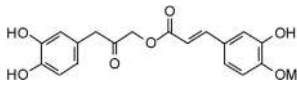
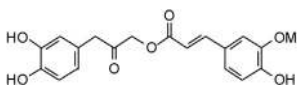
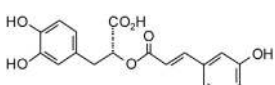
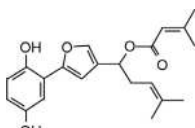
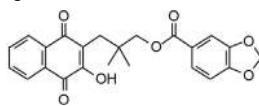
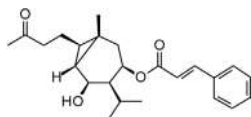
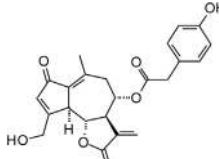
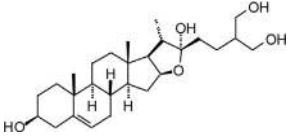
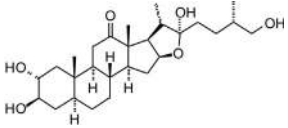
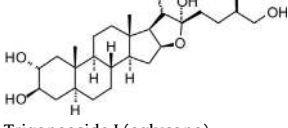
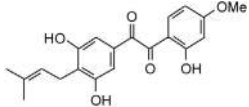
Phytochemical Ligand(MW, NH, HBD, HBA, ClogP) <sup>a</sup>	ZIKV protein	E <sub>dock</sub> <sup>b</sup>	E <sup>c</sup>	E <sup>rd</sup>	Phytochemical and Bioactivity Notes and References
 <p>Kaerophyllin (368.38, 27, 0, 6, 3.57)</p>	Protease	-76.5	-74.3	-78.3	Isolated from the roots of <i>Chaerophyllum maculatum</i> [70].
	Helicase (RNA site)	-115.6	-112.3	-109.0	
	Helicase (ATP site)	-145.8	-141.5	-132.6	
	MTase	-103.2	-100.2	-99.2	
	RdRp	-111.5	-108.3	-105.7	
<p>PHENOLICS</p>  <p>7-(3,4-Dihydroxy-5-methoxyphenyl)-1-phenyl-4-hepten-3-one (326.39, 24, 2, 4, 4.80)</p>	Protease	-67.6	-68.3	-72.9	A constituent of the rhizomes of <i>Alpinia officinarum</i> ; shows antibacterial activity [71].
	Helicase (RNA site)	-108.5	-109.7	-106.0	
	Helicase (ATP site)	-124.3	-125.7	-118.7	
	MTase	-118.7	-120.0	-114.2	
	RdRp	-97.2	-98.3	-96.9	
 <p>Cimicifhenol (358.34, 26, 3, 7, 2.69)</p>	Protease	-88.1	-86.3	-88.1	Found in the rhizome of black cohosh, <i>Actaea racemosa</i> (syn. <i>Cimicifuga racemosa</i> ) [72].
	Helicase (RNA site)	-121.7	-119.2	-115.9	
	Helicase (ATP site)	-123.3	-120.8	-100.2	
	MTase	-124.8	-122.3	-117.1	
	RdRp	-99.9	-97.9	-97.4	
 <p>Cimiracemate B (358.34, 26, 3, 7, 2.69)</p>	Protease	-91.7	-89.8	-90.9	Found in the rhizome of black cohosh, <i>Actaea racemosa</i> (syn. <i>Cimicifuga racemosa</i> ) [73].
	Helicase (RNA site)	-120.5	-118.0	-113.7	
	Helicase (ATP site)	-139.0	-136.2	-128.3	
	MTase	-127.2	-124.7	-119.1	
	RdRp	-107.0	-104.9	-103.1	
 <p>Rosmarinic acid (360.32, 26, 5, 8, 2.24)</p>	Protease	-86.8	-84.9	-87.0	Found in rosemary ( <i>Rosmarinus officinalis</i> ), lemon balm ( <i>Melissa officinalis</i> ), common sage ( <i>Salvia officinalis</i> ), and many other plant species [54]; shows antiviral activity [74].
	Helicase (RNA site)	-108.2	-105.9	-104.0	
	Helicase (ATP site)	-126.3	-123.6	-118.3	
	MTase	-123.5	-120.8	-116.1	
	RdRp	-96.9	-94.8	-95.0	
 <p>Shikonofuran E (356.41, 26, 2, 5, 3.14)</p>	Protease	-95.4	-93.7	-93.9	Isolated from the roots of <i>Lithospermum erythrorhizon</i> [75].
	Helicase (RNA site)	-113.0	-110.9	-107.8	
	Helicase (ATP site)	-135.8	-133.3	-125.9	
	MTase	-115.0	-112.9	-109.4	
	RdRp	-106.6	-104.6	-102.7	
<p>QUINONES</p>  <p>Rhinacanthin D (408.40, 30, 1, 7, 2.83)</p>	Protease	-98.9	-92.8	-93.4	Isolated from <i>Rhinacanthus nasutus</i> ; shows antiviral activity [76].
	Helicase (RNA site)	-118.0	-110.7	-108.0	
	Helicase (ATP site)	-136.6	-128.2	-122.1	
	MTase	-114.8	-107.7	-105.6	
	RdRp	-96.3	-90.4	-91.5	
<p>SESQUITERPENOIDS</p>  <p>Cinnamoylchinoxanthol (384.51, 28, 1, 4, 4.53)</p>	Protease	-76.8	-73.6	-77.9	Isolated from the roots of <i>Echinacea purpurea</i> [77].
	Helicase (RNA site)	-115.4	-110.5	-107.9	
	Helicase (ATP site)	-123.1	-117.8	-113.8	
	MTase	-110.9	-106.1	-104.3	
	RdRp	-114.4	-109.5	-107.1	
 <p>Lactucopicrin (410.42, 30, 2, 7, 1.63)</p>	Protease	-78.9	-73.9	-78.3	Isolated from <i>Lactuca</i> spp. and <i>Cichorium</i> spp. [54]. Shows antiplasmodial activity [78].
	Helicase (RNA site)	-120.8	-113.1	-110.1	
	Helicase (ATP site)	-153.3	-143.6	-134.8	
	MTase	-103.6	-97.0	-97.0	
	RdRp	-109.3	-102.4	-101.4	

Table 2 (Continued)

