IN VITRO ANTIDIABETIC ACTIVITIES OF EXTRACT AND ISOLATED FLAVONOID COMPOUNDS FROM Artocarpus altilis (Parkinson) FOSBERG

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ABSTRACT

The antidiabetic activity test through a mechanism of inhibition of α-glucosidase enzyme was studied against ethanol, n-hexane, ethyl acetate and n-butanol fractions of ethanol extract of Artocarpus altilis (Parkinson) Fosberg (Moraceae) leaves and four flavonoid compounds isolated from ethyl acetate extracts of A. altilis. Ethyl acetate fraction has strongest antidiabetic activity compared to ethanol, n-hexane, and n-butanol fractions with IC₅₀ values 5.98, 6.79, 440.18 and 14.42 μg/mL, respectively. Four flavonoid compounds (1-(2,4-dihydroxyphenyl)-3-[8-hydroxy-2-methyl-2-(4-methyl-3-pentenyl)-2H-1-benzopyran-5-yl]-1-propanone (AC-31), 2-geranyl-2',3,4,4'-tetrahydroxydihydrochalcone (AC-51), 8-geranyl-4',5,7-trihydroxyflavone (AC-33) and cyclocommunol (AA-3), have been isolated from ethyl acetate fraction. AC-31 was the strongest antidiabetic compound compared to AC-51, AC-33 and cyclocommunol with IC₅₀ values are 15.73, 24.41, 49.49, and 72.20 μg/mL. Kinetic studies of AC-31 using Lineweaver-Burk method showed that inhibition mechanism of enzyme α-glucosidase was a non-competitive type. Keywords: Artocarpus altilis; antidiabetic; α-glucosidase; flavonoid

INTRODUCTION

Diabetes mellitus (DM) is a chronic disease caused by inherited and/or acquired deficiency in production of insulin by the pancreas, or by the ineffectiveness of the insulin produced [1]. DM can be classified into 2 types: DM type 1 (insulin-dependent diabetes mellitus or IDDM) and DM type 2 diabetes (non-insulin dependent diabetes mellitus or NIDDM). In DM type 2, insulin-resistance is a characteristic feature and several drugs to increase the insulin sensitivity are currently being used in clinic [2]. At the present time, it is estimated that 150 million people worldwide have diabetes and that this will increase to 220 million by 2010 and 300 million by 2025 [3].

Indonesian health research foundation in 2007 showed that DM is a serious health problem for Indonesian peoples with 14.7% proportion of deaths due to diabetes disease in the age 45-54 years. DM is a debilitating and often life-threatening disorder with increasing incidence throughout the world [2]. Diabetic complications arise partly from glycosylation damage to structural and functional proteins and reflect chronic failure to maintain blood glucose homeostasis. Other complications such as diabetic nephropathy, diabetic retinopathy, diabetic neuropathy and diabetic cardiomyopathy prevail as a result of hyperglycemia. The pathophysiology involves an interaction between metabolic and hemodynamic...
factors. Metabolic factors include advanced glycation, increased formation of polyols and activation of protein kinase-C. Hemodynamic factors include systemic hypertension, intraglomerular hypertension and the role of vasoactive hormones, such as angiotensin II. Clinical course progresses from microalbuminuria to overt proteinuria and then to renal failure [5].

Dietary intake is related to blood glucose levels, and it is considered to be important to suppress hyperglycemia induced by overeating in order to prevent worsening of the symptoms in the hyperglycemic individuals with uncertain diabetic status. Inhibition of α-glucosidase is one way that could do in order to suppressing postprandial hyperglycemia [6-7] and reducing chronic vascular complications [8]. Therefore, by using α-glucosidase inhibitors is a cost-effective means to preventing the progression of diabetes [9-10].

Several inhibitors of this enzyme have been reported such as acarbose, voglibose, miglitol [11], nojirimycin and 1-deoxynojirimycin from plants [12]. Plants as sources of medicinal compounds have long been used by people. With a wealth of biodiversity, Indonesia has potential resources of various plants with various active compounds content which could inhibit the activity of enzyme α-glucosidase to overcome the disease diabetes [3-4]. Artocarpus altillis (Parkinson) Fosberg is a tropical plant, widely distributed in the tropics regions including Indonesia. Various parts of the plant, from leaves, latex, bark, roots were used as traditional medicine for various diseases ranging from infectious diseases (skin, urinary tract, and ear), diarrhea, dysentery, oral inflammation, fever, liver to diabetes and hypertension [13]. In vivo experiments showed that ethanol extract of A. altillis leaves (10 and 20 mg/kg) can reduced blood glucose levels significantly [16]. In this study, we investigated the inhibitory effect of A. altillis leaves extracts and four isolated flavonoid compounds on α-glucosidase activity. We also investigated kinetics inhibition reaction of the compounds.

