Antihypertensive, Antidiabetic, Antioxidant and Cytotoxic Activities of Indonesian Traditional Medicine

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ABSTRACT
Background: Indonesian people have long used herbal medicine (jamu) to overcome various diseases, including hypertension and diabetes mellitus. Hypertension and diabetes mellitus are two diseases that are directly related and require proper and thorough management. Objectives: The present study investigated the antihypertensive, antidiabetic, and cytotoxic activities ethanol extracts of Indonesian traditional medicine (jamu). Material and Methods: Jamu was extracted by maceration using ethanol. Antihypertensive and antidiabetic activity investigated by measurement of ACE inhibitor, an alpha-glucosidase inhibitor, and antioxidant activity at a concentration ranging from 125-1000 µg/mL, respectively, by in vitro method. Cytotoxic evaluation of the extract was carried out using Brine Shrimp Lethality Test (BSLT). Results: measurements of ACE inhibitors, alpha-glucosidase inhibitor and antioxidant activity showed that herbal extracts had ACE inhibitors, alpha-glucosidase inhibitors, and antioxidant activity with IC50 values of 292.15 µg/mL, 36.13 µg/mL, and 24.43 µg/mL respectively. Ethanol extract of herbal medicine (jamu) exerts a cytotoxic effect on larvae of shrimp Artemia salina with an IC50 value of 215.04 µg/mL. Conclusion: Jamu extract has antihypertensive and antidiabetic activity in vitro and cytotoxic effects.

Key words: Jamu, ACE inhibitors, Antioxidant, Alpha-glucosidase inhibitors, Cytotoxic.

INTRODUCTION
Herbal medicines are types of alternative medicine already used by Indonesian people from generation to generation. Traditional medicine represents an essential potential in the development of traditional medicines derived from plants. The use of traditional medicine is preferred because of the assumption that traditional medicine’s side effects are relatively slighter when used correctly. Traditional medicine has a complementary or synergistic effect because traditional medicinal herbs generally consist of several types of medicinal plants that support each other to achieve the effectiveness of treatment.1

Hypertension and diabetes mellitus are two diseases that are directly related and require proper and thorough management. Hypertension does not merely cause heart attack, heart failure, and stroke, but in many cases, it often causes diabetes mellitus. To avoid being exposed to diabetes, hypertension sufferers were asked to maintain his blood pressure by keeping up body weight, levels of glucose, triglycerides, HDL, and LDL. The number of sufferer’s diabetes in Indonesia is increasing every year as increasingly sufferers hypertension and heart disease.2 Several studies have found an association between increased high blood pressure in patients with diabetes mellitus (DM). People who suffer from DM, primarily type 2 have a risk of 2 to 4 times more susceptible to death due to cardiovascular abnormalities than those who do not suffer from DM.

Conversely, the incidence of hypertension occurs two times more susceptible in patients with DM compared to DM in the same age group. In most diabetic populations, 80% of diabetic patient’s complication with hypertension, and vice versa, 70% of hypertensive patients may experience impaired glucose tolerance or develop type 2 diabetes. Targets for reducing blood pressure in hypertensive patients with diabetes are < 130 for systolic blood pressure and < 90 for diastolic blood pressure. Diabetes patients who are accompanied by hypertension will further increase the risk of coronary heart disease, stroke, nephropathy, and retinopathy. DM accompanied by hypertension increases by 75% morbidity and mortality in people who have previous risk factors.3,4 DM causes metabolic changes, including hyperglycemia, excessive expenditure of free fatty acids, and insulin resistance, which cause abnormalities of endothelial cell function because of a decrease in nitric oxide (NO). Impaired glucose tolerance in DM patients causes an increase in free fatty acids which causes endothelial damage and the incidence of hypertension.4

