**Dengue protease inhibition activity of selected Malaysian medicinal herbs**

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**Abstract.** Dengue fever is one of major health problem around the world including Malaysia. It is caused by the arthropode-borne flavivirus and transmitted by the bite of the *Aedes aegypti* or *Aedes albopictus* mosquito infected with one of the four dengue virus serotypes (DENV-1, DENV-2, DENV-3, or DENV-4). In this study, a screening exercise of various Malaysian medicinal plants showed that the extracts of *Lawsonia inermis*, *Dryobalanops aromatica*, *Punica granatum*, *Zizyphus jujuba* L. and *Zingiber zerumbet* exhibited potent inhibitory activity against NS2B-NS3 serine protease. The methanol extracts of *Dryobalanops aromatica* showed inhibition of 99.70 % at concentration of 200 µg/mL with IC₅₀ value of 0.30 ± 0.16 µg/mL.

**INTRODUCTION**

Dengue is a mosquito-borne viral disease and classified as one of the rapidly spread diseases in the world with about 4.54 million dengue infections worldwide in 2016 (World Health Organization, 2007a). Dengue can be transmitted through the saliva of the female mosquito of *Aedes aegypti* and *Aedes albopictus*. According to the World Health Organization (WHO), it was estimated that 2.5 billion people, two-fifths of the world’s population, are susceptible to dengue infection risk. The organization reported increase of 55% of cases from the year of 2010 (2.2 million) to 2015 (3.2 million) (World Health Organization, 2007a). In Malaysia, even though dengue infection cases decreased by 25.3% from January until May 2017 as compared to the same period in 2016, it is still considered to be one of the fatal health threats to Malaysia (World Health Organization, 2007b). Therefore, there is an urgent need to look for effective modes to prevent and cure the disease.

Recently, Sanofi had successfully developed the first WHO-approved vaccine against dengue known as Dengvaxia. It was claimed to be effective against all of the four different dengue serotypes. Nonetheless, the effort of finding potential therapeutics against dengue infection continues as the efficiency of Dengvaxia varies for the different serotypes, with the lowest efficiency was shown against Denv-2 (35%) (Capeding et al., 2014). Developing and producing a vaccine...
against dengue virus is the most challenging part because the vaccine have to be effective towards all the dengue serotypes (Normile, 2013). Additionally, there is a lot of aspects should be accounted for when developing and producing a vaccine such as the unstable current economic situation. Therefore, it is recommended to investigate the local plants that might lead to the development of the vaccine.

Dengue virus contains a single-stranded RNA of positive polarity, encoding for a single polyprotein precursor of 3,391 amino acids comprising three structural and seven non-structural (NS) proteins (Irie et al., 1989). Dengue virus replication is dependent on the correct cleavage of this polyprotein and requires both host cell proteases and the virus-encoded, two-component protease, NS2B-NS3 (Falgout et al., 1991; Yusof et al., 2000). Inhibiting the NS2B-NS3 protease should prevent virus infection, thus, it serves as a target in the development of anti-viral drugs. Several studies have reported the use of natural products in ascertaining their inhibitory potentials against dengue NS2B-NS3 protease. Compounds isolated from the plant Boesenbergia rotunda (L.) Mansf. Kulturpfl. (Zingiberaceae family) had shown inhibition against this protease (Kiat et al., 2006; Othman et al., 2008). Two cyclohexenyl chalcone derivatives namely, 4-hydroxypanduratin A and panduratin A, exhibited competitive inhibitions against DENV-2 NS2B-NS3 protease with reported inhibition constant, $K_i$ values of 25 and 21 µM, respectively, while pinostrobin (flavonoid) exhibited non-competitive inhibition with IC$_{50}$ value of 90 mg/mL. From computational investigations, it was suggested that pinostrobin bind into an allosteric binding site via interaction with Lys74, adjacent to one of the catalytic triad residues, Asp75, of the NS3 protein. (de Sousa et al., 2015) had performed screening assays of flavonoids against NS2B-NS3 protease of DENV-2 and DENV-3. The results obtained showed that the flavonoids exhibited inhibitory activities with IC$_{50}$ values ranging from 15 to 44 µM. Agathisflavone and myricetin turned out to be non-competitive inhibitors with $K_i$ values of 11 and 4.7 µM, respectively.

