

Pharmaciana

Vol.11, No.2, July 2021, Page. 273-282

ISSN: 2088 4559; e-ISSN: 2477 0256

DOI: 10.12928/pharmaciana.v11i2.20110

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In-vitro studies on antioxidant and antidiabetic potential of Sesoot (*Garcinia picrorrhiza* Miq.) fruit ethanolic extract from Indonesia**Sri Utami^{*1}, Sonny Pamuji Laksono¹, Qomariyah Romadhiyani Sachrowardi¹, Dewi Nurul Mustaqimah¹, Susi Endrini¹, Ndaru Andri Damayanti¹, Said Nafik², Betharie Cendera Arrahmani³, Hanna Sari Widya Kusuma⁴, Wahyu Widowati⁵**¹Faculty of Medicine, YARSI University,

Jl. Letjend Suprpto, Cempaka Putih, Jakarta Pusat 10510, Indonesia

²Directorate General of Intellectual Property, Ministry of Law and Human Rights,

Jl. H.R. Rasuna Said, Kuningan, Jakarta Selatan, Indonesia

³Master Program of Pharmaceutical Bioprocessing Engineering, School of Life, Sciences,

Technical University of Munich, Alte Akademie 8, 85354 Freising, Germany

⁴Aretha Medika Utama, Biomolecular and Biomedical Research Center

Jl. Babakan Jeruk 2 No. 9, Bandung 40164, Indonesia.

⁵Faculty of Medicine, Maranatha Christian University,

Jl. Prof. Drg. Surya Sumantri No. 65 Bandung 40164, Indonesia

Submitted: 24-02-2021

Reviewed: 24-05-2021

Accepted: 30-07-2021

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 μ g/mL) than xanthone (IC₅₀ = 404.30 μ g/mL). GpKar showed IC₅₀ = 109.32 μ g/mL for α -glucosidase inhibitory activity, while xanthenes had a better activity (IC₅₀ = 33.97 μ g/mL). GpKar also showed lower α -amylase inhibitory activity and β -Glucosidase (IC₅₀ = 126.01 and 9432.09 μ g/mL) compared to xanthone (IC₅₀ = 44.32 and 405.03 μ g/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***Corresponding author:**

Sri Utami

Faculty of Medicine, YARSI University

Jl. Letjend Suprpto, Cempaka Putih, Jakarta Pusat 10510, Indonesia

Email: uutsuyono@yahoo.com

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 metabolisms 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 xanthenes, benzophenones, hydroxycitric acid, and anthocyanins (Espirito Santo et al., 2020). Xanthenes are one of the compounds highly found in natural sources, including plants, fungi, ferns, and lichens. Xanthenes could be found in several plant families, including *Garcinia*. Their pharmacological properties have raised a great interest (Negi et al., 2013) and xanthenes 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 *Garcinias* 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 rotatory 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, xanthenes 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 MultiscanTM GO Microplate Spectrophotometer, Thermo Scientific (Prahastuti et al., 2020). The ABTS^{•+}- reducing activity (%) was then determined as median inhibitory concentration (IC₅₀) (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). Na₂CO₃ 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:

$$\text{Inhibition (\%)} = \frac{C - Sx}{c} \times 100 \quad (1)$$

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 $\mu\text{g/mL}$). 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 Na_2CO_3 (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:

$$\text{Inhibition (\%)} = \frac{C - Sx}{c} \times 100 \quad (2)$$

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 $\mu\text{g/mL}$) 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:

$$\text{Inhibition (\%)} = \frac{C - Sx}{c} \times 100 \quad (3)$$

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 \pm 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 xanthenes, flavonoids, triterpenoids, and benzophenones (Gutierrez-Orozco and Failla, 2013). Xanthenes are polyphenol groups commonly found in mangosteen. Xanthenes contain α -mangostin, γ -mangostin, and another minor compound (Mohammad et al., 2019). Xanthenes 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.