Management of hypertension can be done by non-pharmacologically and pharmacologically. Nonpharmacological therapy includes a low salt diet, regular exercise, limiting fat, sugar, and alcohol intake. Pharmacological therapy is done by giving antihypertensive drugs. The inappropriate selection of antihypertensive drugs in diabetic patients can worsen glycemic control. Some antihypertensive drugs can affect blood glucose level parameters, insulin sensitivity, and HbA1c.
Beta-blockers and thiazide diuretics can increase blood glucose levels and cause hyperglycemic coma at high doses. ACE inhibitors (ACEI) and Angiotensin Receptor Blocker (ARB) in hypertensive patients with diabetes complications possess a potential effect in lowering blood pressure, and is beneficial in preventing microalbuminuria, worsening kidney function, and other microvascular complications. Studies show that ACE inhibitors work by inhibiting the effects of angiotensin II, which are as vasoconstrictors. ACE inhibitors obtain the drug of choice in hypertension with diabetes because, in addition to the treatment of hypertension, it equally affects renal hemodynamics which can reduce glomerular hydraulic pressure. ACE inhibitors can lower glomerular hypertension and proteinuria by modifying pressure capillaries and glomerular permeability. ACE inhibitors beneficial for reducing protein excretion urine in diabetic or non-diabetic kidney disease. In patients with diabetes and hypertension with albuminuria, ACE inhibitors represent the first choice.  

The use of traditional medicines has been thoroughly developed and is counted as an integral component in essential health services. Traditional medicine, as part of the treatment of hypertension, has increased in the last decade. The contributing factor is mainly the price of traditional medicines, which are considered cheaper with fewer side effects. Some medicinal plants in Indonesia possess antihypertensive activities, including Persea Americana, Phyllanthus niruri, Morinda citrifolia, Phyllanthus niruri, Centella asiatica, Zingiber officinale, Alyxia reinwardtii, and Hibiscus rosasinensis.  

Extract preparation and standardization

A total of 500 grams of herbal medicine powder was extracted by maceration using 70% ethanol and stirred for 3 hours. The maceration process is carried out for 24 hours. The filtrate is carefully separated from the residue using filter paper. The residue is macerated again, and the process is repeated until a clear filtrate is obtained. The filtrate contains a combination of the solvent and the compound. Standardization of non-specific parameters includes checking ash content, ash content, which is insoluble in acid, water content, levels of extracts that are soluble in water, compounds that are soluble in ethyl alcohol, and coliform contamination and mold/yeast rates utilizing the ALT method. Standardization of specific parameters involves organoleptic, extract chemical content, and identification of the infrared spectrum.

**In-vitro ACE inhibitor activity assay**

Measurement of in vitro ACE inhibitor activity is carried out using a spectrophotometer under aerobic conditions. A total of 50 μL of the sample solution was added into a test tube, 50 μL of HHL substrate was added and pre-incubated at 37°C for 10 minutes. After that, 100 μL of ACE enzyme solution was added to the test tube and homogenized with a vortex. The mixture was incubated at 37°C for 90 minutes. To the solution, the mixture was included 250 μL HCl 1 N to prevent the reaction. The formed hippuric acid was extracted using 1.5 mL ethyl acetate and centrifuged for 10 minutes. The ethyl acetate layer is taken and evaporated at 100°C for 5 minutes. The enzyme solution is dissolved with 3 μL aqueous. The absorption of the solution is measured using a spectrophotometer at a wavelength of 246 nm. A similar procedure was carried out with Captopril as a standard. The absorbance of the sample is measured at a wavelength of 575 nm. The results determining ACE inhibitor activity compared with a Captopril.

**In-vitro antioxidant activity assay**

A total of 500 μL extracts at concentrations of 125, 250, 500, 750, and 1000 ppm added with 500 μL DPPH 0.125 mM. The mixture is then vortexed and incubated for 30 minutes. The absorption of test solutions and blanks was measured at a wavelength of 517 nm. The results of determining antioxidants compared with a vitamin C.

