Regulatory T cells and anti-inflammatory cytokine profile of mice fed a high-fat diet after single-bulb garlic (*Allium sativum* L.) oil treatment

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**Abstract**

**Purpose:** To analyze the effects of a single-bulb garlic (*Allium sativum*) oil (SGO) on the activation of regulatory T cells and anti-inflammatory cytokines IL-10 and TGF-β in mice fed a high-fat diet (HFD).

**Methods:** The study was conducted with 24 BALB/c male mice divided into six groups consisting of four mice each, i.e., control group (non-HFD, no treatment); HFD group without treatment, HFD administered with simvastatin, and three HFD groups administered SGO doses of 12.5, 25.0 and 50.0 mg/kg, respectively, and continuously treated (with SGO) for 45 days. The relative number of regulatory T cells, IL-10 and TGF-β were measured using flow cytometry.

**Results:** HFD decreased the expression of regulatory T cells, and the production of IL-10 and TGF-β compared to the control group (p < 0.05). SGO (50 mg/kg) significantly enhanced the activation of regulatory T cells and production of TGF-β in mice fed high-fat diet compared to simvastatin group (p < 0.05). The dose of 25 mg/kg SGO significantly increased the level of IL-10 in mice fed a HFD (p < 0.05).

**Conclusion:** These results suggest that SGO inhibits inflammatory processes in mice fed a HFD, by enhancing regulatory T cells and the anti-inflammatory cytokine IL-10 and TGF-β. Thus, SGO a promising food supplements and/or therapeutic agent for management of inflammation disorders caused by a HFD.

**Keywords:** Anti-inflammatory agent, *Allium sativum*, High-fat diet, Interleukin-10, Regulatory T cells, Transforming Growth Factor-β

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**INTRODUCTION**

Atherosclerosis is one of the metabolic disease characterized by blocked arteries [1]. In 2030, the prevalence of atherosclerosis disease is expected to increase to 23.3 million people. Atherosclerosis has a mortality rate of almost 50% worldwide and is one of the largest cause of death in the developed country such as Indonesia [2]. It involves several cell interactions such as endothelial cells, macrophages, smooth muscle cells and lymphocytes. The accumulation of LDL (low density lipoprotein) is the main factor that causes atherosclerosis [3].
The accumulation of LDL, especially oxidatively modified LDL (Ox-LDL) in artery walls, triggers innate immune responses through the recruitment and activation of macrophages. This process then triggers the adaptive immune response through the activation of pro-inflammatory T cells (Th1 subset) [4,5]. The accumulation of inflammatory cells in arteries causes the release of chemokines, interleukin, and protease such as interferon gamma (IFN-γ), tumor necrosis factor alpha (TNF-α) and ligand CD40 membranes. The release of these molecules lead to increased immune response and atherosclerotic lesion progression [5]. For this reason, the role of regulatory T cells is important for modulating the pro-inflammatory process into a pro anti-inflammatory process [3].

A medicinal plant that has often been used as an alternative treatment for reducing the risk of coronary heart disease is garlic (Allium sativum L.) [6]. The cardiovascular properties of garlic are attributed to organosulfur compounds. Garlic contains 33 sulfur compounds (allii, allicin, ajoene, allypropyl disulfide, and others), several enzymes (allinase, peroxidase and others), amino acids and minerals. Allicin is only produced when the garlic bulb is crushed or cut, activating the allinase enzyme, which metabolizes alliin to allicin [7]. Allicin has an anti-inflammatory effect [8]. Allicin reduces the immune response due to oxLDL in atherosclerosis plaque formation. Free radicals can be suppressed in the presence of antioxidants. Administration of allicin has been reported to enhance the expression of SOD (superoxide dismutase) enzyme [9].

The single bulb garlic variety is usually used as traditional medicinal plant in Indonesia. This study aimed to analyze the effect of single bulb/clove garlic oil extract (SGO) on the activation of regulatory T cells and anti-inflammatory cytokines IL-10 and TGF-β in mice fed a high-fat diet. Regulatory T cells and anti-inflammatory cytokines can be used as indicators to examine immune response related to atherosclerosis disease, in mice fed a high-fat diet.

**EXPERIMENTAL**

**Plant material**

Fresh bulbs of *Allium sativum* were collected and identified at UPT Materia Medica Batu, Malang, Indonesia on August 2017. A voucher specimen (no. TO-T0975) was prepared and deposited in the herbarium of the UPT Materia Medica Batu, Malang, Indonesia.

