Phytochemical assay and Antiplatelet Activity of Fractions of Velvet Bean Seeds (Mucuna pruriens L.)

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Platelet aggregation is an important factor contributing to the formation of thrombus due to an uncontrolled blood clotting. An antiplatelet agent is a compound which decreases platelet aggregation and inhibits thrombus formation. The objectives of this study were to determine the class of compound employing phytochemical assay and to determine the in vitro antiplatelet activity of four fraction, namely hexane, ethyl acetate, butanol, and water fractions of velvet bean seeds (Mucuna pruriens L.) using epinephrine (EPN) as agonist of platelet aggregation. The antiplatelet activities were tested in human platelet rich plasma with hyperaggregation. To determine the activities, EPN was arranged at 4 level of concentrations (300, 150, 75, and 30 µM), and antiplatelet agents were at 500 µg/ml. The results indicated that ethyl acetate, butanol and water fraction contained high flavonoids and moderate phenols. The water, butanol and ethyl acetate fractions of velvet bean seeds exhibited potential inhibition of EPN-induced platelet aggregation at all concentrations. The strongest antiplatelet agent was water fraction and had the same antiplatelet activity as aspirin at level 150, 75, and 30 µM of EPN. Butanol fraction had the same antiplatelet activity as aspirin at the lowest EPN (30 µM).

Key words: platelet, Mucuna pruriens L., epinephrine, flavonoid, aspirin

INTRODUCTION
Platelet aggregation is an important factor contributing to the formation of thrombus due to an uncontrolled blood clotting. An antiplatelet agent is a compound which decreases platelet aggregation and inhibits thrombus formation. The objectives of this study were to determine the class of compound employing phytochemical assay and to determine the in vitro antiplatelet activity of four fraction, namely hexane, ethyl acetate, butanol, and water fractions of velvet bean seeds (Mucuna pruriens L.) using epinephrine (EPN) as agonist of platelet aggregation. The antiplatelet activities were tested in human platelet rich plasma with hyperaggregation. To determine the activities, EPN was arranged at 4 level of concentrations (300, 150, 75, and 30 µM), and antiplatelet agents were at 500 µg/ml. The results indicated that ethyl acetate, butanol and water fraction contained high flavonoids and moderate phenols. The water, butanol and ethyl acetate fractions of velvet bean seeds exhibited potential inhibition of EPN-induced platelet aggregation at all concentrations. The strongest antiplatelet agent was water fraction and had the same antiplatelet activity as aspirin at level 150, 75, and 30 µM of EPN. Butanol fraction had the same antiplatelet activity as aspirin at the lowest EPN (30 µM).

It has been found that platelet inhibitory drugs, such as aspirin, physiologically reduce the incidence of myocardial infarction, stroke, and death from cardiovascular diseases in prevention trials. A number of prospective randomized clinical trials have shown that 50-500 mg of aspirin per day reduces the risk of primary or secondary cardiovascular events (Olas & Wachowicz 2005). The usage of aspirin has negative side effects: low dose aspirin increases the risk of major bleeding and intestinal ulceration, and for a prolong therapy, it usually causes many side effect (Schror 1997; Ohmori et al. 2006). The aspirin resistance is a clinical inability of aspirin to protect individuals from arterial thrombotic events or when laboratory methods indicate the failure of aspirin to inhibit platelet activity (Cattaneo 2004; Ohmori et al. 2006). It is estimated that between 8-45% of patients suffering an ischemic stroke or cardiovascular disease are aspirin resistant (McKee et al. 2002; Ohmori et al. 2006). It is an urgent need to search natural compounds having antiaggregation properties with minimal side effects and safer. One of the natural compounds with antiplatelet activity is flavonoid. It can prevent atherosclerosis, endotelial damage, leukocyte activation, adhesive
aggregation, and platelet secretion (Koshy et al. 2001; Bucki et al. 2003). Natural antioxidants having minimal side effects can be obtained from plants containing phenolic and polyphenolic compounds, such as flavonoids (Bucki et al. 2003; Koleckar et al. 2008). Flavonoids bioavailability and its relationship with antioxidant and antiplatelet activity, resulting in cardiovascular protection (Violi 2002).

The natural compounds including flavonoids can be obtained ubiquitously in Indonesia, such as in velvet bean Mucuna pruriens L., and it has not been used optimally. Previous research demonstrated the total polyphenolic content of the methanol extract of M. pruriens was 33.04 mg/g gallic acid, a noticeable amount of total phenols (Rajeshwar et al. 2005). M. pruriens contains isoflavones such as daidzin 0.041; glycitein 0.011; genistein 0.050; aglucones total 0.131 mg/l (Handajani 2001). Methanol extract of M. pruriens (MEMP) has high antioxidant activity (Rajeshwar et al. 2005).

