In-vitro studies on antioxidant and antidiabetic potential of Sesoot (Garcinia picrorrhiza Miq.) fruit ethanolic extract from Indonesia

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ABSTRACT

Diabetes mellitus (DM) is a disease that can be identified by high levels of blood glucose. Garcinia plants have been widely used for many traditional medicines as antioxidant, anticancer, antidiabetics, and antiinflammation. The antioxidant and antidiabetic activities of (Garcinia picrorrhiza Miq.) or sesoot fruit extract were evaluated in this study and compared with xanthone. The antioxidant and antidiabetic of ethanolic ripe sesoot (G. picrorrhiza Miq.) fruit extract (GpKar) was evaluated by 2,2-Azinobis 3-ethyl benzothiazoline 6-sulfonic acid (ABTS•+) reducing activity, α-Glucosidase, β-Glucosidase, and α-amylase inhibitor activity. GpKar showed higher ABTS•+ -reducing activity (IC₅₀ = 49.30 µg/mL) than xanthone (IC₅₀ = 404.30 µg/mL). GpKar showed IC₅₀ = 109.32 µg/mL for α-glucosidase inhibitory activity, while xanthones had a better activity (IC₅₀ = 33.97 µg/mL). GpKar also showed lower α-amylase inhibitory activity and β-Glucosidase (IC₅₀ = 126.01 and 9432.09 µg/mL) compared to xanthone (IC₅₀ = 44.32 and 405.03 µg/mL, respectively). The compounds of GpKar are proven to have antioxidant and antidiabetic activities. Therefore, it will be industrially relevant to develop a natural medicine for decreasing DM risk, thus evaluating the antioxidant and antidiabetic effect of G. picrorrhiza by a pre-clinic study is needed.

Keywords: antidiabetic, antioxidant, Garcinia picrorrhiza, xanthone

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INTRODUCTION

In 2019, there are about 463 millions people around the world have been diagnosed with diabetes. Moreover, the International Diabetes Federation (IDF) said that diabetes cases are anticipated to grow higher up to 700 millions by 2040 (Pasupuleti et al., 2020). Diabetes is a disease stimulated by high levels of blood glucose (hyperglycemia). Hyperglycemia is a condition in which insulin secretion and/or insulin action experiences some defects (American Diabetes Association, 2013). Diabetes is divided into two types and Type 2 diabetes mellitus (T2DM) can lead to various illnesses and premature death (Pasupuleti et al., 2020). Defect of insulin action and production leads to long-term health problems. In addition to insulin, defects of carbohydrates, lipids, proteins, and biomolecules also play a role in health problems (American Diabetes Association, 2013). α-amylase and α-glucosidase are enzymes that stimulate the starch breakdown which is the main source of increased blood sugar levels. Blood glucose levels can be decreased by inhibiting the enzymes and it is an essential strategy in diabetes (Lordan et al., 2013; Widowati et al., 2018).

Diabetes can cause activation of several pathological diseases, chronic hyperglycemia, and free radical generation (Yaribeygi et al., 2019). Free radicals can be neutralized by electron donation from antioxidants. Catalase (CAT), glutathione peroxidase (GPX), superoxide dismutase (SOD) are some antioxidants that counter the deleterious effect of free radicals. These antioxidants are produced in the body or provided through the diet. Furthermore, traditional medicinal plants also have antioxidants (Kumar et al., 2021). Antioxidant plays a protective role in diabetic pathologies and cardiovascular disease.

Garcinia is one of the Clusiaceae family which is native to Asia, Australia, and tropical regions. Garcinia fruit is a nutrient-rich source, minerals, vitamins, and dietary fiber. In several studies, Garcinia plants have been widely used as a traditional natural medicine for antioxidant, anticancer, antidiabetic, and antiinflammation, due to their good source of bioactive compounds (Aisha et al., 2013; Murthy et al., 2019). G. picrorrhiza Miq. grows in tropical to temperate climates and in the original region known as sesoot (Utami et al., 2017). It contains bioactive compounds including xanthones, benzophenones, hydroxycitric acid, and anthocyanins (Espirito Santo et al., 2020). Xanthones are one of the compounds highly found in natural sources, including plants, fungi, ferns, and lichens. Xanthones could be found in several plant families, including Garcinia. Their pharmacological properties have raised a great interest (Negi et al., 2013) and xanthones themselves are polyphenolic compounds that have been widely used in the treatment of DM (Putri, 2015).

