ATHEROPROTECTIVE EFFECT OF SOLANUM BETACEUM ON RAT EXPOSED TO CIGARETTE SMOKE

by Hanik Badriyah Hidayati
ATHEROPROTECTIVE EFFECT OF SOLANUM BETACEUM ON RAT EXPOSED TO CIGARETTE SMOKE

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Received: 19 December 2018, Revised and Accepted: 17 July 2019

ABSTRACT

Objective: The objective of this research was to determine the atheroprotective effect of Solanum betaceum ethanol extract toward carotid artery intima-media thickness (cIMT) and the level of malondialdehyde (MDA) on rats exposed to cigarette smoke.

Methods: Thirty adult Rattus norvegicus strain Wistar were divided into five groups and exposed to cigarette smoke, 3 pc cigarette/day and simultaneously administered with S. betaceum in Group K2, K3, and K4 (100 mg/kg b.w/day, 200 mg/kg b.w/day, and 400 mg/kg b.w/day, respectively). The duration of treatment for all groups was 28 days. Blood was withdrawn from the cardiac to determine the MDA level. Histological slide from carotid artery intima-media was collected to determine cIMT.

Results: S. betaceum ethanol extract administration could significantly prevent the development of atherosclerosis due to oxidative stress by decreasing the level of MDA (p<0.05) and reducing the degree of cIMT changes (p<0.05).

Conclusion: The present study found that S. betaceum ethanol extract could prevent the development of atherosclerosis due to smoke exposure through the reduction of the MDA level, i.e., the marker of oxidative stress, which is associated with the reduced of cIMT changes. However, further studies on other bioactivity of S. betaceum as an antioxidant are warranted.

Keywords: Solanum betaceum, Malondialdehyde, cIMT, Atherosclerosis, Smoke exposure.

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INTRODUCTION

Cardiovascular disease (CVD) is the primary cause of death in both developed and developing country. Coronary artery disease occurs in as much 42% of males and 25% of females, with an even higher percentage in people with other complications. It is estimated that until 2030, as much as 23.3 million people will die due to CVD [1,2]. One of the main causes of this disease is oxidative stress, which is 25% imbalance between oxidants and antioxidants inside the body [3]. Evidence shows that oxidative stress plays a significant role in the pathogenesis and development of CVD, among them being atherosclerosis [3,4].

Atherosclerosis is a systemic disease responsible for most incidences of CVD's and stroke [5]. Atherosclerosis is a complex phenomenon that is the main cause of coronary artery disease, which can be illustrated as an excess of fibro-fatty substance, or the accumulation of atheroma plaques, is an inflammatory response toward arterial wall damage [6,7]. Exposure to cigarette smoke is one of the main risk factors toward the early development of atherosclerosis and other CVD complications. Several main components of cigarette smoke are nicotine, tar, and carbon monoxide. These substances also have a negative influence on the body, both for the respiratory, and the cardiovascular system [8]. Cigarette smoke also contains free radicals. These free radicals can initiate the development of reactive oxygen species (ROS), which are reactive molecules that are constantly formed by the enzymatic reactions within the body [9,10]. When the level of ROS increases past that of the body's antioxidant defense system, oxidative stress occurs [9,11]. Oxidative stress creates substances that can cause cytotoxicity in the endothelial cells and smooth muscles of the blood vessel, which includes several variations of aldehyde, one of which is malondialdehyde (MDA) [11,12].

Oxidative stress is reported to have a significant role in the initiation and development of atherosclerosis by stimulating the production of inflammatory factors and cytokines [13]. Atherosclerosis is characterized by the weakening of the vessel walls, inflammatory response, the formation of foam cell plaques, and the formation of blood clots. The oxidative stress process causes endothelial dysfunction, and on the other hand, endothelial dysfunction increases the state of constant oxidative stress, thus causing the development of atherosclerosis [4].