Phytochemical Ligand(MW, NH, HBD, HBA, ClogP) <sup>a</sup>	ZIKV protein	E <sub>dock</sub> <sup>b</sup>	E' <sup>c</sup>	E'' <sup>d</sup>	Phytochemical and Bioactivity Notes and References
<b>STEROIDS</b>					
 Furost-5-ene-3,22,26,27-tetrol (= Hypoglaucin F aglycone) (448.64, 32, 4, 5, 3.48)	Protease	-34.2	-31.1	-43.7	Isolated from the rhizomes of <i>Dioscorea collettii</i> [79].
	Helicase (RNA site)	-105.3	-95.7	-96.7	
	Helicase (ATP site)	-106.8	-97.1	-97.8	
	MTase	-112.9	-102.7	-102.3	
	RdRp	-118.1	-107.4	-106.2	
 Tribufuroside I (aglycone) (464.64, 33, 4, 6, 3.45)	Protease	-71.8	-64.5	-71.1	Isolated from the fruits of <i>Tribulus terrestris</i> [80].
	Helicase (RNA site)	-109.6	-98.5	-99.0	
	Helicase (ATP site)	-106.3	-95.5	-96.5	
	MTase	-110.7	-99.5	-99.8	
	RdRp	-124.6	-111.9	-110.0	
 Trigoneoside I (aglycone) (450.65, 32, 4, 5, 4.06)	Protease	-75.7	-68.7	-74.6	Isolated from the seeds of fenugreek ( <i>Trigonella foenum-graecum</i> ) [81].
	Helicase (RNA site)	-102.6	-93.2	-94.7	
	Helicase (ATP site)	-116.1	-105.4	-104.7	
	MTase	-98.0	-89.0	-91.2	
	RdRp	-125.5	-113.9	-111.6	
<b>STILBENOIDS</b>					
 Licoagrodione (356.37, 26, 3, 6, 3.08)	Protease	-87.5	-85.9	-87.6	Isolated from <i>Glycyrrhiza glabra</i> hairy root cultures; shows antimicrobial activity [82].
	Helicase (RNA site)	-115.8	-113.7	-110.0	
	Helicase (ATP site)	-128.7	-126.4	-120.3	
	MTase	-113.3	-111.2	-108.1	
	RdRp	-111.2	-109.2	-106.4	

<sup>a</sup> MW = molecular weight (g/mol); NH = number of non-hydrogen atoms; HBD = number of hydrogen-bond-donating atoms; HBA = number of hydrogen-bond-accepting atoms; ClogP = calculated octanol/water partition coefficient.

<sup>b</sup> E<sub>dock</sub> = Molegro "rerank" docking scores (kJ/mol).

<sup>c</sup> E' = Normalized docking score:  $E' = 6.96 \times E_{\text{dock}} / \text{MW}^{1/3}$ .

<sup>d</sup> E'' = Alternative normalized docking score:  $E'' = E_{\text{dock}} - 18.3 - E_{\text{dock}} \times \text{NH} / 125$ .

helicases, an RNA binding site and an ATP binding site [83,84]. Most phytochemical ligands showed more exothermic docking energies with the ATP site over the RNA site of ZIKV NS3 helicase. For example, the isoquinoline alkaloid cassiarin D showed very strong docking to the ATP site (E<sub>dock</sub> = -150.7 kJ/mol). In addition, cassiarin D showed some docking selectivity toward the ATP site of helicase (Fig. 4A). Similarly, the flavonoids 3'-O-methyl-diplacone and exiguafavanone A, and the sesquiterpenoid lactucopicrin docked selectively with the ATP site of helicase. Several other phytochemical ligands (e.g., the aurones kanzonol V, the chalcones angusticornin B, balsacone B, and kanzonol Y, and the lignans hibactone and kaerophyllin) also showed remarkably strong interactions with the ATP site of helicase. Interestingly, the bis-indole alkaloid 4,4'-dimethylisoborverine was selective for the RNA site of ZIKV NS3 helicase (Fig. 4B). The intermolecular interactions between the ligands and the helicase protein are illustrated in Fig. 4.

### 3.3. Zika virus NS5 methyltransferase

Non-structural protein 5 is made up of two domains. The N-terminal domain contains the methyltransferase (MTase) domain, while the C-terminus contains the RNA-dependent RNA polymerase (RdRp) domain. The methyltransferase domain is comprised of two binding sites, the GTP binding site and the SAM

binding site [85]. The phytochemical ligands in this study docked preferentially to the GTP site rather than the SAM site.

The best docking ligands for ZIKV NS5 MTase from this study were polyphenolic compounds with at least two phenolic groups, preferably connected with a flexible linker. Thus, for example, cimiphenol, cimracemate B, and rosemarinic acid exhibited relatively strong docking energies. Prenyl substituents also decreased docking energies; the prenylated chalcones kanzonol Y and kuraridin showed excellent docking properties. Likewise, the prenylated aurone kanzonol V and the geranylated flavone solophenol D also showed strong docking to ZIKV MTase. Interestingly, the lignan (-)-asarinin showed good docking to MTase (E<sub>dock</sub> = -126.4 kJ/mol), but its epimer, (+)-sesamin, showed weaker docking (E<sub>dock</sub> = -101.1 kJ/mol). Key interactions between the phenolic ligands and the MTase binding site are Lys2619, Ile2661, Gly2959, Thr2618, and Glu2625 (see Fig. 5).

### 3.4. Zika virus NS5 RNA-dependent RNA polymerase

The ZIKV NS5 RdRp homology-modeled structure based on the Japanese encephalitis virus NS5 (PDB 4K6M) [30] has a very large binding cavity (~2852 Å<sup>3</sup>). There were four major docking sites of phytochemical ligands with this protein structure (see Fig. 6), site A, with major interactions between the docked ligands and Asp3054, Asp3179, Tyr3123, Arg2987, and Ser3177; site B, with major interacting residues Trp3311, Arg3245, Ser3312, and Ser3226; site C,