EXPERIMENTAL SECTION

Materials

Artocarpus altillis leaves was collected from Parung, Bogor, Indonesia in March 2006 and determined at Herbarium Bogoriense, Research Center for Biology, Indonesian Institute of Sciences.

Four flavonoid compounds isolated from A. altillis (collection of Research Center for Chemistry, Indonesian Institute of Sciences) AC-31 (1-(2,4-dihydroxyphenyl)-3-[8-hydroxy-2-methyl-2-(4-methyl-3-pentenyl)-2H-1-

benzopyran-5-yl]-1-propanone), AC-51 (2-geranyl-2',3,4,4'-tetrahydroxydihydrochalcone), AC-33 (8-geranyl-4',5,7-trihydroxyflavone) and cyclocommunal (AA-3) [13-14]. Ethanol, n-hexane, ethyl acetate, n-butanol,

enzyme α-glucosidase, p-nitrofenil-α-D-glukopiranoidia (p-NPG), bovine serum albumin and querctin from Sigma. Solution of sodium bicarbonate and Dimethylsulfoxide (DMSO) from Merck.

Instrumentation

In this study, we used various instruments to support our research such as: macerator, rotary evaporator, and ELISA reader.

Procedure

Extraction

Powdered and dried of A. altillis leaves (4.95 Kg) were extracted exhaustivly with ethanol 70%. The ethanol extracts (250 g) were concentrated using vacuum rotary evaporator and then partitioned with n-hexane, ethyl acetate and n-butanol.

α-Glucosidase activity. Xa-glucosidase enzyme activity was analyzed using Sancheti method (Sancheti et al., 2009) with modifications. 0.001% enzyme stock solution dissolved in phosphate buffer solution (pH 7) contained 0.2% bovine serum albumin. Enzymatic reactions by mixing 25 μL 20 mM p-NPG as the substrate, 45 μL phosphate buffer solution and 5 μL samples were then added to the homogeneous α-glucosidase enzyme was diluted 50 times in phosphate buffer solution. The reaction mixture then water-bath incubated at 37 °C for 30 min. The reaction was stopped by the addition of 100 μL 0.2 M of Na2CO3 solution. The resulted p-nitrophenol was measured at λ 400 nm using ELISA reader. The experiments were performed with 3 replicates and % inhibition was calculated using the following formula:

\[ \text{%Inhibition} = \left( \frac{C - S}{C} \right) \times 100\% \]

C absorbance without sample

S absorbance with the sample

Kinetics inhibition against α-glucosidase enzyme

Kinetics inhibition against α-glucosidase enzyme was analyzed by measuring the formation of p-NP on the time interval 0 to 20 min with a variety of p-NPG substrate concentrations between 1 to 20 mM with or without samples. Linewaver-Burk method used to determine the type of inhibition to create a graph plot 1/v (speed of reaction) against 1/[S] (substrate concentration changes are directly proportional to the product formation).
RESULT AND DISCUSSION

α-Glucosidase inhibition activities of each sample was compared by IC$_{50}$ value resulted in the experimental condition. Smaller IC$_{50}$ values mean stronger inhibition activity. Ethanol extract showed stronger inhibition activity against α-glucosidase compared with quercetin as positive control with IC$_{50}$ 6.79 μg/mL. Further fractionation of the ethanol extract showed that the ethyl acetate fraction was the strongest fraction with IC$_{50}$ 5.98 μg/mL (Table 1). This result suggests that the active compounds from the ethanol extract concentrated on semi polar fraction, and flavonoids were the active major compounds contained in A. altilis plant.