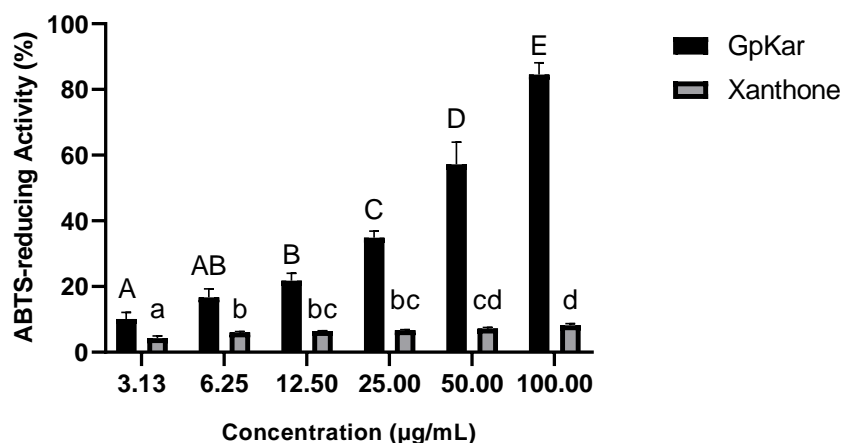


Figure 1. The ABTS^{•+} -reducing activity of ethanolic extract of ripe sesoot (*G. picrorrhiza* Miq.) fruit extract (GpKar) and xanthone. The data are explained as means \pm standard deviation with different letters (A, AB, B, C, D, and E) for GpKars and different letters (a, b, bc, cd, and d) for xanthone, representing significant differences in Tukey HSD post hoc with p value ≤ 0.05

Table 1. The IC₅₀ value of ABTS^{•+} -reducing activity of ethanolic extract of ripe sesoot (*G. picrorrhiza* Miq.) fruit extract (GpKar) and Xanthone

Sample	IC ₅₀ (µg/mL)
GpKar	49.30
Xanthone	404.30

Based on [Figure 1](#), both GpKar and xanthone showed an ABTS^{•+} reducing activity percentage which depended on the concentration-manner. The higher concentration of the sample shows the higher ABTS^{•+} reducing activity. Therefore, as seen in [Table 1](#), GpKar was shown to have a lower IC₅₀ value at 49.30 µg/mL than xanthone at 404.30 µg/mL. The IC₅₀ is the the sample concentration to collect 50% of the free radical. The lower the IC₅₀ value indicates the higher antioxidant activity, and GpKar was categorized highly-active antioxidant ([Marjoni and Zulfisa, 2017](#)).

This result conforms with the previous study where ABTS^{•+} antioxidant activity of methanolic extract *G. talbotii* from India had an IC₅₀ value of 40.09 µg/mL ([Kureshi et al., 2020](#)) which was in line with its ABTS^{•+} -reducing activity. Meanwhile, our previous study revealed that the H₂O₂ inhibitory scavenging activity of GpKar had a higher activity than xanthone ([Utami et al., 2017](#)). The high antioxidant activity of GpKar is probably caused by secondary metabolites. The major secondary metabolites in mangosteen are xanthenes and more than 68 xanthenes derivatives are isolated. The α -mangosteen and γ -mangosteen are the major xanthone derivates that show powerful antioxidant activity ([Aizat et al., 2019](#); [Tjahjani and Widowati, 2013](#)).

Plant extracts are commonly used to treat various diseases including diabetes because of their effectiveness, safety, and abundant sources ([Taher et al., 2016](#)). The α -glucosidase inhibitors delay the digestion as well as the complex carbohydrates absorption to keep blood glucose at lower levels ([Nisha, 2017](#)). The α - glucosidase and β -Glucosidase inhibitors are promising therapeutic agents, such as for diabetes ([Kumar et al., 2011](#)), breast cancer ([Zhou et al., 2017](#)), antitumor ([Li et al., 2018](#)), hepatitis B, and human immunodeficiency virus (HIV) ([Abid et al., 2016](#)). The result of the α -glucosidase inhibitory activity assay of GpKar and xanthone can be seen in [Figure 2](#). [Table 2](#) illustrates the results of the α -Glucosidase, β -Glucosidase, and α -amylase inhibitory activity by the IC₅₀ value.

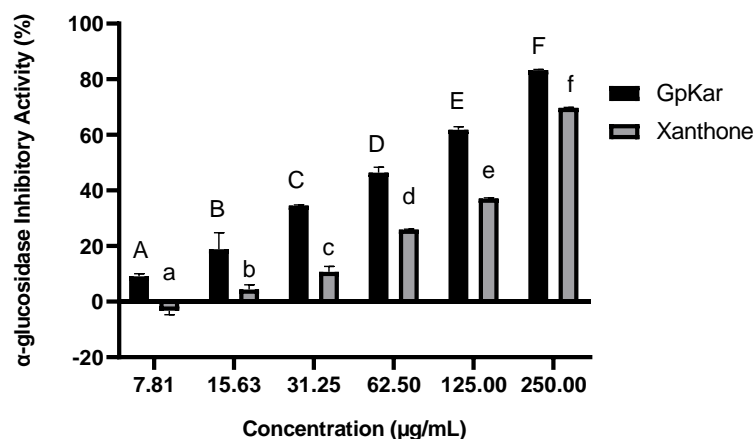


Figure 2. The α -Glucosidase inhibitory activity of ethanolic ripe sesoot (*G. picrorrhiza* Miq.) fruit extract (GpKar) and xanthone. The data are explained as means \pm standard deviation with different letters (A, B, C, D, E, and F) for GpKar and different letters (a, b, c, d, e, and f) for xanthone,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