This attention of the endothelial function plays an important role in the pathophysiology of atherosclerosis [14]. A published research on the topic has managed to prove the presence of foam cells in the blood vessels, from the tunica intima to the tunica media in the aorta of rats that have been exposed to cigarette smoke for 29–30 days [15]. The early phase of atherosclerosis is an alternative response of vascular cells that can be assessed by carotid artery intima-media thickness (cIMT) measurement, and it is an indicator of atherosclerotic plaques [14]. Measurement of the intima-media thickness (IMT) of the common carotid artery is an early morphological marker for the occurrence of atherosclerosis [5]. IMT is an indicator of atherosclerotic risk factors and the treatment given to reduce the influence of those risk factors [11,14,16].

The consumption of vegetable-based food is also linked to a low risk of CVD and hypertension. These last decades, several researchers have
focused on secondary metabolite substances of plants, for example, flavonoids, as a substance that may help prevent CVDs [17,18]. Tamarillo (Solanum betaceum), which is also known in Indonesian as Temeng belanda is a plant originating from the Solanaceae family. A majority of our citizens are not familiar with Tamarillo, often preferring eggplants as their daily vegetable of choice. Prior researches have shown that S. betaceum has a neuroprotective effect. The phytochemical component reported to be present in S. betaceum includes flavonoid, tannin, and terpenoid [19].

This research was done to analyze the atheroprotective effects of S. betaceum ethanol extract, measured through observed histopathological changes of the cIMT and arterial wall thickness parameters measured through the presence of MDA, as a possible early preventative measure against CVD due to cigarette smoke exposure.

METHODS

Experimental design

The present study was carried out in accordance with the guidelines of Ethical Clearance provided by the Animal Care and Use Committee, Faculty of Airlangga University with certificate number 0742-KE. This experimental study used Rattus norvegicus strain Wistar (3 months old, 180 g) in a 12:12 h light-dark cycle with room temperature 22°C. The animals were divided into groups and acclimatized to the laboratory conditions and maintained under 12 h light and dark cycles at room temperature 22°C. All the animals received a standard diet and water ad libitum. K1: Negative control group. K2: Positive control group (smoke exposure). K3: Treatment Group 1 (smoke exposure and ethanol extract of S. betaceum 100 mg/kg BW) [20]. K4: Treatment Group 2 (smoke exposure and ethanol extract of S. betaceum 200 mg/kg BW) [20], and K5: Treatment Group 3 (smoke exposure and ethanol extract of S. betaceum 400 mg/kg BW) [20].

The measurement of carotid artery intima medina thickness (cIMT)

Carotid artery tissue that has been isolated is fixed with formalin 10%. It is then made into a histologic slide and treated with Hematoxylin-Eosin (H&E) coloring. Observation is done with an Olympus microscope, with x100. The IMT is evaluated by measuring the tunica intima and media from the innermost endothelial boundary to the outermost tunica media using the cells software.

The measurement of malondialdehyde levels (MDA)

MDA level is obtained from the serum of R. norvegicus rats. Wistar strain, and is calculated with the thiobarbituric acid reactive substance method, and observed with a spectrophotometer at a wavelength of 529 nm, measured in mmol/ml [21]. Level of MDA was obtained from Zeptometrix, Buffalio, New York, USA, with Cat No. 8801192.

The preparation of S. betaceum ethanol extract

S. betaceum was purchased from farmland in Wonosobo, Central Java, Indonesia, and identified from Indonesian Institute of Science. S. betaceum was dried by fresh dryer. Dry powder was extracted by maceration using ethanol solvent for 3 x 24 h times at room temperature. The ethanol extract of S. betaceum was added to the treatment diet in the form of a suspension using 1% CMC with a dose of 2 ml/200 g b.w. S. betaceum extract was simultaneously administered to the treatment group before exposure to cigarette smoke during the experiment.

Cigarette smoke exposure

Exposure of cigarette smoke in experimental animals was given to groups K1, K2, K3, and K4 by 3 pc cigarettes per day for 28 days, at 6:00 p.m. The exposure system consisted of a glass box with two holes; several holes to insert cigarette smoke and other holes to remove cigarette smoke. A 50 cc syringe is pulled out to suck the cigarette placed on the iron pipe, so the cigarette smoke entered the syringe. The process of pumping 50 cc syringe is done repeatedly until the cigarette burned out. Locally available brand of cigarette, manufactured in Surabaya, East Java, Indonesia, was used in this study.