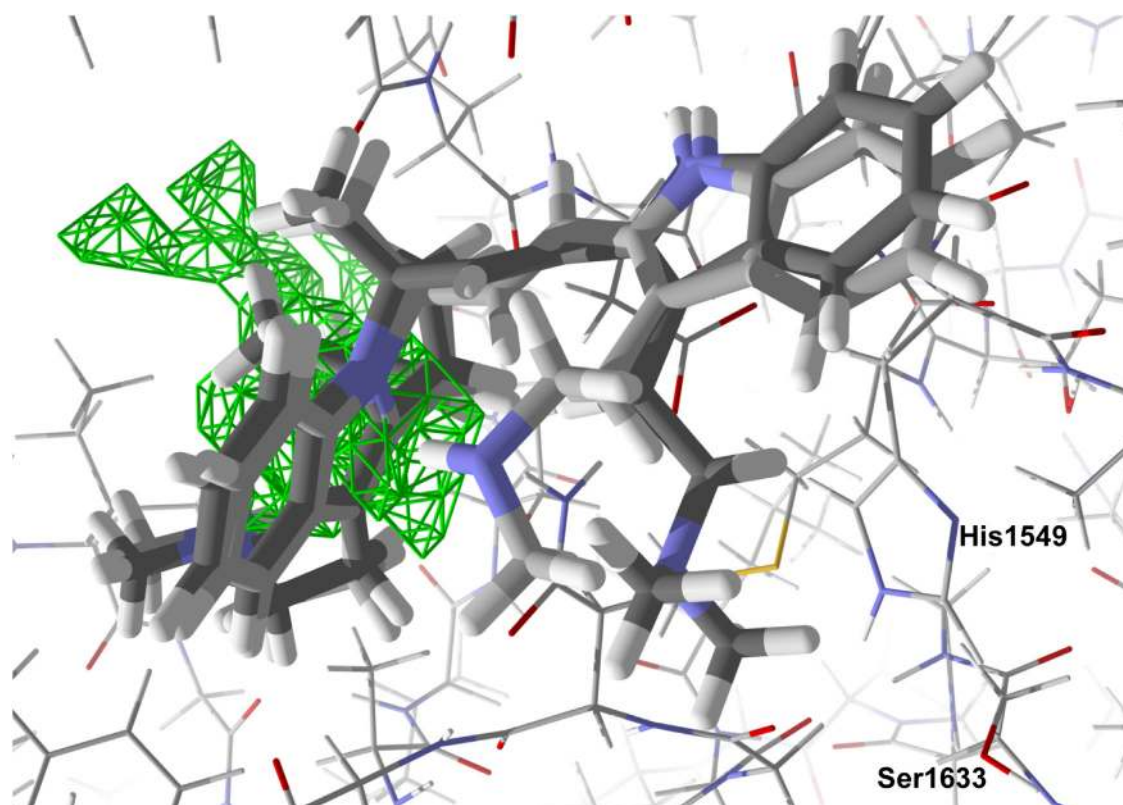


Fig. 2. Lowest-energy docked poses of flinderole A and B with ZIKV NS2B-NS3 protease. The hydrophobic cavity is shown with green hashmarks.

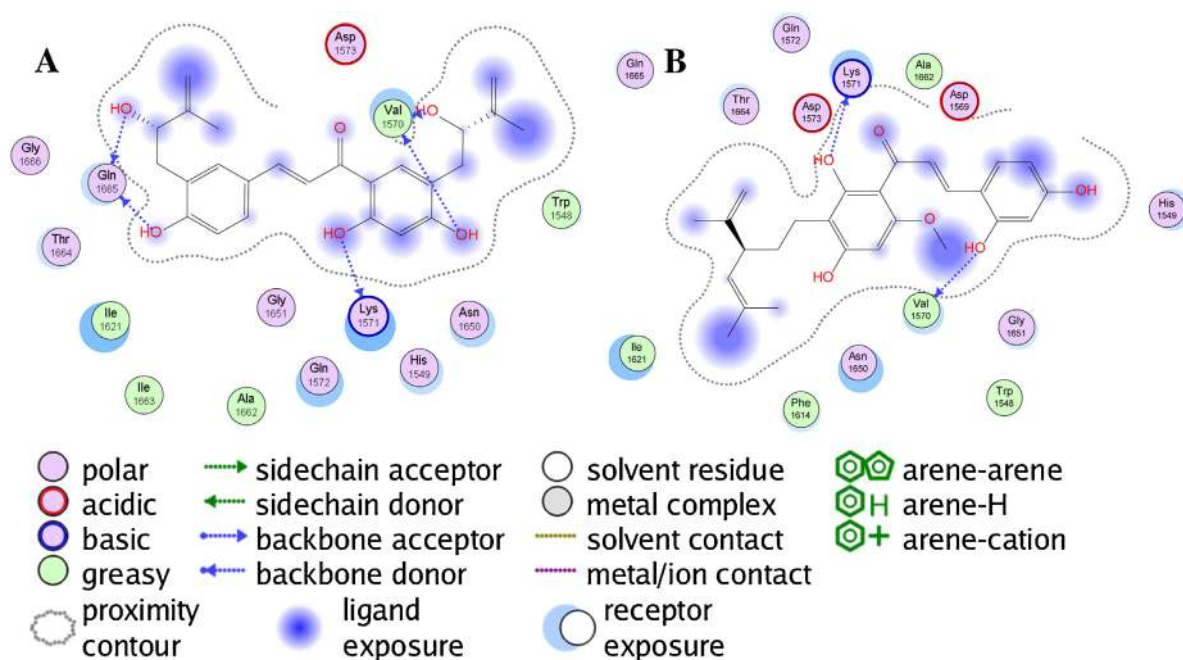


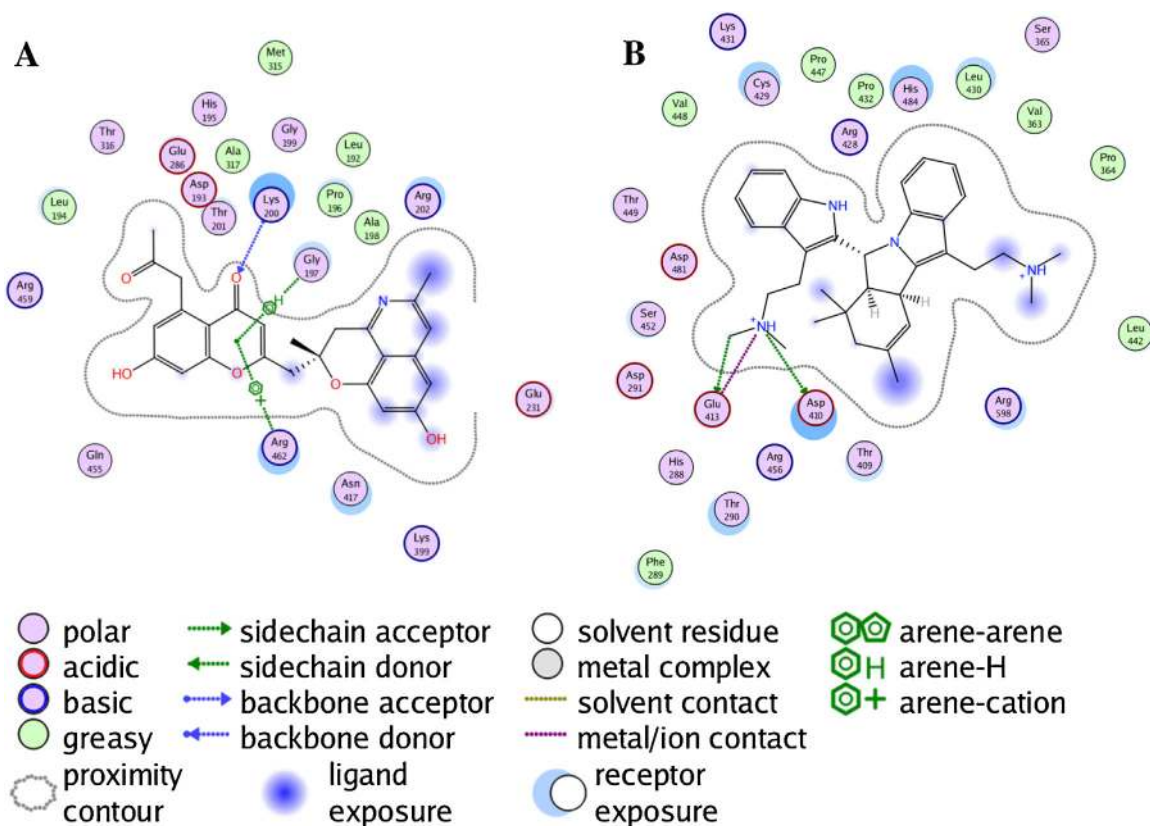
Fig. 3. Ligand-receptor interaction map of angusticin B (panel A) and kuraridin (panel B) with the Zika virus NS2B-NS3 protease.

with residues Arg3245, His3246, His3227, His3228, and Trp3362; and site **D**, bounded by residues Met2970, Leu3096, Pro2863, and Gln2829.