*Allium sativum* extract preparation

Preparation of single bulb garlic extract used a Soxhlet method with N-hexane as the solvent about 0.9 kg of single bulb garlic powder (dried simpisia) was dissolved in 4.5 L of hexane. The final mass of the garlic oil extract was 6 mg. The extraction was done at UPT. Materia Medica Batu, Malang, Indonesia.

**Animal studies**

Twenty-four normal Balb/c male mice aged 12 weeks old (38 ± 5 g) were obtained from Gadjah Mada University, Yogyakarta, Indonesia. The mice were maintained in a pathogen-free facility. The experimental protocol was approved by ethical clearance from the Research Ethics Committee, Medicinal Faculty, Brawijaya University (no. 880-KEP-UB). All animal experiments were performed according to the Principles of Laboratory Animal Care (NIH publication no. 85-23) [10].

Twenty-four Balb/c male mice were randomly divided into six groups (four animals in each): N—control group (normal diet); HFD—high-fat diet group; Simvas—HFD group receiving the Simvastatin 2.6 mg/kg BW; A1—HFD group receiving 12.5 mg/kg BW of SGO; A2—HFD group receiving 25.0 mg/kg BW of SGO; A3—HFD group receiving 50.0 mg/kg BW of SGO.

**Atherosclerosis mice model and SGO extract treatment**

All mice were fed with standard diet for 1 week for acclimatization before study. For the HFD group, the mice were orally received HFD feed once a day for 45 days. HFD feed consisted of a Hi-Grow Medicated 551 feed (30 %), duck yolk (10 %), coconut oil (30 %), wheat flour (5 %), corn (24.9 %), and cholic acid (0.1 %). The control group was fed the standard diet for 45 days. The various doses of single bulb garlic oil extract (SGO) were administered orally for four weeks.

**Lymphocyte isolation**

The mice were sacrificed and sectioned on week 12; spleens were then isolated and separated by gentle pipetting. The spleen suspension in propylene was added to Phosphate-buffered saline (PBS) up to 10 mL and then centrifuged at 2500 rpm, 4 °C for 5 min. The supernatant was removed and the pellet was resuspended in 1 mL of sterile PBS to produce the lymphocyte suspension used for further analysis.
**Lymphocyte count**

Lymphocyte count was calculated using a haemocytometer. A total of 10 μL of the lymphocyte suspension and 90 μL of Evans Blue were combined and homogenized in a microtube. The number of live lymphocytes was calculated using the formula:

\[
\Sigma \text{Lymphocyte} = \Sigma \text{lymphocyte count} \times 4 \times \text{dilution factor} \times 10^4 \text{cell/mL}
\]

**Antibody staining and flowcytometry analysis**

The lymphocyte suspension was co-incubated with FITC-conjugated rat anti-mouse CD4 and phycoerythrin (PE) anti-mouse CD-25 monoclonal antibodies for 15 min. Antibodies used for intracellular staining were PE/Cy5 anti-mouse Foxp3, PE/Cy5 anti-mouse TNF-α and PE/Cy5 anti-mouse IL-10. Antibodies were purchased from BioLegend, Inc (San Diego, CA). Next, 50 μL cytofix-cytospersm was added to the pellet and incubated for 20 min at 4 °C. Then, 500 μL washperm was added and centrifuged at 2500 rpm at 4 °C, for 5 min. The pellet was resuspended using 50 μL of antibodies in sterile PBS. Next, the pellet was re-suspended in 500 μL PBS and assessed via a BD FACS Calibur™ flowcytometer (BD Biosciences, San Jose, CA, USA). The data were then processed using the BD Cell Quest Pro™ software.

**Statistical analysis**

Data are presented as mean ± standard deviation (SD) and were analyzed using SPSS 16.0 for Windows. A one-way ANOVA was used to assess the statistical difference between the treatments. \( P < 0.05 \) was defined as being statistically significant. Significant treatment effects was further analyzed for between-treatment differences with the Tukey HSD Test.

**RESULTS**

**Effect of single bulb garlic oil (SGO) extract on level of regulatory T cells (CD4+CD25+Foxp3+)**

The effect of single bulb garlic oil (SGO) extract in mice fed a HFD could be seen clearly in the relative number of CD4+CD25+Foxp3+ T cells (Figure 1). A high-fat diet suppressed the activation of regulatory T cells in mice. The relative number of CD4+CD25+Foxp3+ in the normal group was significantly decreased (\( P < 0.05 \)) after a HFD treatment for 45 days (19.62 vs 7.59 %).