Statistical analysis

The statistical analysis was performed by one-way ANOVA and Kruskal-Wallis test. The statistical significance between groups was assessed by the least significant difference (LSD) and Mann-Whitney U-test with p<0.05. Data analysis was used by SPSS ver23.

RESULTS AND DISCUSSION

The average result of cIMT measurement in control and experimental animals are summarized in Table 1 and Fig. 1a. A significant increase (p<0.05) in cIMT was observed in K1 group (rats exposed to cigarette smoke) when compared to Group K0 (control group rats). S. betaceum-administered rats (Group K2, K3, and K4) showed a significant decrease in their cIMT (p<0.05) compared to Group K1 rats, which shows the atheroprotective effect of the substance against cigarette smoke exposure.

The level of MDA was observed by TBAIC method from the serum. The level of MDA in the different groups is shown in Table 2 and Fig. 1b. A significant increase (p<0.05) in MDA levels was observed in Group K1 (rats exposed to cigarette smoke) when compared to rats belonging to

Table 1: Mean, standard deviation, and ANOVA test of carotid artery intima-media thickness of rats control and treatment groups (mm)

| Variable | Group   | K0     | K1     | K2     | K3     | K4     | SI     | ANOVA  |
|----------|---------|--------|--------|--------|--------|--------|--------|--------|
| cIMT     |         | 32.76±5.06 | 46.77±4.14 | 38.18±5.17 | 40.09±2.43 | 41.05±10.58 | μm     | 0.024* |

*Significantly with p<0.05, **Mean±SD, ***Different superscript means significant between groups, cIMT: Carotid artery intima-media thickness, SD: Standard deviation, SI: System International of unit.
Table 2: Mean, median, minimum, maximum, and Kruskal-Wallis test of malondialdehyde of rats control and treatment groups

| Variable | Categori | Group | K0 | K1 | K2 | K3 | K4 | S1 | Kruskal-Wallis |
|----------|----------|-------|----|----|----|----|----|----|----------------|
| MDA      | Mean     | 1.83* | 5.86* | 2.46* | 3.41** | 3.55* | 0.002* |
|          | Median   | 1.80  | 6.86  | 2.33  | 2.96  | 1.98  | 1.98  |
|          | Minimum  | 1.52  | 2.78  | 2.99  | 5.31  | 6.45  |
|          | Maximum  | 2.21  | 8.06  | 3.13  | 6.45  |

*Significantly with p<0.05. **Different superscript means significant between groups. MDA: Malondialdehyde, S1: System International of Unit

Fig. 2: Histochromy of carotid artery intima-media thickness

Group K0. Rats belonging to S. beteacum – administered groups (i.e., K2, K3, and K4), exhibited a decrease in their MDA levels when compared to Group K1. This shows the antioxidiant effect of S. beteacum against cigarette smoke exposure.

The histochemistry of the carotid artery sample was used to evaluate cIMT. The cIMT of the common carotid artery is an early morphological marker for the occurrence of atherosclerosis. The research shows that exposure to cigarette smoke alters the cIMT significantly (Fig. 2b) when compared to the normal group (Fig. 2a). The administration of S. beteacum ethanol extract could prevent smoke-derived cIMT alteration (Fig. 2c), which is significantly compared with positive control group (Fig. 2b) not significantly different when compared to the normal group (Fig. 2a).

The present study disclosed that administration of S. beteacum ethanol extract on rats exposed to cigarette smoke could prevent the development of atherosclerosis by reducing the level of MDA and degree of cIMT changes in the samples. Phytochemistry screening analysis in previous studies [18] found that the ethanol extract of S. beteacum contains flavonoid, tanin, gallic tannin, and terpenoid which suggested an important role as antioxidant in the present study.