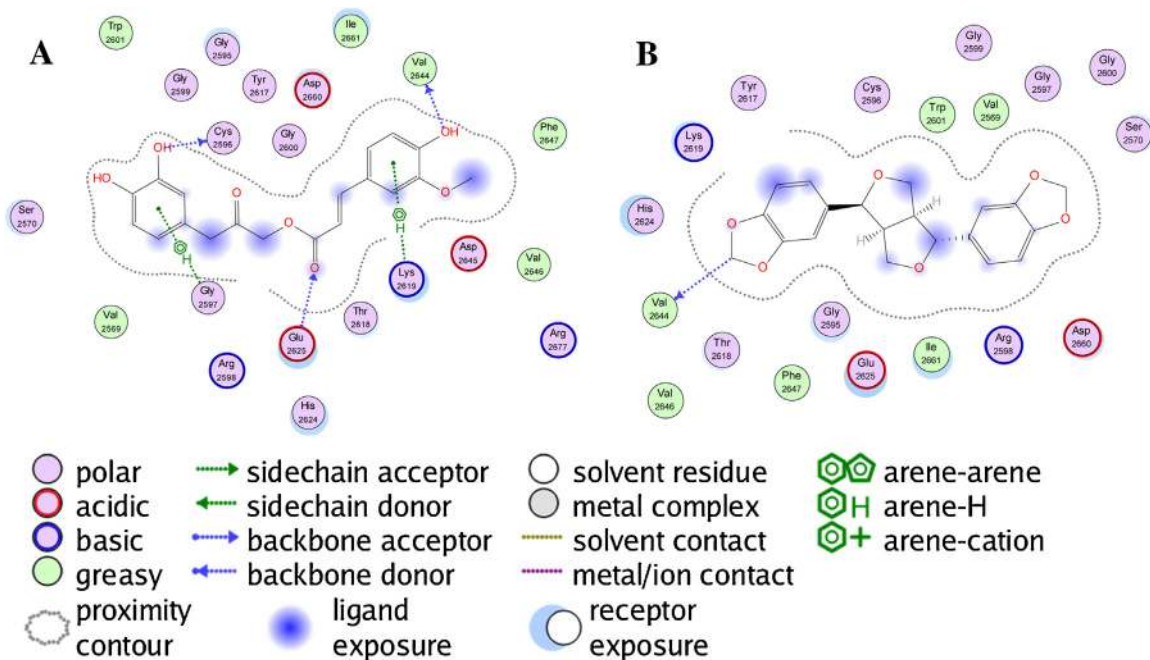
Small hydrophobic ligands such as the monoterpenoids and naphthoquinones docked preferentially in sites **C** (58%) and **D** (38%). Large hydrophobic ligands, the triterpenoids, on the other hand, tended to concentrate in sites **A** (20%), **B** (51%),

and **C** (18%). The best docking ligands to NIKV NS5 RdRp were aromatic ligands, particularly polyphenolics. The prenylated chalcone, 2',4,4'-trihydroxy-3,3'-diprenylchalcone, docked preferentially into site **A** (Fig. 7A) while the bis-indole alkaloid flinderole B docked into site **B** (Fig. 7B). The polyphenolic compound 4',7-digalloylcatechin docked into site **C** (Fig. 7C), and the lignan di-O-demethylisoguaiacin docked into site **D** (Fig. 7D).





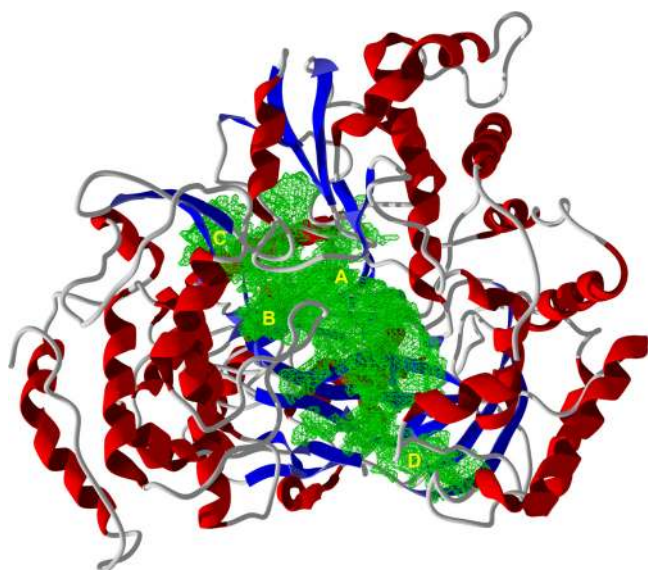
**Fig. 4.** Ligand-receptor interaction map of cassiarin D with the with the ATP binding site (panel A) and 4,4'-dimethylisoborverine (panel B) and with the RNA binding site of the Zika virus NS3 helicase (PDB 5JMT [23]).



**Fig. 5.** Protein-ligand interaction map between Zika virus NS5 methyltransferase and the lowest-energy docked pose of the phenolic ligand cimracemate B (panel A) and the lignan (-)-asarinin (panel B).

Notably, the homology model of ZIKV RNA-dependent RNA polymerase based on the West Nile virus template (PDB 2HFZ) has a much smaller binding cavity than the model based on the Japanese encephalitis virus NS5 template (PDB 4K6M), 302 Å<sup>3</sup> and 2852 Å<sup>3</sup>, respectively. Because of this, nearly all of the docking ener-

gies (2164 of the total 2263 ligands) were lower for the 4K6M model than the 2HFZ model; an average  $E_{\text{dock}}$  difference of 11.8 kJ/mol per ligand. It would appear, based on these data, that the homology model based on WNV RdRp is a poorer model than that based on JEV.