Malaysia is rich with many medicinal plants inspired from religious, traditional beliefs and practices. Amongst these plants, five plants were chosen to be the subject of this study, which are Lawsonia inermis (Inai), Punica granatum (Delima), Dryobalanops aromatica (Kapur barus), Zizyphus jujuba Lam. (Bidara) and Zingiber zerumbet (Halia Bara). All these plants are mentioned in Al-Quran and AHadith, and they are used and consumed in Malaysia as food and for various traditional medicinal purposes.

Lawsonia inermis, recognized as henna or inai was used as cosmetic purposes and highly demanded for the production of halal hair and nail dye products (Alia et al., 1995; Badoni Semwal et al., 2014; Dasgupta et al., 2003). Besides that, it is traditionally used for treatment of inflammatory conditions, ringworm, dandruff, jaundice, calculus affliction, rheumatism, stomach disorder, skin diseases and sore throat (Alia et al., 1995; Badoni Semwal et al., 2014; Sultana et al., 2009). Lawsonia inermis has been shown to possess numerous biological activities such as analgesic, hypoglycemic, hepatoprotective, immunostimulant, anti-inflammatory, anti-bacterial, anti-microbial, anti-fungal, anti-viral, anti-parasitic, anti-trypansomal, anti-dermatophytic, anti-oxidant, anti-fertility, tuberculostatic and anti-cancer (Agarwal et al., 2014; Muhammad, 2005). This plant was reported to contain lawsone, gallic acid, tannic acid, $\alpha$-D-glucose, esculetin, fraxetin, isoplumbagin, scopoletin, betulin, betulinic acid, hennadiol, lupeol, lacoumarin, laxanthone (Nik et al., 2012; Sultana et al., 2009). It also contains carbohydrates, proteins, flavonoids, tannins and phenolic compounds, alkaloids, terpenoids, quinones, coumarins, xanthones, flavone glycosides, pentacyclic triterpenes and fatty acids (Agarwal et al., 2014; Badoni Semwal et al., 2014).

Pomegranate or scientifically known as Punica granatum is distributed throughout Iran, Mediterranean countries, tropical
Africa, India, Malaysia and to some extent, in the United States (Viuda-Martos et al., 2010; Zhang et al., 2010). This plant was traditionally held sacred by several of the world’s major religions such as Islam, Christian and Buddhist (Langley, 2000). In Malaysia, the young leaves of *Punica granatum* was burned and the ash is mixed with water and drank to treat stomach ache (Ong & Norzalina, 1999). A mixture of juice, pounded fruit peel, sugar and warm water is drank to treat vaginal white discharges. Ong et al. (2011) reported that the juice can also be used to treat leucorrhea, expel intestinal worms and to counter obesity for slimming purpose, the juice from the fruits is mixed with a little salt and drank regularly (Ong & Nordiana, 1999; Ong et al., 2011). The extracts from different parts of this plant had been studied for numerous pharmacological activities, such as anti-oxidant, anti-tumor, anti-bacterial, astringent, anti-diarrheal and anti-obesity activities (Zhang et al., 2010). Many researchers revealed that *Punica granatum* is a rich source of anthocyanins, polyphenols, tannins, phenolics, flavonoids, ellagitannins, proanthocyanidin and many minerals, including potassium, nitrogen, calcium, magnesium, phosphorus, and sodium (Heftmann et al., 1966; Moneam et al., 1988; Poyrazoglu et al., 2002; Viuda-Martos et al., 2010).

*Dryobalanops aromatica* have been reported to yield terpenoid compounds like d-borneol, terpenin-4-ol, α-terpineol, α-pinene and caryophyllene, which are known to exhibit various activities such as anti-microbial, antiviral, anti-inflammatory and cytotoxic effects (Le et al., 2016). Examples of compounds that have been isolated from this plant are malaysianol A, flexuasol A, vaticanol B, vaticanol C, laevifonol, amelpopsin E, α-viniferin, ε-viniferin, diptoidonesin A, bergenin, oleanolic acid acetate, hedragonic acid, dryobalanonoloc acid, methyl 11-oxoasiatate and dryobalanolide (Cheung & Wong, 1972; Wibowo et al., 2011).