**In vitro alpha-glucosidase inhibitor activity**

A total of 10 μL of extract solution each at concentrations of 125, 250, 500, 750, and 1000 ppm was added 50 μL of phosphate buffer solution pH 7, and 25 μL p-nitrophenyl-α-D-glucopyranoside 10 mM. Next to the solution was added 25 μL of 0.04 U/mL enzyme solution and incubated at the time and temperature optimum. Then 100 μL NaCO3 200 mM was added. The absorbance was measured with a microplate reader at a wavelength of 410 nm. Percent inhibition is calculated using the formula:

\[
\%\text{Inhibition} = \frac{A_0 - A_1}{A_0} \times 100
\]

Where \(A_0\) is the absorbance of the control, \(A_1\) is the absorbance of the sample. \(IC_{50}\) is calculated using linear regression by plotting the concentration of percent inhibition. The results of determining alpha-glucosidase inhibitor activity compared with an Acarbose.

**Brine shrimp lethality bioassay**

A total of 50 mg of extract was added to 5 ml of dimethyl sulfoxide (DMSO) until a 10 mg/ml stock solution was obtained. Concentration series of test samples were made at concentrations of 5, 10, 25, 50, and 100 ppm by taking a specific volume of stock solution using a micropipette and place it in flacons. *A. salina* larvae eggs are hatched in a dark room in seawater media and aerated using aerators. The eggs will hatch about 24 hours into larvae. 48-hour-old larvae can be used and 100 ppm added with 500 μL DPPH 0.125 mM. The mixture is then vortexed and incubated for 30 minutes. The absorption of test solutions and blanks was measured at a wavelength of 517 nm. The results of determining antioxidants compared with a vitamin C.

### MATERIAL AND METHODS

Herbal medicine (Jamu) were obtained from herbal Market in Jakarta, Ethanol (Brachatem), Aquadest (Brachatem), Aquademineral (Brachatem), Ethyl Acetate (Merck), DMSO (Merck), Hippuric Acid (Sigma), Captopril (Sigma), HHL (Sigma), ACE from rabbit lung (Sigma), NaOH (Merck), Boric Acid (Sigma), DPPH (Merck), vitamin C (Sigma). A-Glucosidase (Sigma), p-nitrophenyl-α-D-glucopyranoside substrate (Sigma), dimethyl sulfoxide (DMSO) (Merck), Bovine Serum Albumin (BSA) (Sigma), Na2CO3 (Merck), buffer phosphate pH 7, acarbose (Sigma), Anemia salina Leach. (Brine eggs), NaCl (Merck).

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flacon filled with 1 mL of seawater, then vortexed for approximately to homogenize the sample. Ten tails of A. salina Leach. Age 48 hours a good (active move) randomly selected, put into flacons containing solvent-free samples using a dropper pipette then added seawater to 5 mL. One drop of yeast suspension Saccharomyces cerevisiae (3 mg/10 ml of seawater) is added to it as A. salina Leach food. The flacons are placed under the lighting for 24 hours, and the number of A. salina Leach larvae dead is counted.14,15 Percent mortality of larvae is calculated following the formula16:

\[ \% \text{ Death} = \frac{\text{number of A. salina larvae dead}}{\text{number of test larvae}} \times 100\% \]

RESULTS

Result of standardization both specific and nonspecific parameter presented in Table 1 and Table 2.

Data ACE inhibitory, α-glucosidase inhibitory, and antioxidant activity are presented in Table 3.

Results of Brine shrimp lethality bioassay presented in Table 4.