MATERIALS AND METHODS

The four fractions (hexane, ethyl acetate, n-butanol, and water fraction) were obtained from ethanol extract of velvet bean seeds (M. pruriens L.).

The four fractions of velvet bean seeds were tested by phytochemical assay including flavonoid assay, phenolic, saponin, triterpenoid, steroid, terpenoid, tanin, and alkaloid assay. The four fractions and aspirin as positive control were diluted with dimethyl sulfoxide (DMSO 1%) achieving at level 500 ìg/ml. Epinephrine (EPN) was diluted with distilled water at various concentrations of 300, 150, 75, 30 ìM. Platelet rich plasma (PRP) obtained from hyperaggregation individual (Helena Laboratories 2008).

9 ml blood was collected from hyperaggregation individual and added with 1 ml 3.8% sodium citrate as anticoagulant. The blood was centrifuged at 100 x g for 10 minutes. The platelet rich plasma (PRP) was removed from the cells with a plastic pipette and placed in a plastic tube. The PRP was maintained at room temperature for 30 minutes. Platelet poor plasma (PPP) was prepared by recentrifuging the remaining blood samples at 1600 x g for 10 minutes. PPP was then removed, placed in a plastic tube, and the tube was maintained at room temperature (Chun-Han et al. 1993; Helena Laboratories 2008).

Aggregation activity was measured by Platelet Aggregation Chromogenic Kinetic System (PACKS-4). 450 ìl of PPP was pipetted into a cuvette. 450 ìl PRP and 40 ìl the antiplatelet agents (aspirin, hexan fraction, ethyl acetate fraction, butanol fraction, water fraction) were also pipetted into cuvettes with stir bar and incubated at 37 °C for 3-5 minutes. The PPP and PRP cuvettes were inserted into appropriate channels and the instrument was set up. 50 ìl of the aggregating reagent dilutions (Epinephrine) at level 300, 150, 75, 30 ìM was added to the PRP cuvette and the percent aggregation was recorded (Helena Laboratories 2008).

All treatments consisted of 20 levels arranged in a factorial design. The first factor was the inducing concentration (four level concentrations) and the second factor was the antiplatelet agent (five level antiplatelet agents). The treatments were replicated three times. To verify the statistical significance of the parameters, the data was calculated for the values of means and standard deviation (M ± SD) and 95% confidence interval (CI) of means. To compare several treatments, we used analysis of variance (ANOVA) with two-factorial completely randomized design. P-values of less than 0.05 were considered as statistically significant. Furthermore, to know the difference of means among treatments and to know the best treatment, we used Duncan’s Post-Hoc test 95% confidence interval. Statistical analysis used SPSS 16.0 program.

RESULTS

The phytochemical assay showed that hexane fraction of velvet bean seeds contained lowest flavonoids and phenolics compound compared with the other fractions (Table 1). Ethyl acetate, butanol and water fractions contained high flavonoid and moderate phenolic compounds, but they did not contain terpenoid and triterpenoid compounds. Butanol fraction contained the highest tanin compared with the other fractions. Water fraction contained moderate alkaloids and butanol and ethyl acetate fractions contained less alkaloids. Hexane, ethyl acetate and butanol fractions contained less steroids. Butanol and water fractions contained less saponins, whereas hexane and ethyl acetate fractions did not contain saponins (Table 1).

Observing the antiplatelet activity, epinephrine (EPN) agonist induced platelet aggregation at all level concentrations, the lower concentration of EPN as agonist caused the reduction of platelet aggregation (Table 2). Aspirin as positive control decreased platelet aggregation at all concentrations of EPN. After aspirin, the most effective fraction showing the antiplatelet activity was

<table>
<thead>
<tr>
<th>Sample (fractions)</th>
<th>Compound content</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Tanins</td>
</tr>
<tr>
<td>Hexane</td>
<td>-</td>
</tr>
<tr>
<td>Ethylacetate</td>
<td>++</td>
</tr>
<tr>
<td>Butanol</td>
<td>+++</td>
</tr>
<tr>
<td>Water</td>
<td>+++</td>
</tr>
</tbody>
</table>

+++: high content, ++: moderate content, +: less content, -: undetected.
the water fraction followed by butanol fraction at all concentrations of EPN, except at 30 mM of EPN. The antiplatelet aggregation at 30 mM concentration of EPN showed similar activity among aspirin, water fraction and butanol fraction of velvet bean (Table 2).