As seen in Figure 1, single bulb garlic oil (SGO) extract was able to promote the activation of regulatory T cells in mice fed a high-fat diet. The relative number of CD4+CD25+Foxp3+ was increased significantly (\( P < 0.05 \)) after administering SGO at a dose of 50.0 mg/kg BW (A3) (34.90 %) compared to other doses. In A2 treatment, the relative number of CD4+CD25+Foxp3+ also increased by 21.1 % but not as high as in A3 treatment. The dose of 12.5 mg/kg SGO did not restore the relative number of regulatory T cells to the normal level (17.29 %).

Administration of simvastatin to HFD mice also increased the activation of Treg cells by 21.56 %, but not as high as A3 treatment (\( P < 0.05 \)). SGO at 50.0 mg/kg was also effective in enhancing regulatory T cells that express Foxp3 in mice fed a HFD. Regulatory T cells that express Foxp3 have anti-inflammatory effect and secrete anti-inflammatory cytokines IL-10 and TGF-β.

**Figure 1**: Effect of single bulb garlic (A. sativum) oil (SGO) extract on the activation of regulatory T cells in mice fed a high-fat diet (HFD). A) The relative number of Treg cells (CD4+CD25+Foxp3+) in mice fed a HFD after 45 days of SGO treatment. B) The percentage of CD4+CD25+Foxp3+ after administration of SGO extract in mice fed a HFD. Data are mean ± SD (n = 4). Note: N: normal mice (normal diet); HFD: high-fat diet mice; Simvas: high-fat diet mice receiving simvastatin dose of 2.6 mg/kg BW; A1: HFD mice receiving SGO 12.5 mg/kg BW; A2: HFD mice receiving SGO 25.0 mg/kg BW; A3: HFD mice receiving SGO 50.0 mg/kg BW

**Effect of single bulb garlic oil (SGO) extract on IL-10 production (CD4+CD25+IL-10+)**

IL-10 is a suppressive cytokine known as being effective in reducing inflammation in atherosclerosis disease. We observed that a
high-fat diet suppressed the production of IL-10. The relative number of CD4+CD25+IL-10- was significantly decreased (p < 0.05) in mice fed a HFD compared to normal mice (11.90 vs 23.39 %) (Figure 2). Flowcytometry analysis showed that administration of SGO significantly increased the production of IL-10 by Treg cells in all treatment doses of SGO (A1: 26.03 %; A2: 38.48 %; A3: 34.87 %). This results suggests that A2 group (SGO dose of 25.0mg/kg BW) had a higher level of IL-10 compared to other doses. As seen in Figure 2 B, A2 group was not significantly different in the relative level of IL-10 as compared to the simvastatin group (Simvas: 46.44 %).

The A3 group (50.0 mg/kg of SGO) appeared to have been administered the optimal dose, which increased the anti-inflammatory TGF-β compared to other doses of SGO and simvastatin group in atherosclerosis mice model. Increasing levels of TGF-β is crucial for the progression of atherosclerosis to result in inflammation, fibrosis, chemotaxis, proliferation, and apoptosis.

**Figure 2:** The effects of single bulb garlic (A. sativum) oil (SGO) extract on IL-10 production in mice fed a high-fat diet (HFD). A) The relative level of IL-10 (CD4+CD25+IL-10-) in mice fed a HFD after 45 days of SGO extract treatment. B) The percentage of CD4+CD25+IL-10- after administration of SGO extract in mice fed a HFD. Data are mean ± SD (n = 4). **Note:** N: normal mice (normal diet); HFD: high-fat diet mice; Simvas: high-fat diet mice receiving simvastatin at dose of 2.6 mg/kg; A1: HFD mice receiving SGO 12.5 mg/kg; A2: HFD mice receiving SGO 25.0 mg/kg; A3: HFD mice receiving SGO 50.0 mg/kg

**Effect of single bulb garlic oil (SGO) extract on TGF-β production (CD4+CD25+ TGF-β*)**

The administration of SGO to mice fed HFD showed significant increase in the production of TGF-β. TGF-β is important in controlling inflammation. Flow cytomtery analysis showed that the relative level of CD4+CD25+ TGF-β* decreased significantly (p < 0.05) in mice fed a high-fat diet and treatment with 50.0 mg/kgBW dose of SGO (A3: 20.10 % vs HFD: 15.25 %) reversed it (Figure 3). The relative level of TGF-β at dose 50.0 mg/kg of SGO was increased by 20.10 % compared to normal mice (19.99 %).