DISCUSSION

Based on the Regulation of Indonesia National Agency of Drug and Food Control No. 12/2014 concerning quality requirements for traditional medicines, it is seen that the extract of herbal medicine meets non-specific or specific parameters. Standardization of pharmaceutical products is carried out so that an extract meets the established chemical, biological, and pharmaceutical standards requirements. Standardization will guarantee that the extract or extract product has a constant parameter value. Standardization is the process of determining properties based on specific parameters to achieve the same degree of quality. Standardization of extracts is done by using two parameters, namely specific parameters, and non-specific parameters. Water content is one of the essential characteristics of the extract. Determination of the water content in the extract aims to provide a minimum limit or range of the amount of water content in the extract. Extracts that have high water content are natural to grow with mold and yeast to reduce the biological activity of the extract in the storage period. Water content depends on the plant’s drying time, the drier, the smaller the water content. Based on the Regulation of Indonesia National Agency of Drug and Food Control No. 12/2014, it is seen that the extracts of herbs meet the water content requirements of <10% (6.82±0.2). Ash content is a mixture of inorganic components or minerals found in an ingredient. Organic materials in the combustion process will burn but not the inorganic components. The ash content was determined by inserting the extract into the furnace at 450°C until ash was formed. Total ash content can also be used to know the internal and external mineral content from the initial process to the end of extracting. The extract ash content of 4.63 ± 0.08 and an acid insoluble ash content of 0.28 ± 0.02. This value is the following required in Herbal Pharmacopoeia, which is less than 7.8%. Small ash content values mark if the remaining material is small. The remaining material includes physiological ash derived from the plant tissue itself and non-physiological ash, a residue from foreign material attached to the plant’s surface, such as sand and soil. The smaller the value of ash content can be, the smaller the impurity in the resulting fraction.

Water-soluble and ethanol extract levels are indicators of active compounds that can be detected, both by water and ethanol solvents. The level of active compound in a plant is influenced by plant age, harvest time and climate, and place of growth. The test results showed that herbal medicine’s extracted content in aqueous solution was 28.38 ± 0.68%, and ethanol soluble was 57.62 ± 0.55%. These results are quite indicative that the active compound in the herbal extract is quickly scattered into water and ethanol. The low water content will prevent the growth of microorganisms and mold (fungus). Aspergillus flavus will produce aflatoxin, which is very toxic and can cause liver cancer. According to traditional medicine requirements, it is stated that the rate of yeast or mold does not exceed 104 CFU/gr. The extract must be

### Table 1: The specific parameters.

| Parameter | Value |
|-----------|-------|
| Organoleptic (colour and texture) | yellowish-brown and paste |
| Chemical content | Flavonoid, tannin, saponin, steroid |
| Identification of infrared spectrum | C=O, -OH; C=C; C-O’ C-H (aromatic); C-H (aliphatic) |

### Table 2: The nonspecific parameters.

| Parameter | Value |
|-----------|-------|
| Water content (v/w) | 6.82±0.2 |
| Ash content (w/w) | 4.63±0.08 |
| Ash insoluble acid content | 0.28±0.02 |
| Level of substances dissolved in alcohol (%) | 57.62±0.55 |
| Level of substances dissolved water (%) | 28.38±0.68 |
| Coliform microbial contamination (colony/g) | negative |
| Mold / Yeast numbers (colony/g) | negative |

### Table 3: ACE inhibitory, α-glucosidase inhibitory and antioxidant activity (IC₅₀).

| Sample | ACE inhibitor (µg/ml) | α-glucosidase (µg/ml) | antioxidant (µg/ml) |
|--------|-----------------------|-----------------------|---------------------|
| Extract | 292.15 | 36.13 | 24.43 |
| Standard | 0.64 | 0.268 | 5.13 |