**DISCUSSION**

Previous research showed that velvet bean seeds are rich in novel alkaloids, saponins, and sterols. The seeds of all *Mucuna* species contain a high concentration of L-dopa; velvet bean seeds contain 7-10% L-dopa (Del Carmen et al. 1999). Crude content of velvet bean showed the presence of alkaloid, flavonoid, tannin, saponin, quinone, and steroid/terprenoid (Yulizia 2008). The velvet bean seeds contain the bioactive alkaloids mucunine, mucucaridine, mucuadinine, pruriendine and nicotine, ß-quinone, and steroid/triterpenoid (Yulizia 2008). The previous research showed that infused epinephrine in doses of 0.1 and 0.2 µg/kg per minute on 40 healthy men aged 20 to 40 years increased in the capacity for TXB₂ production by platelets (Kjeldsen et al. 1995). Infused epinephrine up to 0.04 µg/kg per minute in healthy men, increased platelet count and size (Lande et al. 1988). Epinephrine in the presence of fibrinogen and Ca²⁺ (in vitro) induces both primary and secondary aggregation, potentiates aggregation (Kjeldsen et al. 1995).

Aspirin is the best antiplatelet agent at all level of inducer concentrations and as an anti-thrombotic compounds through the inhibition of platelet cyclooxygenase-1 (COX-1) by irreversible acetylation of a specific serine moiety, thereby blocking the formation of thromboxane A₂ (TXA₂) for the lifetime of the platelets (McKee et al. 2002; Ohmori et al. 2006).

Water fraction of velvet bean seeds is comparable to aspirin because of the moderate content of phenols and the high content of flavonoids (Table 1). Phenols and flavonoids show commonly high antioxidant and antiplatelet activities. The previous research showed that

<table>
<thead>
<tr>
<th>Antiplatelet agents</th>
<th>EPN 300 µM</th>
<th>EPN 150 µM</th>
<th>EPN 75 µM</th>
<th>EPN 30 µM</th>
</tr>
</thead>
<tbody>
<tr>
<td>Control (Hyperaggregation)</td>
<td>82.633 ± 8.228b</td>
<td>74.067 ± 3.761ab</td>
<td>68.267 ± 9.215a</td>
<td>66.067 ± 7.732a</td>
</tr>
<tr>
<td>Aspirin</td>
<td>31.333 ± 9.172b</td>
<td>28.600 ± 4.029a</td>
<td>27.433 ± 1.756a</td>
<td>27.000 ± 1.825a</td>
</tr>
<tr>
<td>Hexane fraction</td>
<td>68.233 ± 9.172b</td>
<td>60.767 ± 3.272ab</td>
<td>55.033 ± 4.168a</td>
<td>53.933 ± 4.509a</td>
</tr>
<tr>
<td>Ethyl acetate fraction</td>
<td>69.733 ± 5.064b</td>
<td>69.000 ± 4.784b</td>
<td>69.800 ± 5.237b</td>
<td>58.867 ± 2.950a</td>
</tr>
<tr>
<td>Butanol fraction</td>
<td>54.433 ± 8.315b</td>
<td>54.367 ± 2.318b</td>
<td>51.033 ± 10.934b</td>
<td>31.133 ± 3.134a</td>
</tr>
<tr>
<td>Water fraction</td>
<td>47.600 ± 8.0206b</td>
<td>34.067 ± 4.350a</td>
<td>31.267 ± 3.027a</td>
<td>27.500 ± 4.939a</td>
</tr>
</tbody>
</table>

The data showed means ± standard deviation. The different small letters at the same row (among EPN concentrations) and capital letters at the same column (among antiplatelet agents) show significant at the 5% (Duncan’s Post Hoc test).

### Table 2. The antiplatelet activity of velvet bean seed fractions

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methanol extract of velvet bean seeds contained a noticeable amount of total phenols (Rajeshwar et al. 2005). Phenolic compounds exhibit a wide range of biological effects, including antiplatelet, anti-inflammatory, anticancer, antimutagenic, and antifungal properties. It is also a potent antioxidant, reactive oxygen species scavenger and metal chelators (Olas & Wachowicz 2005). The extract contains many compounds including flavonoids, tanins, phenols, saponins, terpenoids, triterpenoids, steroids, alkaloids. Each fraction of velvet bean contains different compounds (Table 1). This result confirm the previous study that M. pruriens contained flavonoids (the subclass of flavonoid) (Handajani 2001). Eriodictyol and patuletin are flavonoids from Leuzea carthamoides inhibiting platelet aggregation with agonist collagen (COL) and arachidonic acid (AA) (Koleckar et al. 2008).