**DISCUSSION**

Regulatory T cells (CD4+CD25+FOXP3*) are the main factors involved in immune tolerance and induction [11]. Treg cells CD4+CD25+FOXP3* from the thymus are found in atherosclerotic lesions [12]. Treg cells have a suppressive effect and level of Foxp3 expression determines their suppressive role [13]. Treg cells-expressing Foxp3 secreting the anti-inflammatory cytokines IL-10 and TGF-β [14].

The function of Treg cells is closely related with the effect of suppressive cytokines such as TGF-β and IL10. Some research has reported that Treg cells secrete several cytokines such as TGF-β and IL10 that inhibit the activation of certain cells [15]. IL-10 is an anti-inflammatory cytokines that is effective in atherosclerosis.
Therapy with suppressive cytokines is important for decreasing the progression of inflammatory disease, especially atherosclerosis. The present study found that IL-10 cytokines by Treg cells are crucial factor for reducing the progression of atherosclerosis in mice fed a high-fat diet. The increasing levels of IL-10 due to administration of SGO is one therapy that may be applied to individuals with atherosclerosis disease. Many studies have revealed that TGF-β may inhibit the progression of atherosclerosis disease. One of the possible pathways that it may affect is the inhibition of proliferation, migration and apoptosis of smooth muscle cells and endothelial cells and also potentially inhibits the immune system. Activated Treg cells have a large effect on IL-10 and TGF-β secretion [3].

This study showed that the increase of Treg cells activation followed by the production of the anti-inflammatory cytokines IL-10 and TGF-β in mice fed a HFD and administered SGO indicated that there are active compounds in single bulb garlic that may activate Treg cells. Furthermore, the level of regulatory T cells activated by SGO was higher than that of simvastatin group. Simvastatin is a drug used commonly to treat hypercholesterolemia and coronary heart disease; this study used simvastatin as a positive control; simvastatin is known to inhibit 3-hydroxy-3-methyl-glutaryl-coenzyme A (HMG-CoA) reductase, which catalyzes cholesterol formation. Suppression of this enzyme (HMG-CoA reductase) have an effect on the ROS reduction and increasing of oxidised LDL resistant [20].

Single clove garlic (Allium sativum L.) has been used for traditional medicine, likely because of efficacy of organosulfur compounds present in it such as diallyl disulfide, S-allylcysteine and diallyl trisulfide, or allicin, which is enzymatically produced upon injury to the bulb [21, 22]. Recent studies have demonstrated a pharmacological effect of A. sativum, especially the active compound allicin. Some studies suggest that allicin and sulfur compounds in single clove garlic possess' antifungal, antibacterial, antiparasitic, antiviral, anticancer and cardiovascular protective effects. Some cardiovascular protective effects may be anti-hypertensive [27], antiatherosclerotic and antithrombotic [23]. Research by Wang and Ng [24] reported that garlic has protective effect for atherosclerosis disease. Abramovitz et al [25] have shown that allicin compounds in garlic clove reduce the serum cholesterol and LDL levels. Allicin significantly suppressed oxidized LDL and atherosclerosis progression [22, 23]. The active compound of garlic that has an anti-atherogenic role are mostly found in the oil fraction [23]. Currently, there is no research on the effect of garlic in the activation of immune cells. SGO reduce the migration of neutrophils by endothelial cells and have an effect on T lymphocyte function in extravascular inflammatory sites [23]. This study suggests that garlic oil extract may be a suitable alternative herbal treatment for the prevention of atherosclerosis.

CONCLUSION

These results suggest that SGO inhibits inflammatory processes in mice fed a HFD, by enhancing regulatory T cells and the anti-inflammatory cytokine IL-10 and TGF-β. Thus, SGO can be developed as a food supplement and/or therapeutic agent to ameliorate inflammation disorders caused by HFD.

DECLARATIONS

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Conflict of interest

No conflict of interest is associated with this study.

Contribution of authors

We declare that this work was done by Sri Rahayu Lestari and Muhaimin Rifa’i and all liabilities pertaining to claims relating to the contents of this article will be borne by the authors. Sri Rahayu Lestari performed the study design and drafting the manuscript and Muhaimin Rifa’i performed the final approval of the manuscript.
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