### Table 4: Effect of ethanol extract jamu of various concentrations on mortality against Brine shrimp.

| Concentration (µg/ml) | Log concentration(µg/ml) | % mortality | Probit value | LC₅₀ (µg/ml) |
|-----------------------|--------------------------|-------------|-------------|--------------|
| 1000                  | 3                        | 80          | 5.8416      | 215.04       |
| 500                   | 2.6899                   | 66.7        | 5.4316      | 4.6415       |
| 100                   | 2                        | 36          | 4.1684      |              |
| 50                    | 1.6990                   | 20          | 3.3046      |              |
| 10                    | 1                        | 4.5         |             |              |
uncontaminated with pathogenic microbes, and the aflatoxin content is no more than 30 parts per million (ppm). Examination results of mold/yeast extract uncontaminated with mold/yeast. A natural medicinal product should not contain microorganism contamination, but this is difficult to avoid. The maximum limit of microorganism contamination required depends on the dosage form and is determined by the Total Plate and Yeast Fungi Numbers' determination. Natural medicinal products are unpermitted to contain pathogenic microorganisms like *Pseudomonas aeruginosa*, *Staphylococcus aureus*, and *Clostridia sp*. *Shigella sp.* and *Salmonella sp.* The permitted value or range is related to purity and contamination. The results showed that the extract was uncontaminated with microbial coliform.

Pathophysiology of Diabetes mellitus and hypertension occurs through several mechanisms such as improper activation of the angiotensin aldosterone (RAAS) system, secondary oxidative stress that causes over-production of reactive oxygen species (ROS), inflammation, vasoconstriction due to insulin, increased activation of the sympathetic nervous system and kidney abnormalities in secreting sodium.20 The key factors causing the coexistence of diabetes mellitus with hypertension are obesity and increased visceral adiposity. Mild chronic inflammation and oxidative stress in adipose tissue cause an increase in angiotensinogen and angiotensin II production, which results in the activation of RAAS tissue. Overexpression of angiotensinogen in adipose tissue causes an increase in blood pressure.22 Hypertension in diabetic patients can increase both microvascular and macrovascular complications. Proper management of hypertension is needed to minimize the occurrence of complications and inhibit disease progression. Barriers to the renin-angiotensin aldosterone system by ACE inhibitors or angiotensin receptor blockers can increase glucose metabolism by preventing the formation of angiotensin II or preventing the activation of Angiotensin II receptors.24 ACE inhibitors are antihypertensive drugs that have been used for the past several decades. Captopril, Lisinopril, Enalapril, and Ramipril are some examples of drugs that target ACE inhibitors. The use of these drugs in the long term can cause side effects like dizziness, coughing, and angioneurotic edema.25 Research to seek new drugs continues to be done, one of which represents the search for bioactive compounds from nature as a target. Some natural bioactive compounds that have been investigated to have ACE inhibitor activity are peptides, anthocyanins, flavonoids, and triterpenes.7

Table 3 shows that the extract has an ACE inhibitor, a-glucosidase inhibitory, and antioxidant activity with an IC50 value of 292.15 µg/ml, 36.13 µg/ml, and 24.43 µg/ml, respectively. Herbal extracts containing flavonoids, tannins, saponins, and steroids (Table 1). Secondary metabolites produced by plants are a group of natural compounds that are identified as potential ACE inhibitors. Some terpenoids and polyphenolic compounds, including flavonoids, hydrolyzable tannins, xanthones, procyanidin, caffeoylquinic acid derivatives, turned out to obtain an effective natural ACE inhibitor. Almost all studies show that plant extracts which are rich in phytochemicals are useful as ACE inhibitors.7

The development of insulin resistance, hypertension, and diabetes mellitus occurs due to oxidative stress. ROS can cause impaired endothelial function, tissue injury, reduction of bioavailable Nitrite oxide (NO), and impaired NO-mediated vasodilatation.

The herbs used in this study consisted of plants, namely *Morinda citrifolia* Fructus, *Phyllanthi niruri* Herb, *Centella asiatica* Herb, *Zingiberis officinale* rhizome, *Imperata* radix, and *Alyxiea* cortex. These six plants contribute to their antihypertensive and antidiabetic activities.