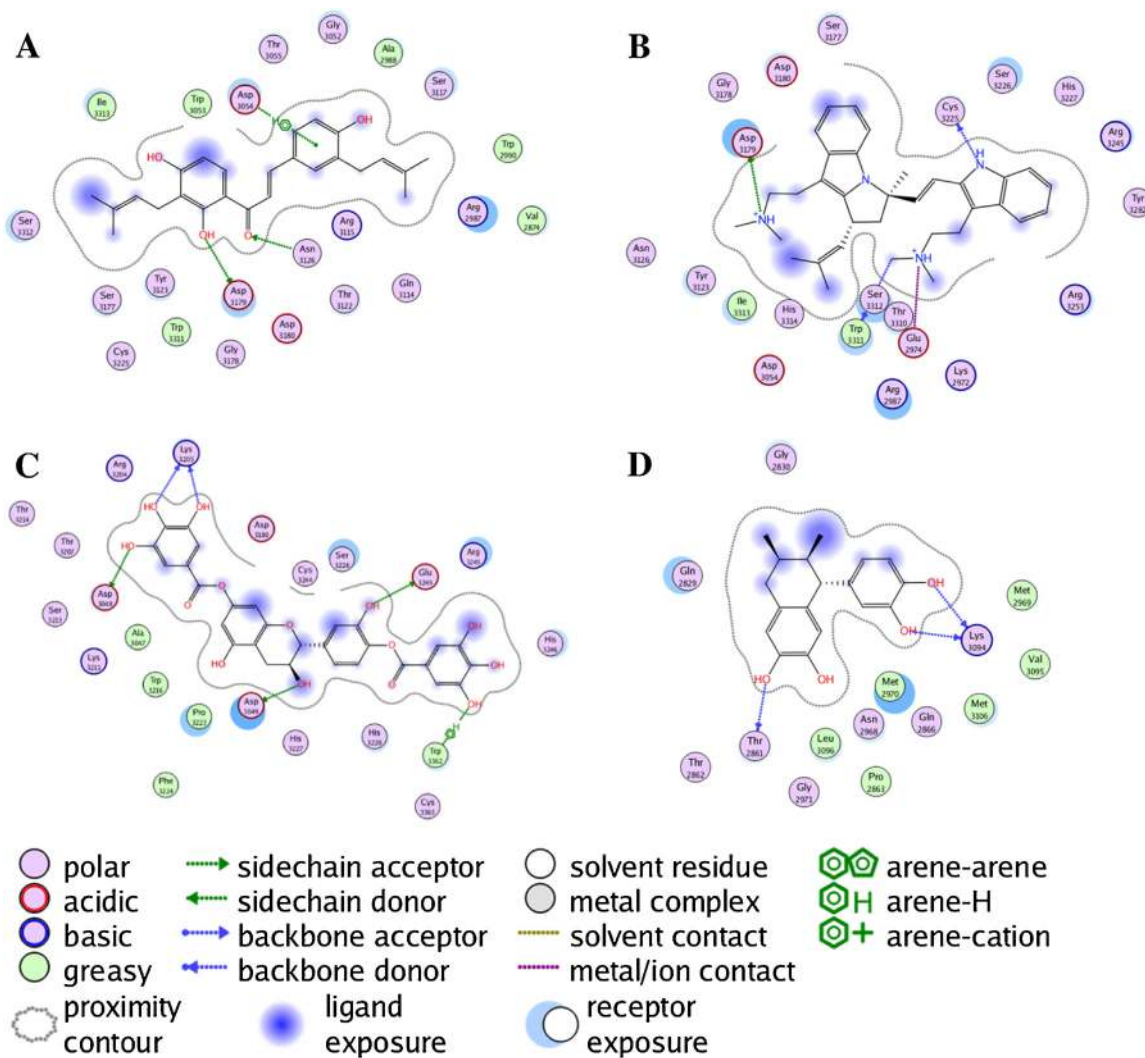


**Fig. 6.** Ribbon structure of Zika virus NS5 RNA-dependent RNA polymerase homology model based on the Japanese encephalitis virus NS5 (PDB 4K6M [30]). The binding cavity is shown in green with the preferred ligand docking sites, A–D.

#### 4. Conclusions

The best docking phytochemical ligands were the polyphenolics, which generally docked strongly to several of the protein targets. The poorest docking ligands were the terpenoids, in particular the monoterpenoids, owing to their small size, and the triterpenoids, likely due to their paucity of functional groups.

There are several phytochemical ligands that had shown good selective docking properties and that are found in relatively common medicinal herbs. For example, balsacone B is found in the buds of balsam poplar (*Populus balsamifera*), kanzonol V is found in licorice root (*Glycyrrhiza glabra*), cinnamoylechinoxanthol is found in *Echinacea* root; cimiphenol is found in black cohosh (*Actaea racemosa*, syn. *Cimicifuga racemosa*); and rosmarinic acid is found in several common herbs including rosemary (*Rosmarinus officinalis*), lemon balm (*Melissa officinalis*), and common sage (*Salvia officinalis*). Hence, there are several common medicinal plants that may serve as antiviral agents themselves or ready sources of potential antivirals. We hope that this study encourages research into antiviral screening of medicinal plants and phytochemicals to identify antiviral agents and lead structures for structure-based design for new chemotherapeutics to treat Zika viral infections as well as other neglected tropical diseases.



**Fig. 7.** Protein-ligand interaction map between Zika virus NS5 RNA-dependent RNA polymerase and the lowest-energy docked pose of the prenylated chalcone, 2',4,4'-trihydroxy-3,3'-diprenylchalcone (panel A), the bis-indole alkaloid flinderole B (panel B), the polyphenolic compound 4',7-digalloyl catechin (panel C), and the lignan di-O-demethylisoguaiacin (panel D).



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**Appendix A. Supplementary data**

Supplementary data associated with this article can be found, in the online version, at <http://dx.doi.org/10.1016/j.jmngm.2016.08.011>.

**References**

[1] P.Y. Shi, *Molecular Virology and Control of Flaviviruses*, Calister Academic Press, Norfolk, UK, 2012.

[2] G.W. Dick, S.F. Kitchen, A.J. Haddow, Zika virus. I. Isolations and serological specificity, *Trans. R. Soc. Trop. Med. Hyg.* 46 (1952) 509–520.

[3] E.B. Hayes, Zika virus outside Africa, *Emerg. Infect. Dis.* 15 (2009) 1347–1350.

[4] D. Musso, E.J. Nilles, and V.M. Cao-Lormeau: rapid spread of emerging Zika virus in the Pacific area, *Clin. Microbiol. Infect.* 20 (2014) O595–O596.

[5] G.S. Campos, A.C. Bandeira, S.I. Sardi, Zika virus outbreak, Bahia, Brazil, *Emerg. Infect. Dis.* 21 (2015) 1885–1886.

[6] A.J. Rodríguez-Morales, Zika: the new arbovirus threat for Latin America, *J. Infect. Dev. Ctries.* 9 (2015) 684–685.

[7] M. Hennessey, M. Fischer, J.E. Staples, Zika virus spreads to new areas –Region of the Americas, May 2015–January 2016, *Morb. Mortal. Wkly. Rep.* 65 (3) (2016) 55–58.

[8] D. Musso, C. Roche, E. Robin, T. Nhan, A. Teissier, V.M. Cao-Lormeau, Potential sexual transmission of Zika virus, *Emerg. Infect. Dis.* 21 (2015) 359–361.

[9] J.A. Tetro, Zika and microcephaly: causation, correlation, or coincidence? *Microbes Infect.* 18 (2016) 167–168.

[10] E. Oehler, L. Watrin, P. Larre, I. Leparc-Goffart, S. Lastère, F. Valour, L. Baudouin, H.P. Mallet, D. Musso, F. Ghawche, Zika virus infection complicated by Guillain-Barré syndrome –case report, French Polynesia, December 2013, *Euro Surveill.* 19 (9) (2014), ID = 20720.