Based on the previous study, the extract of G. picrorrhiza Miq. has antioxidant activity and anticholinesterase activity which is potentially used to decrease Alzheimer’s disease progression (Utami et al., 2017). Another study related to the Garciniaceae reported that extract of G. mangostana L. has antidiabetic activity by decreasing cholesterol levels and lipidic peroxidation (Husen et al., 2017); however, the effect of G. picrorrhiza Miq has not been much reported. The genus Garcinia has more than fifty species. However, G. picrorrhiza Miq. is one of the species which is rarely studied. Based on these facts, more comprehensive studies are needed to investigate another bioactivity on G. picrorrhiza Miq. extract. The potential of ripe fruit G. picrorrhiza Miq. for DM can be investigated by evaluating the antioxidant activity with the ABTS+ method, while the antidiabetic activity was evaluated by α-glucosidase, α-amylase, and β-Glucosidase assay.

MATERIALS AND METHOD

Material

Ripe fruits of G. picrorrhiza Miq. were harvested from Bogor Botanical Garden and identified in the herbarium of the School of Life Science and Technology, Bandung Institute of Technology, West Java, Indonesia. The enzymes were α-glucosidase from Saccharomyces sp. (Sigma Aldrich G5003), β-Glucosidase (Sigma-Aldrich G4511), and α-amylase enzyme (Sigma Aldrich A7595). p-nitrophenyl α-D-glucopyranoside (Sigma Aldrich N1377), p-nitrophenyl-β-D-glycopyranoside...
(Sigma-Aldrich N7006). ABTS$^+$ (Sigma Aldrich, A1888). Pure xanthone (Sigma Aldrich X0626) was used in this experiment as a standard compound.

**Method**

**Preparation of G. picrorrhiza extract**

Ripe fruits of G. picrorrhiza Miq. (1330 g) were ground and immersed in ethanol 70% for 24 hours. The extraction was carried out twice. This procedure was redone until the colorless filtrate was obtained. The collected filtrates were subsequently evaporated at 40°C using a rotary evaporator, and the extract (GpKar) was kept at -20 °C (Prahastuti et al., 2020; Rusmana et al., 2017; Widowati et al., 2016, 2018).

**ABTS$^+$ reducing-activity**

The ABTS$^+$ radical (ABTS$^+$) has a bluish-green color (More and Makola, 2020) and absorbs at 743 nm. The color was formed when an electron loses because of the nitrogen atom of ABTS$^+$. ABTS$^+$ itself can be oxidized using potassium persulfate or manganese dioxide (Xiao et al., 2020). Then, 14 mM ABTS$^+$ was added into 4.9 mM potassium persulfate to obtain 7 mM ABTS$^+$ in 2.45 mM potassium persulfate. It was then incubated for 16 hours at room temperature and dark conditions. After that, 5.5 mM PBS (pH 7.4) was added to dilute the ABTS$^+$ solution, and the absorbance at 745 nm using a microreader was measured. 198 µL of ABTS$^+$ solution was added with 2 µL of GpKar, xanthones in various concentration (3.13, 6.25, 12.5, 25, 50, and 100 µg/mL). The mixture was kept for 6 minutes at 30°C while the absorbance was measured at 745 nm using Multiscan™ GO Microplate Spectrophotometer, Thermo Scientific (Prahastuti et al., 2020). The ABTS$^+$- reducing activity (%) was then determined as median inhibitory concentration (IC$_{50}$) (Prahastuti et al., 2020; Widowati et al., 2016, 2018).

**α-Glucosidase inhibitory activity assay**

The activity of α-glucosidase inhibitory was evaluated with a method taken from the previous study (Gondokesumo et al., 2017; Widowati et al., 2018) with certain modifications. The solution of the α-glucosidase enzyme was produced from 1 mg of α-glucosidase from Saccharomyces sp. (Sigma Aldrich G5003), diluted in 100 mL phosphate buffer (pH 7.0) that contained 200 mg bovine serum albumin (BSA). The reaction mixture contained 20 mM p-nitrophenyl α-D-glucopyranoside (Sigma Aldrich N1377), 45 µL phosphate buffer, and 5 µL of GpKar, xanthone (7.81, 15.63, 31.25, 62.5, 125, and 250 µg/mL). After the reaction mix was homogenized, 25 µL α-glucosidase was added before incubation (37°C in 30 minutes). Na$_2$CO$_3$ was added after the incubation. The α-glucosidase will hydrolyze the p-nitrophenyl-α-D-glucopyranoside to glucose and p-nitrophenol (yellow color) (Widowati et al., 2018). The released p-nitrophenol was then determined using a microplate reader at 400 nm.