People widely use *Morinda citrifolia* L. as natural medicine. The content of flavonoids in this plant function as an antioxidant that can withstand the rate of absorption blood glucose from the digestive tract to the blood vessels so that able to withstand the rate of increase in blood glucose levels. Bioactive components such as flavonoids, triterpenoids, and saponins a significant amount has a hypoglycemic effect.20 *Morinda citrifolia* has antioxidant activity that can inhibit the rate of formation of Advanced Glycation End Products (AGES) and dicarbonyl compounds. The binding of AGEs to the AGEs (RAGE) receptor triggers reactive oxygen species (ROS) and NF-B activation of target cells, endothelium, mesangial cells, and macrophages an increase in permeability vascular.21 A study conducted by Mutia et al., 2017 shows that ethanol extract of *Morinda citrifolia* has ACE inhibitor activity with a percent inhibition against ACE of 66.64 + 2.32%.22 Scopeolin phenolic compounds and routine have an antihypertensive effect through the ACE inhibitor mechanism and antioxidant activity.23

*Phyllanthi niruri* L. has been empirically used in the management of diabetes and hypertension. Bioactive substances like glycosides, flavonoids, saponins, tannins, sterols, and carbohydrates are thought to be responsible for their actions. We already know certain flavonoids have hypoglycemic effects because they can regenerate pancreatic beta-cell regeneration. Studies in experimental animals have shown that sterol compounds can reduce blood sugar, whereas cardiac glycosides have cardio-protective and cardiotoxic activity. A study conducted by Bharati et al., 2017 shows that ethanol extract of leaves and fruit of *P. niruri* decreases blood pressure in diabetic hypertensive animals.24 *P. niruri* fruit juice comprises ACE inhibitory compounds, and daily consumption of the juice is useful for prevention and healing against hypertension.20

*Centella asiatica* (L.) is a plant that has long been used in traditional medicine for various indications like skin disorders, vascular disorders, microangiopathy, and inflammation. This plant also has vigorous antioxidant activity with the chemical content of phenolic compounds, namely flavonoids (quercetin, kaempferol, catechin, routine, apigenin, and naringin), triterpenoids (asiaticoside, madecassoside, asiatic acid, madecassic acid), glycosides, flavonoids, alkaloids, steroids, volatile and fatty oils. *C. asiatica* extract can reduce blood pressure in rats that are made hypertensive with L-NNAME. Quercetin may act in components sufficient for the antihypertensive effect of *C. asiatica* extract.25 Ethanol extract of freeze-dried and juice *C. asiatica* has ACE inhibitor activity with IC50 of 1 mg and 1.6 mg, respectively, advising possible usage as natural antihypertensive medicine.26 *C. asiatica* also has antiangiogenic activity in Alloxan induced diabetic rats. *C. asiatica* could restore the damage of the pancreatic β-cells and improve insulin synthesis, thus reducing plasma glucose level. The two glycosides, brahmoside and brahminoside, which are essential components of *C. asiatica*, have been confirmed to exert sedative and hypoglycemic effects in rats.26

Ginger (*Zingiber officinale* Roscoe) is a tropical plant used as a spice and has a therapeutic effect on various diseases like anticancer, anticoagulant, and antiemetic hypolipidemic and antioxidant. The main bioactive compounds contained in ginger obtain gingerol, with other gingerol analogs like shogoals, paradol, and zingerone. Gingerol compounds have antioxidant activity in vitro by protecting HDL-60 cells against oxidative stress. Ginger oil protects DNA from damage by H2O2. Ginger oil can serve an oxygen radical scavenger and can be used as an antioxidant.27-29 The administration of *Z. officinale* juice at a dose of 4 ml/kg once daily decreases blood glucose levels and increases plasma insulin in mice induced by streptozotocin. *Z. officinale* juice also lowers cholesterol, triglycerides, and blood pressure.20