[11] A.S. Oliveira Melo, G. Malinger, R. Ximenes, P.O. Szejnfeld, S.A. Sampaio, A.M. Bispo de Filippis, Zika virus intrauterine infection causes fetal brain abnormality and microcephaly: tip of the iceberg? *Ultrasound Obstet. Gynecol.* 47 (2016) 6–7.

[12] C.V. Ventura, M. Maia, V. Bravo-Filho, A.L. Góis, R. Belfort, Zika virus in Brazil and macular atrophy in a child with microcephaly, *Lancet* 387 (2016) 228.

[13] E.E. Petersen, J.E. Staples, D. Meaney-Delman, M. Fischer, S.R. Ellington, W.M. Callaghan, D.J. Javieson, Interim guidelines for pregnant women during a Zika virus outbreak –United States, 2016, *Morb. Mortal. Wkly. Rep.* 65 (2) (2016) 30–33.

[14] S.M. Tomlinson, R.D. Malmstrom, S.J. Watowich, New approaches to structure-based discovery of dengue protease inhibitors, *Infect. Disord. Drug Targ.* 9 (2009) 327–343.

[15] M.J. Stoermer, K.J. Chappell, S. Liebscher, C.M. Jensen, C.H. Gan, P.K. Gupta, W.J. Xu, P.R. Young, D.P. Fairlie, Potent cationic inhibitors of West Nile virus NS2B/NS3 protease with serum stability: cell permeability and antiviral activity, *J. Med. Chem.* 51 (2008) 5714–5721.

[16] H. Steuber, M. Kanitz, F.G.R. Ehler, W.E. Diederich, Recent advances in targeting dengue and West Nile virus proteases using small molecule inhibitors, *Top. Med. Chem.* 15 (2015) 93–142.

[17] D. Luo, S.G. Vasudevan, J. Lescar, The flavivirus NS2B-NS3 protease-helicase as a target for antiviral drug development, *Antiviral Res.* 18 (2015) 148–158.

[18] B.J. Geiss, A.A. Thompson, A.J. Andrews, R.L. Sons, H.H. Gari, S.M. Keenan, O.B. Peersen, Analysis of flavivirus NS5 methyltransferase cap binding, *J. Mol. Biol.* 385 (2009) 1643–1654.

[19] H. Dong, B. Zhang, P.Y. Shi, Flavivirus methyltransferase: a novel antiviral target, *Antiviral Res.* 80 (2008) 1–10.

[20] S.P. Falk, B. Weisblum, Aptamer displacement screen for flaviviral RNA methyltransferase inhibitors, *J. Biomol. Screen.* 19 (2014) 1147–1153.

[21] B.J. Geiss, H. Stahla, A.M. Hannah, H.H. Gari, S.M. Keenan, Focus on flaviviruses: current and future drug targets, *Future Med. Chem.* 1 (2009) 327–344.

[22] S.P. Lim, C.G. Noble, P.Y. Shi, The dengue virus NS5 protein as a target for drug discovery, *Antiviral Res.* 119 (2015) 57–67.

[23] H. Tian, X. Ji, X. Yang, W. Xie, K. Yang, C. Chen, C. Wu, H. Chi, Z. Mu, Z. Wang, H. Yang, The crystal structure of Zika virus helicase: basis for antiviral drug design, *Protein Cell* 7 (2016) 450–454.

[24] P.R. Gerber, K. Müller, MAB: a generally applicable molecular force field for structure modelling in medicinal chemistry, *J. Comput. Aided Mol. Des.* 9 (1995) 251–268.

[25] J. Wang, P. Cieplak, P.A. Kollman, How well does a restrained electrostatic potential (RESP) model perform in calculating conformational energies of organic and biological molecules? *J. Comp. Chem.* 21 (2000) 1049–1074.

[26] D.A. Case, T.A. Darden, T.E. Cheatham, C.L. Simmerling, J. Wang, R.E. Duke, R. Luo, M. Crowley, R.C. Walker, W. Zhang, K.M. Merz, Amber 10, University of California, San Francisco, 2008.

[27] D. Luo, N. Wei, D.N. Doan, P.N. Paradkar, Y. Chong, A.D. Davidson, M. Kotaka, J. Lescar, S.G. Vasudevan, Flexibility between the protease and helicase domains of the dengue virus NS3 protein conferred by the linker region and its functional implications, *J. Biol. Chem.* 285 (2010) 18817–18827.

[28] R. Assenberg, E. Mastrangelo, T.S. Walter, A. Verma, M. Milani, R.J. Owens, D.I. Stuart, J.M. Grimes, E.J. Mancini, Crystal structure of a novel conformational state of the flavivirus ns3 protein: implications for polyprotein processing and viral replication, *J. Virol.* 83 (2009) 12895–12906.

[29] P. Erbel, N. Schiering, A. D’Arcy, M. Renatus, M. Kroemer, S.P. Lim, Z. Yin, T.H. Keller, S.G. Vasudevan, U. Hommel, Structural basis for the activation of Flaviviral NS3 proteases from dengue and West Nile virus, *Nat. Struct. Mol. Biol.* 13 (2006) 372–373.

[30] G. Lu, P. Gong, Crystal Structure of the full-length Japanese encephalitis virus NS5 conserved methyltransferase-polymerase interface, *PLoS Pathog.* 9 (2013) e1003549.

[31] H. Malet, M.P. Egloff, B. Selisko, R.E. Butcher, P.J. Wright, M. Roberts, A. Gruez, G. Sulzenbacher, C. Vonnheim, G. Bricogne, J.M. Mackenzie, A.A. Khromykh, A.D. Davidson, B. Canard, Crystal structure of the RNA polymerase domain of the West Nile virus non-structural protein 5, *J. Biol. Chem.* 282 (2007) 10678–10689.

[32] R. Thomsen, M.H. Christensen, MolDock: a new technique for high-accuracy molecular docking, *J. Med. Chem.* 49 (2006) 3315–3321.

[33] T.A. Halgren, Merck molecular force field. I. Basis form, scope, parameterization, and performance of MMFF 94, *J. Comput. Chem.* 17 (1996) 490–519.

[34] Y. Pan, H. Huang, S. Cho, A.D. MacKerell, Consideration of molecular weight during compound selection in virtual target-based database screening, *J. Chem. Inf. Comput. Sci.* 43 (2003) 267–272.

[35] Y.M. Yang, T.W. Shen, A pharmacophore-based evolutionary approach for screening selective estrogen receptor modulators, *Proteins Struct. Funct. Bioinf.* 59 (2005) 205–220.

[36] C. Abad-Zapatero, J.T. Metz, Ligand efficiency indices as guideposts for drug discovery, *Drug Disc. Today* 10 (2005) 464–469.

[37] G. Carta, A.J.S. Knox, D.G. Lloyd, Unbiasing scoring functions: a new normalization and rescoring strategy, *J. Chem. Inf. Model.* 47 (2007) 1564–1571.