Formula 1 was used to calculate α-glucosidase assay as below:

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

Where $C$ = absorbance value of the negative control

$S$ = absorbance value of the sample

**β-Glucosidase inhibitory activity assay**

The activity of β-Glucosidase inhibitory was evaluated with a method taken from the previous study (Gondokesumo et al., 2017; Widowati et al., 2015) with certain modifications. The solution of the β-Glucosidase enzyme was produced from 1 mg of β-Glucosidase (Sigma Aldrich G4511), diluted in 100 mL phosphate buffer (pH 7.0) that contained 200 mg BSA. The reaction mixture contained 200 mM p-nitrophenyl-β-D-glycopyranoside (Sigma Aldrich N7006), 990µL of 100 mM phosphate buffer
(pH 7.0), and 10 µL of GpKar, and xanthone (7.81, 15.63, 31.25, 62.5, 125, and 250 µg/mL). For 5 minutes, the reaction mixture was kept at 37°C, and 500 µL of enzyme mixture was added before the second incubation (15 minutes at 37°C). The reaction was then ended by supplementation of 200 µL Na₂CO₃ (200 mM). The β-Glucosidase hydrolyzed the p-nitrophenyl-β-D-glucopyranoside, and the product was measured at 405 nm (Gondokesumo et al., 2017; Widowati et al., 2018).

A microplate reader was used to calculate the absorbance at 405 nm while the inhibition of β-Glucosidase was evaluated according to formula 2:

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

Where  
C = absorbance value of the negative control
S = absorbance value of the sample

α-amylase inhibitory activity assay

The assay of α-amylase inhibitory activity was done according to the breakdown of the substrate to generate a colored product and the absorbance was measured over a certain period (Utami et al., 2019; Widowati et al., 2015, 2018). For this assay, dimethyl sulfoxide (DMSO) was used as the blank, 30 µL of GpKar, xanthone (20.83, 41.67, 83.33, 166.67, 333.33, and 666.67 µg/mL) and 10 µL of α-amylase enzyme 250 units/g (Sigma Aldrich 7595) were put into the well. The reaction mixture was then incubated for 10 minutes at 37°C. For the treatment group, 40 µL starch solution was added after incubation while for the control group 40 µL phosphate buffers were supplemented. The reaction mixture was again kept for another 15 minutes at 37°C. The 100 µL acidic iodine solution (except black) was also added to terminate the enzymatic reaction after incubation. Using a microplate reader, the absorbance was then measured at 565 nm (Utami et al., 2019; Widowati et al., 2018). Formula 3 was used to calculate α-amylase assay, seen as below:

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

Where  
C = absorbance value of the negative control
S = absorbance value of the sample

Data Analysis

All data measurements were done three times. Data are shown as means ± standard deviation. In this study, the statistical analysis was performed using the SPSS Statistics 20.0 computer program. The analysis used the One Way ANOVA, followed by Tukey’s HSD Post Hoc Test. The significance value was determined with a p-value of ≤ 0.05.

RESULT AND DISCUSSION

Sesoot fruit is one of those natives to Asia and Africa which contains many bioactive compounds including xanthones, flavonoids, triterpenoids, and benzophenones (Gutierrez-Orozco and Failla, 2013). Xanthones are polyphenol groups commonly found in mangosteen. Xanthones contain α-mangostin, γ-mangostin, and another minor compound (Mohammad et al., 2019). Xanthones from Garcinia fruit were known to exhibit pharmacologic properties such as antioxidant (Gondokesumo et al., 2019), antidiabetic (Ewenighi et al., 2015), and anti-inflammatory (Mohan et al., 2018). The ABTS•⁺-reducing activity of GpKar and xanthone is shown in Figure. 1, and the IC₅₀ of samples in ABTS•⁺-reducing activity are shown in Table 1.
In-vitro Studies ... (Utami et al.)
Based on Figure 2, both GpKar and xanthone showed α-glucosidase inhibitory activity depending on concentration-manner. Based on Table 2, GpKar showed higher IC$_{50}$ value (109.32 µg/mL) compared to xanthone (33.97 µg/mL); xanthone was categorized highly active (Marjoni and Zulfisa, 2017).

Table 2. The IC$_{50}$ value of α-glucosidase, β-Glucosidase, and α-amylase inhibitory activities of ethanolic ripe sesoot (G. picrorhiza Miq.) fruit extract (GpKar) and xanthone

| Samples | α-glucosidase inhibitory activity | β-Glucosidase inhibitory activity | α-amylase inhibitory activity |
|---------|----------------------------------|----------------------------------|-------------------------------|
|         | IC$_{50}$ (µg/mL) | IC$_{50}$ (µg/mL) | IC$_{50}$ (µg/mL) |
| GpKar   | 109.32             | 126.01            | 9432.09                       |
| Xanthone| 33.97              | 44.32             | 405.03                        |

Based on Figure 2, both GpKar and xanthone showed α-glucosidase inhibitory activity depending on concentration-manner. Based on Table 2, GpKar showed higher IC$_{50}$ value (109.32 µg/mL) compared to xanthone (33.97 µg/mL); xanthone was categorized highly active (Marjoni and Zulfisa, 2017).