*Imperata cylindrica* contains potassium, flavonoids, graminone B, and cylindrene, which is useful for treating various diseases, including hypertension. Infusion *I. cylindrica* in 15 healthy volunteers reduced systolic and diastolic blood pressure significantly lower (p < 0.01) than before drinking infusion. The presence of flavonoids on *I. cylindrica* exerts the effect of inhibiting the Angiotensin Converting Enzyme.
Potassium obtains anti-renin and inhibits aldosterone secretion. Potassium can decrease membrane potential, causing relaxation of vascular smooth muscle.\textsuperscript{19} Ethanol extract of \textit{I cylindrica} dose of 500 mg/kg has an antidiabetic effect in alloxan-induced mice.\textsuperscript{21} 

\textit{Alyxia reinwardtii} is a medicinal plant used to manage various diseases. The fruit can reduce fever, and the flower is effective in treating mental confusion and hallucinations, and the stem is used for heart failure and stomach discomfort due to gas distension. Chemical compounds in the cortex are iridoids, coumarin, and isolated lignans that have antioxidant activity.\textsuperscript{23}

Our previous study showed that herbal extracts containing \textit{Phaleria macrocarpa, Cynura procumbens, Imperata cylindrica, Centella asiatica, and Syzgium polyanthum} had ACE inhibitors activity with IC\textsubscript{50} values of 18.37 ppm.\textsuperscript{24} 

Toxic activity assay is one of the prerequisites for a plant to be developed as a drug, especially as an anticancer. Brine shrimp lethality test (BSLT) is a simple, inexpensive, non-aseptic, and high potentiality cytotoxicity test for bioactive compounds and used as a preliminary test to determine the activity of a substance or a compounds contained in an-aqueous extract or plant isolate.\textsuperscript{25} Acute toxicity can be measured as a concentration that can kill 50% of the tested animal population. LC\textsubscript{50} lethal concentrations were assessed at 95% confidence intervals using probit analysis. Toxicity assays using \textit{Artemisia Salina} can be used in filtering out various chemical compounds that have bioactivity.\textsuperscript{25,26} 

Table 4 shows that the jamu extract possesses cytotoxic activity with an LC\textsubscript{50} value of 215.04 μg/mL. Larvae mortality was seen to be directly proportional to the extract concentration, starting from the lowest (10 μg/ml) to the highest (100 μg/ml). An extract is said to be toxic if it has an LC\textsubscript{50} < 1000 μg/ml. If the BSLT test results indicate that a plant extract is toxic, it can be developed to isolate its bioactive compounds.\textsuperscript{26}

**CONCLUSION**

Jamu extract has ACE inhibitors, alpha-glucosidase inhibitors, and antioxidant activity with IC\textsubscript{50} values of 292.15 μg/mL, 36.13 μg/mL, and 24.43 μg/mL, respectively. Ethanol extract of jamu exerts a cytotoxic effect on larvae of shrimp \textit{Artemia salina} with an IC\textsubscript{50} value of 215.04 μg/mL.

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**CONFLICTS OF INTEREST**

No conflicts of interest to be declared by authors.

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**GRAPHICAL ABSTRACT**

Herbal medicine (*jamu*) were extracted with ethanol 70% by maceration

- Identification of the specific and non-specific parameters
- Nonspecific parameter: ash content, acid insoluble ash content, water content, water-soluble extract content, ethanol-soluble compound, and microbial contamination
- Specific parameters: organoleptic, extract chemical content and identification of infrared spectrum

- In vitro ACE Inhibitor activity assay: The examination was conducted using a spectrophotometer
- In vitro Antioxidant activity assay: Free radical scavenging activity. The absorbance was measured with using a spectrophotometer
- In vitro alpha-glucosidase Inhibitor activity: The absorbance was measured with using a microplate reader at a wavelength of 410 nm

- *Jamu* extract has ACE inhibitors, alpha-glucosidase inhibitors, and antioxidant activity with IC50 values of 202.15 µg/mL, 36.13 µg/mL, and 54.43 µg/mL, respectively.
- Ethanol extract of *jamu* exerts a cytotoxic effect on larvae of shrimp *Artemia salina* with an IC50 value of 215.04 µg/mL.

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