*Zizyphus jujuba*, a species from the Rhamnaceae family, is an indigenous plant possessing an array of medicinal properties, attributed by a diverse group of secondary metabolites such as alkaloids, flavonoids, terpenoids and glucosides (Cheng et al., 2000). *Zizyphus jujuba* has been used traditionally to treat insomnia, anxiety, sleep-related problems, hemorrhage and diarrhoea (Bencao, 1999; Chopra et al., 1956; Lee, 1986; Sun et al., 2011). Many researchers have reported multiple medicinal values of this plant such as anti-allergic, anti-ulcer, anti-fertility, wound healing and anti-oxidant properties (Ansari et al., 2006; Chopda, 2009; Chopda & Mahajan, 2009; Ganachari & Kumar, 2004; Kim & Han, 1996; Su et al., 2002). It is also found to be a potential natural pesticide in controlling *Aedes aegypti* mosquitoes (Devī & Bora, 2015).

*Zingiber zerumbet* is a species belonging to the Zingiberaceae family that has long been used in Asian popular medicine with many reported biological activities such as anti-cancer, anti-inflammation, anti-Alzheimer's disease, anti-viral, anti-tumor, anti-oxidant, anti-allergic, and anti-microbial activities (Abdul et al., 2008; Abdul Hamid et al., 2012; Matthes et al., 1980; Singh et al., 2012). The dichloromethane (DCM) and methanol (MeOH) extracts of the rhizome of *Zingiber zerumbet* demonstrated potent larvicidal activity against *Aedes nuneztovari* larvae and *Aedes aegypti* larvae, suggesting that *Zingiber zerumbet* could be an alternative to prevent the spread of mosquito vectors diseases (Bucker et al., 2013). Examples of some compounds isolated from the essential oil of *Zingiber zerumbet* are zerumbone, zerumbone epoxide, humulene, zederone, camprene, diferuloylmethane, feruloyl-p-coumaroylmethane and di-p-coumaroylmethane.

In pursuing our interest in the studies of Malaysian medicinal plants (Abdul et al., 2008; Ablat et al., 2014; Awang et al., 2010; Hamdi et al., 2015; Othman et al., 2006), this manuscript reports the inhibition of NS2B-NS3 serine protease (IC₅₀) by five traditional medicinal plants, namely *Lawsonia inermis* (Inai), *Punica granatum* (Delima), *Dryobalanops aromatica* (Kapur barus), *Zizyphus jujuba* Lam. (Bidara) and *Zingiber zerumbet* (Halita Bara).
MATERIALS AND METHODS

General procedures and plant materials
All plant materials were purchased from the local people and local traditional herbs suppliers. The voucher specimens were deposited at the Herbarium of the Department of Chemistry, Faculty of Science, University of Malaya. The industrial grade solvents were distilled prior to use for extraction.

Extraction
All plant materials were thoroughly rinsed with water. Each part of the plants (bark, leaves and rhizomes) were sliced and air-dried until constant mass. Dried plant materials were ground and the extraction process was carried out using maceration method, of which dried plant materials were first defatted with hexane and then soaked in ethyl acetate (EA) with periodical stirring for three days. Upon day 3, the EA extracts were separated from the plant materials through filtration. The filtrates were then dried using rotary evaporator to give dried extracts. To maximize extraction yield, the plant materials were subjected to the same extraction process for two more times. The dried extracts of each extraction were then combined to give the total EA extraction yield. After that, the same extraction process as above was carried out using methanol, MeOH (ethanol, EtOH for Zingiber zerumbet) and then water (except for Zingiber zerumbet) to give methanol, ethanol and water extracts, respectively.

Denv-2 protease inhibition assay
Protease inhibition study was performed using purified Denv-2 NS2B-NS3 protease (Abdul et al., 2016; Nawi, 2015), and Boc-Gly-Arg-Arg-MCA as the substrate. The bioassay protocol was employed as described by Nawi (2015) and Abdul H.W. et al. (2016) with minor modification (Abdul et al., 2016; Nawi, 2015). The enzyme concentration used in the assay was 0.5 µM and substrate concentration was 10 mM in 200 mM Tris-HCl (buffer) with pH 8.5. The plant extract concentration used in this assay was 200 µg/mL. All tests were performed in quadruplicate. Firstly, a Tris buffer (pH 8) was pipetted to the wells, followed by 1 µL of plant extract and 3.1 µL of enzyme. Before adding the substrate, the enzyme and inhibitor were incubated at 37°C for 10 minutes. After adding the substrate, the reaction mixtures were incubated at 37°C for 60 minutes. All reactions were performed in 96-well plates with final volume of 100 µL per well and fluorescence was detected using the Promega Glomax Multi Detection System microplate reader with excitation and emission wavelengths at 365 and 410-460 nm, respectively. For IC50 determination, the same protocol was used as described before with serial dilutions of plant extract with concentration in the range of 0.78 to 200 µg/mL.