### Table 1. Enzyme α-glucosidase inhibition activity of A. altilis leaves ethanol extract and its fractions.

<table>
<thead>
<tr>
<th>Compound</th>
<th>IC$_{50}$ (μg/mL)</th>
</tr>
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<tbody>
<tr>
<td>Quercetin</td>
<td>11.60</td>
</tr>
<tr>
<td>Ethanol</td>
<td>6.79</td>
</tr>
<tr>
<td>n-Hexane</td>
<td>440.18</td>
</tr>
<tr>
<td>Ethylacetate</td>
<td>5.98</td>
</tr>
<tr>
<td>n-Butanol</td>
<td>14.42</td>
</tr>
</tbody>
</table>

AC-31 (1-(2,4-dihydroxyphenyl)-3-[8-hydroxy-2-methyl-2-(4-methyl-3-pentenyl)-2H-1-benzopyran-5-yl]-1-propanone), yellow crystal, ¹³C-NMR (in CDCl$_3$, ppm) δC 165.2 (C-1), 103.6 (C-2), 163.48 (C-3), 108.3 (C-4), 132.45 (C-5), 113.6 (C-6), 204.1 (C-7), 39.8 (C-8), 26.7 (C-9), 128.8 (C-10), 121.29 (C-11), 114.7 (C-12), 143.1 (C-13), 139.7 (C-14), 119.2 (C-15), 119.5 (C-16), 130.3 (C-17), 78.9 (C-18), 40.9 (C-19), 22.9 (C-20), 123.9 (C-21), 132.1 (C-22), 25.8 (C-23), 17.8 (C-24), 26.2 (C-25). Molecular weight with [M+] 409.5460 [17].

AC-51 (2-geranyl-2',3,4,4'-tetrahydroxydihydro chalcone), yellow gum, ¹³C-NMR (in CDCl$_3$, ppm) δC 131.1 (C-1), 126.1 (C-2), 142.4 (C-3), 142.6 (C-4), 112.9 (C-5), 121.3 (C-6), 113.5 (C-1'), 164.9 (C-2'), 103.5 (C-3'), 163.0 (C-4'), 108.0 (C-5'), 132.2 (C-6'), 204.0 (C=O), 39.6 (α), 27.7 (β), 25.8 (C-1'), 121.7 (C-2'), 138.6 (C-3'), 39.6 (C-4'), 26.3 (C-5'), 123.7 (C-6'), 132.1 (C-7'), 25.7 (C-8'), 17.7 (C-9'), 16.2 (C-10') [19].

AC-33 (8-geranyl-4',5,7-trihydroxyflavone), white crystal, ¹³C-NMR (in CD$_3$OD, ppm) δC 5.3 (1H, dd, H-2); 2.70 (1H, dd H-3); 3.06 (1H, dd, H-3b); 5.93 (1H, s, H-6); 3.18 (2H, dd, H-9); 5.14 (1H, t, H-10); 1.91 (2H, t, H-12); 2.02 (2H, t, H-13); 5.04 (1H, t, H-14); 1.55 (3H, H-20); 5.08 (1H, t, H-21); 1.66 (3H, s, H-23); 1.57 (3H, s, H-24); 1.28 (3H, s, H-25).
Table 2. Enzyme α-glucosidase inhibition activity of the isolated flavonoid compounds from A. altilis leaves

<table>
<thead>
<tr>
<th>Compound</th>
<th>IC₅₀ (µM)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Quercetin</td>
<td>38.40</td>
</tr>
<tr>
<td>AC-51</td>
<td>24.41</td>
</tr>
<tr>
<td>AC-31</td>
<td>15.73</td>
</tr>
<tr>
<td>AC-33</td>
<td>49.49</td>
</tr>
<tr>
<td>AA-3</td>
<td>72.20</td>
</tr>
</tbody>
</table>

Table 3. Vm and Km values of the enzymatic reactions of α-glucosidase with and without inhibitor (AC31)

<table>
<thead>
<tr>
<th>Condition</th>
<th>Vm (mM/mnt)</th>
<th>Km (mM)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Without Inhibitor</td>
<td>0.033±0.009</td>
<td>3.246±2.242</td>
</tr>
<tr>
<td>AC-31 1.5 mM</td>
<td>0.012±0.001</td>
<td>4.262±0.413</td>
</tr>
<tr>
<td>AC-31 3 mM</td>
<td>0.009±0.002</td>
<td>2.934±0.170</td>
</tr>
</tbody>
</table>

CONCLUSION

Enzyme α-glucosidase inhibition assay of Artocarpus altilis leaf extract shown that ethanol extract, ethyl acetate fraction and two flavonoid compounds (AC-31 and AC-51) have antidiabetic activity stronger than quercetin as positive control with IC₅₀ value 6.79 μg/mL, 5.98 μg/mL, 15.73 μM, and 24.41 μM, respectively. From the kinetic inhibition of enzyme α-glucosidase mechanism can be concluded that the type of inhibition of AC-31 compound is a non-competitive.

ACKNOWLEDGEMENT

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REFERENCES