[38] C.A. Lipinski, F. Lombardo, B.W. Dominy, P.J. Feeney, Experimental and computational approaches to estimate solubility and permeability in drug discovery and development settings, *Adv. Drug Deliv. Rev.* 64 (2012) 4–17.

[39] F. Tillequin, M. Koch, M. Bert, T. Sevenet, Plantes de Nouvelle Calédonie. LV. Isoborrévérine et borrévérine: alcaloïdes bis-indoliques de *Flindersia fourieri*, *J. Nat. Prod.* 42 (1979) 92–95.

[40] L.S. Fernandez, M.S. Buchanan, A.R. Carroll, Y.J. Quinn, R.J. Quinn, V.M. Avery, Flinderolones A–C: antimalarial bis-indole alkaloids from *Flindersia* species, *Org. Lett.* 11 (2009) 329–332.

[41] S. Oshimi, J. Deguchi, Y. Hirasawa, W. Ekasari, A. Widyawaruyanti, T.S. Wahyuni, N.C. Zaini, O. Shirota, H. Morita, Cassiarins C–E, antiplasmodial alkaloids from the flowers of *Cassia siamea*, *J. Nat. Prod.* 72 (2009) 1899–1901.

[42] T.S. Wu, L.S. Shi, S.C. Kuo, Alkaloids and other constituents from *Tribulus terrestris*, *Phytochemistry* 50 (1999) 1411–1415.

[43] T. Fukai, B. Cai, K. Maruno, Y. Miyakawa, M. Konishi, T. Nomura, An isoprenylated flavanone from *Glycyrrhiza glabra* and rec-assay of licorice phenols, *Phytochemistry* 49 (1998) 2005–2013.

[44] Y. Asada, W. Li, T. Yoshikawa, The first prenylated biaurone: licoagron from hairy root cultures of *Glycyrrhiza glabra*, *Phytochemistry* 50 (1999) 1015–1019.

[45] V. Kuete, I.K. Simo, B. Ngameni, J.D. Bigoga, J. Watchueng, R.N. Kapgwap, F.X. Etoa, B.N. Tchaleu, V.P. Beng, Antimicrobial activity of the methanolic extract: fractions and four flavonoids from the twigs of *Dorstenia angusticornis* Engl. (Moraceae), *J. Ethnopharmacol.* 112 (2007) 271–277.

[46] S. Lavoie, J. Legault, F. Simard, E. Chiasson, A. Pichette, New antibacterial dihydrochalcone derivatives from buds of *Populus balsamifera*, *Tetrahedron Lett.* 54 (2013) 1631–1633.

[47] S. Yin, C.Q. Fan, Y. Wang, L. Dong, J.M. Yue, Antibacterial prenylflavone derivatives from *Psoralea corylifolia*: and their structure-activity relationship study, *Bioorg. Med. Chem.* 12 (2004) 4387–4392.

[48] T. Narendar, K. Shweta, M.S. Tanvir, K. Srivastava Rao, S.K. Puri, Prenylated chalcones isolated from *Crotalaria* genus inhibits in vitro growth of the human malaria parasite *Plasmodium falciparum*, *Bioorg. Med. Chem. Lett.* 15 (2005) 2453–2455.

[49] K. Hatayama, M. Komatsu, Studies on the constituents of *Sophora* species. V. Constituents of the root of *Sophora angustifolia* Sieb. et Zucc. (2), *Chem. Pharm. Bull.* 19 (1971) 2126–2131.

[50] B.C.L. Chan, H. Yu, C.W. Wong, S.L. Lui, C. Jolival, C. Ganem-Elbaz, J.M. Paris, B. Morleo, M. Litaudon, C.B.S. Lau, M. Ip, K.P. Fung, P.C. Leung, Q.B. Han, Quick identification of kuraridin: a noncytotoxic anti-MRSA (methicillin-resistant *Staphylococcus aureus*) agent from *Sophora flavescens* using high speed counter-current chromatography, *J. Chromatogr. B* 880 (2012) 157–162.

[51] S.J. Kim, K.H. Son, H.W. Chang, S.S. Kang, H.P. Kim, Tyrosinase inhibitory prenylated flavonoids from *Sophora flavescens*, *Biol. Pharm. Bull.* 26 (2003) 1348–1350.

[52] H.J. Kwon, J.H. Jeong, S.W. Lee, Y.B. Ryu, H.J. Jeong, K. Jung, J.S. Lim, K.O. Cho, W.S. Lee, M.C. Rho, S.J. Park, *In vitro* anti-reovirus activity of kuraridin isolated from *Sophora flavescens* against viral replication and hemagglutination, *J. Pharm. Sci.* 128 (2015) 159–169.