RESULTS AND DISCUSSION
Twenty one plant extracts were screened for their inhibitory activity against Denv-2 NS2B-NS3 protease and the results are shown in Table 1. Thirteen extracts were found to be active at the concentration of 200 µg/mL with percentage of inhibition more than 80%. The IC50 values of these active extracts were determined via serial dilutions which gave IC50 values ranging from 0.30 to 17.55 µg/mL. The percentage inhibition of plant extracts against protease activity at different concentration (0.78 to 200.00 µg/mL) was shown in Figures 1 and 2.

Two extracts were found to have 70% to 75% inhibition at 200 µg/mL; while six extracts exhibited moderate to less inhibition (less than 65% inhibition). Extracts with more than 80% inhibition were subjected to IC50 studies (except for MeOH extract of the barks of Zizyphus jujuba). The MeOH extract from the leaves of Dryobalanops aromatica showed the most potent activity (99.70 %). Moreover, the MeOH extracts of the barks of Punica granatum and Lawsonia inermis exhibited very strong inhibition as compared to their EA extracts. Interestingly, the barks of Lawsonia inermis, Punica granatum and Zizyphus jujuba as well as the leaves of Dryobalanops aromatica indicated an increase in inhibition against Denv-2 NS2B-NS3 protease with increasing polarity.
Table 1. The percentage of inhibition of the crude extracts of *Lawsonia inermis*, *Punica granatum*, *Dryobalanops aromatica*, *Zizyphus jujuba* Lam. and *Zingiber zerumbet* against DENV2 NS2B/NS3 protease

<table>
<thead>
<tr>
<th>Plants</th>
<th>Parts</th>
<th>Extracts</th>
<th>Mean percentage inhibition at 200 (µg/mL)</th>
<th>IC₅₀ ± Standard deviation (µg/mL)</th>
</tr>
</thead>
<tbody>
<tr>
<td><em>Lawsonia inermis</em> (KL 5824)</td>
<td>Leaves (L)</td>
<td>Ethyl acetate</td>
<td>17.62</td>
<td>NT</td>
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<td></td>
<td></td>
<td>Methanol</td>
<td>56.52</td>
<td>NT</td>
</tr>
<tr>
<td></td>
<td>Barks (B)</td>
<td>Ethyl acetate</td>
<td>86.43</td>
<td>0.77 ± 0.37</td>
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<tr>
<td></td>
<td></td>
<td>Methanol</td>
<td>87.99</td>
<td>0.33 ± 0.14</td>
</tr>
<tr>
<td><em>Punica granatum</em> (KL 5826)</td>
<td>Leaves (L)</td>
<td>Ethyl acetate</td>
<td>92.23</td>
<td>9.03 ± 3.21</td>
</tr>
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<td></td>
<td></td>
<td>Methanol</td>
<td>91.02</td>
<td>3.10 ± 0.14</td>
</tr>
<tr>
<td></td>
<td>Barks (B)</td>
<td>Ethyl acetate</td>
<td>85.36</td>
<td>17.55 ± 0.54</td>
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<td></td>
<td></td>
<td>Methanol</td>
<td>91.84</td>
<td>1.72 ± 0.29</td>
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<td><em>Dryobalanops aromatica</em> (KL 5829)</td>
<td>Leaves (L)</td>
<td>Ethyl acetate</td>
<td>95.10</td>
<td>3.49 ± 0.65</td>
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<td></td>
<td></td>
<td>Methanol</td>
<td>99.70</td>
<td>0.30 ± 0.16</td>
</tr>
<tr>
<td></td>
<td>Barks (B)</td>
<td>Ethyl acetate</td>
<td>72.08</td>
<td>NT</td>
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<tr>
<td></td>
<td></td>
<td>Methanol</td>
<td>96.52</td>
<td>0.51 ± 0.42</td>
</tr>
<tr>
<td></td>
<td>Fruits (F)</td>
<td>Ethyl acetate</td>
<td>50.44</td>
<td>NT</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Methanol</td>
<td>75.00</td>
<td>NT</td>
</tr>
<tr>
<td><em>Zizyphus jujuba</em> Lam. (KL 5828)</td>
<td>Leaves (L)</td>
<td>Ethyl acetate</td>
<td>94.44</td>
<td>7.83 ± 0.99</td>
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<td></td>
<td></td>
<td>Methanol</td>
<td>57.67</td>
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</tr>
<tr>
<td></td>
<td>Barks (B)</td>
<td>Ethyl acetate</td>
<td>93.10</td>
<td>2.47 ± 0.72</td>
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<td></td>
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<td>Methanol</td>
<td>94.58</td>
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<tr>
<td><em>Zingiber zerumbet</em> (KL 5835)</td>
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<td>Hexane</td>
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<td>Ethyl acetate</td>
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<td>Ethanol</td>
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<tr>
<td><em>Quercetin</em></td>
<td>Standard</td>
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<td>100.00</td>
<td>3.17 ± 0.65</td>
</tr>
</tbody>
</table>