Pathogenesis of atherosclerosis (based on oxidative stress study) due to smoke exposure

The previous study has indicated that active and passive cigarette smoking is associated with dysfunction of endothelial cells [22]. Cigarette smoke contains so many free radicals that these can initiate the generation of ROS, which are reactive molecules that are constantly formed by enzymatic reactions within the body [9,10]. ROS compounds are produced by several oxidase enzymes, including nicotinamide adenine dinucleotide phosphate oxidase, xanthine oxidase, uncoupled endothelial nitric oxide synthase (eNOS), cyclooxygenase (COX), glucose oxidation, hypoglycaemia, and mitochondrial electron transport [23]. A variety of ROS and reactive nitrogen species (RNS) has been found to be involved in the disease pathology mechanism including radical superoxides (O2•−), radical hydroxyl (OH•), singlet oxygen (O2), hydrogen peroxide (H2O2), radical hydroxyl nitrite (ClO•), benzyl alcohol, and peroxyl nitrite (ONOO•), nitrite oxide (NO), and radical nitrogen dioxide (NOO•) [4,24-26].

This imbalance between oxidants and antioxidants inside the human body has the potential to cause damage through a process known as oxidative stress. It is a cellular, organic, or organism-wide metabolic condition that is characterized by an oxidative excess. The means through which ROS disturbs cellular function cannot be wholly explained as yet; however, one of the most important mechanisms is the occurrence of lipid peroxidation, which causes cell death [24,27]. One of the products of lipid membrane degradation is MDA, which is produced by the process of lipid peroxidation. This marker has often been identified in patients with atherosclerosis, certain types of tumors, neurodegenerative diseases, and lung illnesses, especially those due to inflammation [24].

Some existing evidence points to the fact that oxidative stress plays a significant role in the pathogenesis and development of CVDs, which includes atherosclerosis [4]. Atherosclerosis is an inflammation of the blood vessel due to the accumulation of atheromatous plaques within the walls of the artery causes the dysfunction of the vessel wall’s endothelial cells [18]. Oxidative stress causes endothelial dysfunction and endothelial dysfunction causes constant oxidative stress. This forms a vicious cycle that eventually leads to the formation of atherosclerosis.

Oxidative stress is related to a lack of tetrahydropterin and a decrease in eNOS activity, which may lead to endothelial dysfunction in atherosclerotic patients by causing a decrease in oxidative stress pathways within the body [4].

In patients with high cardiovascular risk level, the decrease in endothelial NO bioavailability can cause the following conditions: (a) reduced eNOS expression, (b) a disruption of cellular signals which causes an eNOS substrate or cofactor deficiency and the reduction of eNOS activation; (c) decrease in the endothelial cell’s capacity to synthesize and/or release NO; and (d) decrease of NO synthesis by ROS. All these abnormalities may lead to endothelial dysfunction, which is commonly seen as an initial step of the pathogenesis of atherosclerosis [4].

The early phase of atherosclerosis that can be assessed by carotid IMT (cIMT) measurements in later stages are mostly consisting of carotid plaques. Increasing of cIMT is typically seen in a common carotid artery due to its flow dynamics, whereas plaque is particularly seen in the internal carotid artery or carotid bulbs. As found in the present study, cIMT elevation is most often related vascular disorder due to smoke exposure [14,28].

Flavonoid: its role as atheroprotective substance

The process of atherosclerosis has long been seen as a systemic chronic inflammatory disease; mainly causing morbidity and mortality through...
its consequent cardiovascular illnesses [17]. A high intake of flavonoids from fruits and vegetables has a known correlation with a lower risk of cardiovascular illnesses. The mechanism to explain the related process is still unclear, but current evidence clearly shows that flavonoid can help reduce cardiovascular risk factors [18].