Figure 2. The α-Glucosidase inhibitory activity of ethanolic ripe sesoot (G. picrorhiza Miq.) fruit extract (GpKar) and xanthone. The data are explained as means ± standard deviation with different letters (A, B, C, D, E, and F) for GpKar and different letters (a, b, c, d, e, and f) for xanthone, representing significant differences in Tukey HSD post hoc with p value ≤ 0.05.

Figure 3. The β-Glucosidase inhibitory activity of ethanolic ripe sesoot (G. picrorhiza Miq.) fruit extract (GpKar) and xanthone. The data are explained as means ± standard deviation with different letters (A, B, C, D, E, and F) for GpKar and different letters (a, b, c, d, and e,) for xanthone, representing significant differences in Tukey HSD post hoc with p value ≤ 0.05.
Based on Figure 3, both GpKar and xanthone showed β-Glucosidase inhibitory activity depending on concentration-manner. Based on Table 2, GpKar showed a higher IC₅₀ value (126.01 µg/mL) than xanthone (44.32 µg/mL) in inhibition of β-Glucosidase. This result indicates that GpKar has a lower β-Glucosidase inhibitory activity than xanthone. Xanthones are categorized as highly active in β-Glucosidase activity based on these results (Marjoni and Zulfisa, 2017). Xanthones from mangosteen have been proven to suppress glucose absorption and potential compounds for in vitro α-glucosidase inhibition (Ryu et al., 2011). The extract of *G. mangostana* pericarp demonstrated a mild lowering effect on glucose level in normoglycemic animals, which was comparable with glibenclamide (standard drug) (Taher et al., 2016). The antioxidant compounds such as epicatechin and xanthones may play a role in this effect. Therefore, *G. mangostana* extract is also a strong candidate for DM management (Taher et al., 2016).

![Figure 4](image_url)

**Figure 4.** The α-amylase inhibitory activity of ethanolic ripe sesoot (*G. picrorrhiza* Miq.) fruit extract (GpKar) and xanthone. The data are explained as means ± standard deviation with different letters (A, AB, BC, C, and D) for GpKar and different letters (a, ab, bc, c, and d) for xanthone, representing significant differences in Tukey HSD post hoc with *p* value ≤ 0.05

Based on Figure 4, the α-amylase inhibitory activity of GpKar and xanthone showed a concentration dependent-manner. Table 2 shows that the IC₅₀ value of GpKar was higher (9432.09 µg/mL) than xanthone (405.03 µg/mL). Xanthone was categorized weak in α-amylase inhibitory activity and GpKar was categorized inactive (Marjoni and Zulfisa, 2017).

There was a report stated that extract of *G. mangostana* can deaccelerate the absorption of glucose by inhibiting the carbohydrate hydrolyzing enzymes like α-amylase (Manaharan et al., 2012). The result in this present study has been in line with the previous study. Inhibition of amylase and glucosidase enzymes is proven to lower the postprandial elevation of blood glucose after a mixed carbohydrate diet (Thamizharasan and Umamaheswa, 2016). Meanwhile, *G. kola* possesses hypoglycemic properties and significantly lowered blood glucose in alloxan-induced diabetic rats (Ewenighi et al., 2015).

Reactive oxygen species (ROS) and reactive nitrogen species (RNS) are included to free radicals. Unstable electrons in free radicals react with lipids, proteins, and/or DNA resulting in various damages. Oxidative damage happens when free radicals and antioxidant agents are imbalanced (Ayepola et al., 2014). However, oxidative stress affects various diseases such as diabetes, vascular and neural disorders (Pasupuleti et al., 2020). Various mechanisms of oxidative stress play a role in diabetes, including excess oxygen radicals, glycated protein formation, and the glycation of antioxidant enzymes. This will limit the ability of antioxidants to prevent free radicals (Pasupuleti et al., 2020). The previous study showed that various group compounds such as phenolic, flavonoids, anthocyanins, and carotenoids could have a contribution to the antioxidant activity in the extracts of *G.*

*In-vitro Studies* ...(Utami et al.,)
parvifolia fruit (Hassan et al., 2013). Moreover, Hassan stated that the sum up phenolic content strongly correlated with the activity of antioxidant (Hassan et al., 2013). The quality of antioxidant properties relates to various factors, and the major factors that affect the activities are the maturation stage, extraction solvent, and materials preparations (Suttirak and Manurakchinakorn, 2014).

CONCLUSION
The extract of sesoot fruit has highly active antioxidants while xanthone has weak antioxidant properties. The extract of sesoot has moderate $\alpha$ and $\beta$-Glucosidase inhibition and inactive $\alpha$-amylase inhibition properties. On the other hand, xanthone has highly active $\alpha$ and $\beta$-Glucosidase inhibition and weak $\alpha$-amylase inhibition properties. Further research to develop its activity in antidiabetic prevention by pre-clinic studies is needed.

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