*NT: not tested

Figure 1. Percentage of inhibition of the ethyl acetate extracts of *Lawsonia inermis*, *Punica granatum*, *Dryobalanops aromatica*, *Zizyphus jujuba* Lam. and *Zingiber zerumbet* against DENV2 NS2B/NS3 protease at concentration ranging from 0.78 to 200.00 µg/mL.
Figure 2. Percentage of inhibition of the methanol extracts of 
*Lawsonia inermis*, *Punica granatum*, 
*Dryobalanops aromatica*, *Zizyphus jujuba* Lam. and *Zingiber zerumbet* against DENV2 NS2B/NS3 protease protease at concentration ranging from 0.78 to 200.00 µg/mL.

of the solvent extracts (from EA to MeOH). This observation suggests that polar functionalities such as hydroxyl and carbonyl groups play an important role in facilitating interactions within the protease enzyme.

Among all the extracts, the MeOH extract of *Dryobalanops aromatica* showed the most potent inhibition against DENV-2 NS2B-NS3 protease with an IC\(_{50}\) value of 0.30 ± 0.16 µg/mL. In addition, leaves and bark of *Punica granatum* (MeOH extracts), barks of *Lawsonia inermis* (EA and MeOH extracts), *Dryobalanops aromatica* (MeOH extract), *Zizyphus jujuba* (EA extract) and the rhizomes of *Zingiber zerumbet* (EA extract) showed higher inhibiting potency as compared to quercetin (IC\(_{50}\) value of 3.17 µg/mL) which is used as a standard because it has demonstrated significant anti-dengue inhibitory activity (de Sousa et al., 2015; Zandi et al., 2011a; Zandi et al., 2011b). Furthermore, Tan has reported the MeOH fraction of *Zingiber zerumbet* were potent against DENV-2 NS2B-NS3 protease at 300 ppm with different extraction method (Tan et al., 2006). However, the extracts of EA for the leaves of *Punica granatum*, *Dryobalanops aromatica*, *Zizyphus jujuba* and the bark of *Punica granatum* were found to be less active. Based on the results obtained, it shows that the medicinal plants has potential to be investigated further for the development of anti-dengue agents.

CONCLUSIONS

The results of these preliminary studies showed that, in general, all the plants tested showed potent activity either in their leaves or bark extracts or both extracts. *Dryobalanops aromatica* (kapur barus), *Zizyphus jujuba* (bidara) and *Punica granatum* (delima) revealed potent activities in both leaves and bark extracts (more than 80% inhibition at 200 µg/mL). The MeOH extract of *Dryobalanops aromatica* leaves was found to exhibit very potent activity against DENV-2 NS2B-NS3 protease with an IC\(_{50}\) value of ten times lower than the standard quercetin. These medicinal plants have been reported to contain compounds such as apigenin (*Lawsonia inermis*), quercetin-3-O-rutinoside (*Zizyphus jujuba*), kaempferol-3-O-methylether (*Zingiber zerumbet*), quercetin (*Punica granatum*) and ß-caryophyllene (*Dryobalanops aromatica*) (Chauhan & Chauhan, 2001; Guha...
et al., 2011; Jang et al., 2004; Le et al., 2017; Wang et al., 2014). The anti-protease activity of the extracts may be attributed to the presence of these compounds as several studies have shown that some flavonoids and bicyclic sesquiterpenes were active against dengue protease activity (Abd Kadir et al., 2013; Kiat et al., 2006; María et al., 2018; Tang et al., 2012).

Therefore, further studies such as bioassay-guided isolation of chemical constituents from the potent extracts especially for Dryobalanops aromatica can be carried out in the future. These extracts, along with pure compounds from the active extracts, can be tested for in vitro cell-based assays, such as plaque reduction or virus yield inhibition assays, to ascertain their anti-dengue properties in our efforts towards the development of anti-dengue therapeutics.

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