- [53] W. Li, Y. Asada, T. Yoshikawa, Flavonoid constituents from *Glycyrrhiza glabra* hairy root cultures, *Phytochemistry* 55 (2000) 447–456.
- [54] Dictionary of Natural Products on DVD version 24:2, CRC Press Boca Raton, Florida, USA, 2015.
- [55] J. Kouam, L.B.K. Mabeku, J.R. Kuate, A.T. Tiabou, Z.T. Fomum, Antimicrobial glycosides and derivatives from roots of *Picralima nitida*, *Int. J. Chem.* 3 (2011) 23–31.
- [56] K. Šmejkal, S. Chudík, P. Klouček, R. Marek, J. Cvačka, M. Urbanová, J. Ondřej, L. Kokoška, T. Šlapetová, P. Holubová, A. Zima, M. Dvorská, Antibacterial C-geranylflavonoids from *Paulownia tomentosa* fruits, *J. Nat. Prod.* 71 (2008) 706–709.
- [57] M.M. Salem, J. Capers, S. Rito, K.A. Werbovetz, Antiparasitic activity of C-geranyl flavonoids from *Mimulus bigelovii*, *Phytother. Res.* 25 (2011) 1246–1249.
- [58] K. Šmejkal, J. Svačinová, T. Šlapetov, K. Schneiderová, S. Dall'Acqua, G. Innocenti, V. Závalová, P. Kollár, S. Chudík, R. Marek, O. Julínek, M. Urbanová, M. Kartal, M. Csöllei, K. Doležal, Cytotoxic activities of several geranyl-substituted flavanones, *J. Nat. Prod.* 73 (2010) 568–572.
- [59] M.L. Barrett, A.M. Scutt, F.J. Evans, Cannflavin A and B: prenylated flavones from *Cannabis sativa* L., *Experientia* 42 (1986) 452–453.
- [60] R. Chanphen, Y. Thebtaranonth, S. Wanauppathamkul, Y. Yuthavong, Antimalarial principles from *Artemisia indica*, *J. Nat. Prod.* 61 (1998) 1146–1147.
- [61] L.A. Mitscher, S.R. Gollapudi, I.K. Khanna, S.D. Drake, T. Hanumaiah, T. Ramaswam, K.V.J. Rao, Antimicrobial agents from higher plants: activity and structural revision of flemiflavanone-D from *Flemingia stricta*, *Phytochemistry* 24 (1985) 2885–2887.
- [62] M. Kuroyanagi, T. Arakawa, Y. Hirayama, T. Hayashi, Antibacterial and antiandrogen flavonoids from *Sophora flavescens*, *J. Nat. Prod.* 62 (1999) 1595–1599.
- [63] M. Fiebig, H. Wagner, Neue antihepatotoxisch wirksame Flavonolignane aus einer weißblühenden Silybum-Varietät, *Planta Med.* 50 (1984) 310–313.
- [64] S. Inui, T. Hosoya, Y. Shimamura, S. Masuda, T. Ogawa, H. Kobayashi, K. Shirafuji, R.T. Moli, I. Kozono, K. Shin-ya, S. Kumazawa, Solophenols B-D and solomonin. New prenylated polyphenols isolated from propolis collected from the Solomon Islands and their antibacterial activity, *J. Agric. Food Chem.* 60 (2012) 11765–11770.
- [65] J.D. Cha, S.E. Moon, J.Y. Kim, E.K. Jung, Y.S. Lee, Antibacterial activity of sophoraflavanone G isolated from the roots of *Sophora flavescens* against methicillin-resistant *Staphylococcus aureus*, *Phytother. Res.* 23 (2009) 1326–1331.
- [66] V.F. de Andrade-Neto, T. da Silva, L.M.X. Lopes, V.E. do Rosário, F.P. Varotti, A.U. Krettli, Antiplasmodial activity of aryltetralone lignans from *Holostylis reniformis*, *Antimicrob. Agents Chemother.* 51 (2007) 2346–2350.
- [67] L.G. Felipe, D.C. Baldoqui, M.J. Kato, V.S. Bolzani, E.F. Guimarães, R.M.B. Cicarelli, M. Furlan, Trypanocidal tetrahydrofuran lignans from *Peperomia blanda*, *Phytochemistry* 69 (2008) 445–450.
- [68] T.J. Schmidt, S. Rzeppa, M. Kaiser, R. Brun, *Larrea tridentata* – absolute configuration of its epoxy lignans and investigations on its antiprotozoal activity, *Phytochem. Lett.* 5 (2012) 632–638.
- [69] C.C. Wen, Y.H. Kuo, J.T. Jan, P.H. Liang, S.Y. Wang, H.G. Liu, C.K. Lee, S.T. Chang, C.J. Kuo, S.S. Lee, C.C. Hou, P.W. Hsiao, S.C. Chien, L.F. Shyur, N.S. Yang, Specific plant terpenoids and lignoids possess potent antiviral activities against severe acute respiratory syndrome coronavirus, *J. Med. Chem.* 50 (2007) 4087–4095.
- [70] G.A. Mikaya, D.G. Turabelidze, E.P. Kemertelidze, N.S. Wulfson, Kaerophyllin: a new lignan from *Chaerophyllum maculatum*, *Planta Med.* 43 (1981) 378–380.
- [71] B.B. Zhang, Y. Dai, Z.X. Liao, L.S. Ding, Three new antibacterial active diaryheptanoids from *Alpinia officinarum*, *Fitoterapia* 81 (2010) 948–952.
- [72] S.O. Kruse, A. Löhning, G.F. Pauli, H. Winterhoff, A. Nahrstedt, Fukiic and piscidic acid esters from the rhizome of *Cimicifuga racemosa* and the *in vitro* estrogenic activity of fukinolic acid, *Planta Med.* 65 (1999) 763–764.
- [73] S. Stromeier, F. Peteret, A. Nahrstedt, Phenolic esters from the rhizomes of *Cimicifuga racemosa* do not cause proliferation effects in MCF-7 cells, *Planta Med.* 71 (2005) 495–500.
- [74] M.J. Parnham, K. Kesselring, Rosmarinic acid, *Drugs Future* 10 (1985) 756–757.
- [75] F. Yoshizaki, S. Hisamichi, Y. Kondo, Y. Sato, S. Nozoe, Studies on shikon. III. new furylhydroquinone derivatives, shikonofurans A, B, C, D and E, from *Lithospermum erythrorhizon* Sieb. et Zucc., *Chem. Pharm. Bull.* 30 (1982) 4407–4411.
- [76] A. Sendl, J.L. Chen, S.D. Jolad, C. Stoddart, E. Rozhon, M. Kernan, W. Nanakorn, M. Balick, Two new naphthoquinones with antiviral activity from *Rhinacanthus nasutus*, *J. Nat. Prod.* 59 (1996) 808–811.
- [77] R.F.X. Bauer, I.A. Khan, H. Lotter, H. Wagner, V. Wray, Structure and stereochemistry of new sesquiterpene esters from *Echinacea purpurea* (L.) Moench, *Helv. Chim. Acta* 68 (1985) 2355–2358.
- [78] T.A. Bischoff, C.J. Kelley, Y. Karchesy, M. Laurantos, P. Nguyen-Dinh, A.G. Arefi, Antimalarial activity of lactucin and lactucopicrin: sesquiterpene lactones isolated from *Cichorium intybus* L., *J. Ethnopharmacol.* 95 (2004) 455–457.
- [79] K. Hu, X.S. Dong, H. Kobayashi, S. Iwasaki, A furostanol glycoside from rhizomes of *Dioscorea colletii* var *hypoglauca*, *Phytochemistry* 44 (1997) 1339–1342.
- [80] T. Xu, Y. Xu, Y. Liu, S. Xie, Y. Si, D. Xu, Two new furostanol saponins from *Tribulus terrestris* L., *Fitoterapia* 80 (2009) 354–357.
- [81] M. Yoshikawa, T. Murakami, H. Komatsu, N. Murakami, J. Yamahara, H. Matsuda, Medicinal foodstuffs. IV. fenugreek seed (1): structures of trigoneosides Ia, Ib, IIa, IIb, IIIa, IIIb, new furostanol saponins from the seeds of Indian *Trigonella foenum-graecum* L., *Chem. Pharm. Bull.* 45 (1997) 81–87.
- [82] W. Li, Y. Asada, T. Yoshikawa, Antimicrobial flavonoids from *Glycyrrhiza glabra* hairy root cultures, *Planta Med.* (1998) 746–747.
- [83] D. Luo, T. Xu, R.P. Watson, D. Scherer-Becker, A. Sampath, W. Jahnke, S.S. Yeong, C.H. Wang, S.P. Lim, A. Strongin, S.G. Vasudevan, J. Lescar, Insights into RNA unwinding and ATP hydrolysis by the flavivirus NS3 protein, *EMBO J.* 27 (2008) 3209–3219.
- [84] C.A. Belon, D.N. Frick, Helicase inhibitors as specifically targeted antiviral therapy for hepatitis C, *Future Virol.* 4 (2009) 277–293.
- [85] C.G. Noble, P.Y. Shi, Structural biology of dengue virus enzymes: towards rational design of therapeutics, *Antiviral Res.* 96 (2012) 115–126.