Rutin is a flavonoid inhibiting platelet aggregation in human platelets stimulated by COL agonist depend on its concentration (250 and 290 μM) (Sheu et al. 2004). The antiplatelet activity of rutin (flavonoid) may involve the following pathways: rutin inhibits the activation of phospholipase C, followed by inhibition of protein kinase C activity and TXA2 formation, thereby leading to inhibition of the phosphorylation of P47 and intracellular Ca2+ mobilization, finally resulting in inhibition of platelet aggregation (Mc Kenzie 2004; Sheu et al. 2004). TXA2 formation cause the bond between platelets is weak and platelet aggregation is reversible (primer aggregation), secondary aggregation need longer time and produces irreversible aggregation platelet (Mc Kenzie 2004). Supplementation studies with polyphenol-rich foods or extracts indicate that the compounds may exert effects in vivo as well. For instance, the ingestion of dealcoholized red wine, grape juice, or polyphenol extracts reduced blood pressure and inhibited platelet aggregation in laboratory animals (Erlund et al. 2008).

The flavonoid-rich beverages such as red wine and purple grape juice are vasodilators and improve endothelial function, probably because of a nitric oxide-dependent mechanism (Lin et al. 2007). Quercetin and catechin are flavonoids antioxidant, synergistically inhibit platelet function (in vitro assay) by blunting the release of hydrogen peroxide (H2O2) from platelets, subsequently reducing phospholipase C activation, calcium mobilization, and inositol phosphate synthesis. Flavonoids inhibit platelet aggregation because of their antioxidant activity, either by inhibiting the formation of endogenous mediators derived from phospholipid peroxidation, by blocking enzymatic free radical production, or by reducing platelet sensitivity to agonists by preventing lipid peroxidation (Murphy et al. 2003). Based on the experimental model of diabetes tested animal, the combination of acetylsalicylic acid and á-tocopherol as antioxidant led to beneficial changes that can help protect tissues from thrombotic incidents (Gonzales-Correa et al. 2006). Various agonists may stimulate platelet reactive oxygen species (ROS) production and aggregation, by regulating AA metabolism or via COX inhibition (Iuliano et al. 1997). In the presence of haemoglobin, ROS-induced platelet aggregation is enhanced (Iuliano et al. 1992). The resting platelets also generated a low amount of ROS. AA stimulates platelet ROS production, which is inhibited by hydroxychavicol flavonoid (HC), supporting HC flavonoid as a ROS scavenger (Chang et al. 2002). Our data (Table 2) showed that water fraction, butanol and ethyl acetate fraction of velvet bean seeds decreased platelet aggregation (as an antiplatelet activity) probably due to antioxidant activities. Previous research showed that M. pruriens had an ability to scavenge DPPH radicals, ABTS radicals and ROS. Velvet bean seeds significantly inhibit the oxidation of lipids and deoxyribose sugar. M. pruriens exhibited bivalent iron chelating activity (Dhanasekaran et al. 2008). Methanol extract of M. pruriens shows strong antioxidant activity by inhibiting DPPH and hydroxyl radicals, nitric oxide and scavenging superoxide anion, scavenging hydrogen peroxide, and reducing power activities when compared with different standards such as BHT, L-Ascorbic acid, curcumin, quercetin, and á-tocopherol (Rajeshwar et al. 2005). AA-induced thromboxane production or induced by the other agonists is not solely mediated by ROS production. The other possible reason
is that platelet ROS production can be mediated by COX as well as other enzymes such as platelet isoforms of NADPH oxidase, xanthine oxidase, mitochondrial respiration (Krotz et al. 2002). HC flavonoid inhibits the enzymes responsible for platelet ROS formation (Finazzi-Agro et al. 1982; Chang et al. 2007).

Aspirin showed higher antiplatelet activity compared with butanol and water fraction of velvet bean seeds, particularly at high concentration of epinephrine. Because aspirin is pure compound, it will produce high antiplatelet activity. The butanol and water fractions of velvet bean seeds contain complex compounds. To obtain the pure compound with high antiplatelet activity is required on further study.

The ethyl acetate, butanol, and water fraction contained high flavonoid and moderate phenol. The water, butanol, and ethyl acetate fraction of velvet bean seeds exhibited potential inhibition of EPN induced platelet aggregation at all concentrations. The strongest antiplatelet was water fraction and had similar antiplatelet activities with aspirin at level 150, 75, and 30 M of EPN inducer. Butanol fraction had similar antiplatelet activity with aspirin at lowest EPN inducer (30 μM).


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