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

Samples	α -glucosidase inhibitory activity	β -Glucosidase inhibitory activity	α -amylase inhibitory activity
	IC_{50} ($\mu\text{g/mL}$)	IC_{50} ($\mu\text{g/mL}$)	IC_{50} ($\mu\text{g/mL}$)
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 $\mu\text{g/mL}$) compared to xanthone (33.97 $\mu\text{g/mL}$); xanthone was categorized highly active (Marjoni and Zulfisa, 2017).

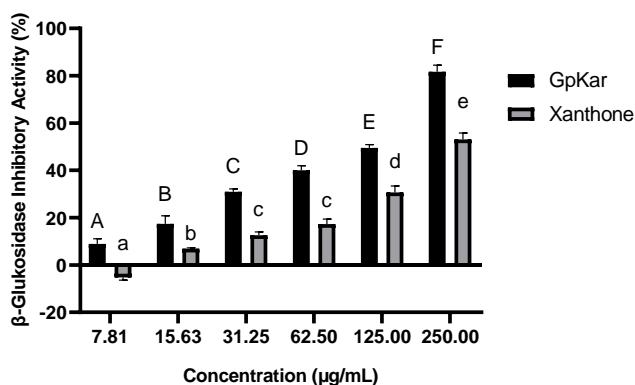


Figure 3. The β -Glucosidase inhibitory activity of ethanolic ripe sesoot (*G. picrorrhiza* Miq.) fruit extract (GpKar) and xanthone. The data are explained as means \pm standard deviation with different letters (A, B, C, D, E, and F) for GpKar and different letters (a, b, c, d, and e) for xanthone, 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 IC_{50} value (126.01 $\mu\text{g/mL}$) than xanthone (44.32 $\mu\text{g/mL}$) 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).

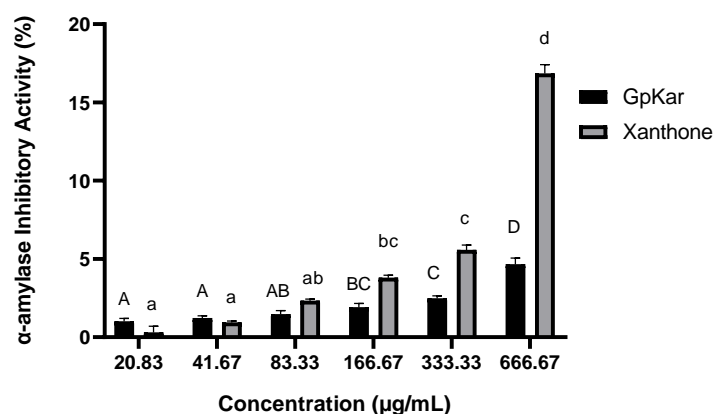


Figure 4. The α -amylase inhibitory activity of ethanolic ripe sesoot (*G. picrorrhiza* Miq.) fruit extract (GpKar) and xanthone. The data are explained as means \pm standard deviation with different letters (A, AB, BC, C, and D) for GpKar and different letters (a, ab, bc, c, and d) for xanthone, representing 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 IC_{50} value of GpKar was higher (9432.09 $\mu\text{g/mL}$) than xanthone (405.03 $\mu\text{g/mL}$). 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.*

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 α and β -Glucosidase inhibition and inactive α -amylase inhibition properties. On the other hand, xanthone has highly active α and β -Glucosidase inhibition and weak α -amylase inhibition properties. Further research to develop its activity in antidiabetic prevention by pre-clinic studies is needed.

ACKNOWLEDGEMENT

This study was supported financially by YARSI University for research grant 2016. Thanks to research methodology and laboratory facilities supported by Aretha Medika Utama Biomolecular and Biomedical Research Center, Bandung, Indonesia. The authors would also like to thank Ervi Afifah, Seila Arumwardana, Cintani Dewi Wahyuni, Cahyaning Riski Wijayanti, Muhamad Aldi Maulana from Biomolecular and Biomedical Research Center, Aretha Medika Utama for all their help in this project.

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