The flavonoid, included within the phenolic compound group, is often found in plant tissues and can act as an antioxidant. Flavonoid’s antioxidant activity stems from its ability to donate hydrogen atoms or form a complex to chelate metal. Multiple research has shown that flavonoid has anti-inflammatory, anti-microbial, anti-cancer, anti-diarrhea, and anti-viral properties [3,24,29,30]. Several studies have proven that flavonoid is positively linked with a significant reduction of coronary heart disease risk. The vascular endothelial cells play a significant role in safeguarding the health of the heart by producing nitric oxide, a substance that relaxes arteries (causing vasodilation). The production of endothelial nitric oxide can inhibit platelet adhesion and aggregation, which is one of the primary factors in the formation of a blood clot. A variety of clinical studies have researched the potential of high flavonoid intake in potentially reducing the number of platelet aggregation sizes; these trials have reported diet results. Food-based flavonoids were potentially reported to be involved in cardiovascular disease prevention, mainly by reducing oxidative stress and improving NO bioavailability. Flavonoid is to modulate genes that are related to metabolism, stress-defense, enzyme metabolism, detoxification, and protein transporters [18].

A majority of flavonoids work as an effective free-radical scavenger. However, this effect may not always be beneficial because flavonoids turn into radical flavonoids after they complete their scavenging functions. However, some radical flavonoids with high-stability will not easily react and will instead act as an antioxidant. The position and total number of hydroxyl chains, its configuration, substitution, and other flavonoid structure is the flavonoid chemical structure contribute to it can ability to chelate metallic ion, which in turns helps them scavenge and inhibit free radicals [3,21,23]. The hydroxyl chain configuration in the B-ring significantly muffles (scavenges) and chelates metallic ion from ROS and RNS due to their function as hydrogen and electron donor to hydroxyl, peroxyl, and radical peroxynitrite chains, which produces relatively stable radical flavonoids or radical aryl (Fl-O) [24,29].

Aside from that, flavonoid also has an antioxidant effect that can improve the length of a cell’s life, as well as induce apoptosis and inhibit cell proliferation, which is why it can also act as an anticancerogenic. In line with this information, it is reported that flavonoids such as epigallocatechin-gallate and resveratrol can inhibit transcription factors such as Nrf2 and AP-1 through an interaction between upstream signaling pathways (ERK phosphorylation, MAPK phosphorylation, and PI3K/Akt phosphorylation) and/or by reducing pro-inflammatory mediators (tumor necrosis factor-a, interleukin, and prostaglandin E2) and the activity of pro-inflammatory enzymes (COX-2, INOS) [31].

The modulation of cell-signaling pathways by flavonoids can help prevent cardiovascular diseases through the following means: (a) reduced inflammation, (b) reduction of the expression of vascular-cell adhesion molecules, (c) increased endothelial nitric oxide synthase (eNOS) activity and (d) reduced platelet aggregation. Nitric oxide is required to maintain vasodilation. Nitric oxide donors are often related to a heightened risk of CVDs. (d) reduced platelet aggregation. Thromboxane synthesis is one of the first steps in the formation of blood clots. These clots may block the coronary arteries or the brain and cause either a myocardial infarction or a stroke. Dealing with platelet aggregation is an important strategy in the primary and secondary prevention of CVDs [18].

Based on the evidence from the current results, our study suggested that the health-promoting capabilities effects of S. betacorum in those animal model of rats exposed to cigarette smoke could be attributed to the anti-inflammatory effect. As preventive from atherosclerosis, we suggest that S. betacorum can direct intercept free radical before any significant oxidation can occur which was located by decreasing level of MDA. As an antioxidant, S. betacorum can retard or slow down the oxidative processes leading to decrease MDA, i.e., end product of lipid peroxidation. Thus, reduce the incidence and prevent atherosclerosis through inhibit in carotid artery IMT changes induced by cigarette smoke exposure.

CONCLUSION
The present study findings showed that exposure to cigarette smoke could increase the level of MDA, i.e., marker of oxidative stress and degree of carotid artery IMT (cIMT) changes in the experimental animal. Ethanol extract of S. betacorum administration exhibited a powerful antioxidant potential toward lowering the level of MDA and thus reduce the development of atherosclerosis which shown by decreasing the degree of cIMT changes induced by cigarette smoke exposure. However, further studies on other bioactivity of S. betacorum are warranted.

ACKNOWLEDGMENT
We would like to thank Airlangga University for all the support they have given toward the completion of this research.

CONFLICTS OF INTEREST
All authors have none to declare.

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