Extracellular-Signal Regulated Kinase Signalling Pathway Mediates the Increased Proliferation of EPCs Treated with Garlic (Allium sativum) Extract, Purple Sweet Potato (Ipomoea batatas) Extract, and Vitamin C

Yudi Her Oktaviono1,*, Alisia Yuana Putri1, Makhyan Jibril Al-Farabi1,2, Yesita Rizky Firmansyah3, Ferry Sandra4,5

ABSTRACT

The endothelial progenitor cell (EPCs) proliferation capability is reduced in the patient with stable coronary artery disease (SCAD). Garlic (Allium sativum), purple sweet potato (Ipomoea batatas), and vitamin C are proven antioxidant which potentially improve EPCs proliferation ability. **Objective:** To investigate the effect of garlic (Allium sativum), purple sweet potato (Ipomoea batatas), and vitamin C in EPCs proliferation from CAD patients and identify the involvement of Extracellular-Signal Regulated Kinase (ERK) Signalling Pathway.

**Material and Method:** Mononuclear cells were isolated from SCAD patients and cultivated with colony-forming unit (CFU)-Hill medium and divided into untreated (control), garlic extract (10 mcg/ml and 100 mcg/ml), purple sweet potato extract (1 mcg/ml and 25 μg/ml), and vitamin C (10 μg/ml and 250 μg/ml). EPCs proliferation was measured using the MTT Assay. **Results:** This research shows that EPCs proliferation was increased in the treatment with garlic extract at 10 mcg/ml and 100 mcg/ml dose (0.267 ± 0.003 and 0.391 ± 0.008; p < 0.05), purple sweet potato extract at 1 mcg/ml and 25 μg/ml dose (0.250 ± 0.005 and 0.356 ± 0.023; p < 0.001), and vitamin C at 10 μg/ml and 250 μg/ml dose (0.259 ± 0.016 and 0.306 ± 0.022; p < 0.001). Increased ERK expression was found in the treatment with garlic extract, purple sweet potato extract and vitamin C. **Conclusion:** Garlic extract, purple sweet potato extract, and vitamin C can increase EPC proliferation through the ERK signaling pathway.

**Key words:** Antioxidant, ERK, Endothelial Progenitor, Proliferation.

INTRODUCTION

Coronary artery disease is a major health problem that causes mortality and reduction of life quality worldwide.1 Endothelial progenitor cells (EPCs) from the patients with stable coronary artery disease (SCAD) had a progressive reduction of proliferation abilities, which worsen as the disease progressed.2 Impaired EPCs proliferation will reduce vascular damage repair.3 Lower EPCs proliferation in the patient with SCAD is also associated with a higher incidence of cardiovascular events, mortality, and morbidity.2

Multiple pathways were suggested to be responsible for EPCs impairment in SCAD patients. It is suggested that oxidative stress play significant roles in EPCs impairment through intracellular damage and balance disruption which will alter the control of apoptosis, proliferation, self-renewal, senescence, and differentiation of EPCs.4,5 Oxidative stress may disrupt the ERK signaling pathway, which is important in EPCs proliferation.6,7 The antioxidant is suggested to have a beneficial effect on impaired EPCs proliferation from patients with cardiovascular disease.7,8

Plants with antioxidant properties such as Chokeberry (Aronia melanocarpa),9 potato shoot (Solanum tuberosum) and Marigold (Calendula officinalis) has been proven to improve impaired EPCs proliferation.10 Similarly, vitamin C also able to prevent lowering EPCs proliferation caused by TNF-α.10 Garlic (Allium sativum), purple sweet potato (Ipomoea batatas) extract and vitamin C have potent antioxidant capabilities.8,9 Previous studies also have shown purple sweet potato extract and vitamin C on the EPCs from SCAD patients and identify its mechanism.4 Hence, this research aims to identify the effect of garlic extract, purple sweet potato extract and vitamin C treatment on the EPCs and identify the involvement of ERK phosphorylation in the SCAD patient.

MATERIAL AND METHODS

Garlic extract, purple sweet potato extract, and vitamin C preparation

Garlic and Purple sweet potato were obtained from UPT Materia Medica Batu, Indonesia and vitamin C powder was obtained from Sigma-Aldrich, USA. Purple sweet potato and garlic extract were produced with aqueous extraction method as described...
previously. The L-ascorbic acid dose referred to previous research that use dose 250 mcg/mL to improve adipocyte stem cell proliferation. Briefly, PSP chunks were mixed in water with a 1:1 ratio and blended. The mixture was filtered then boiled for 30 min and dried up using a rotary evaporator. PSP extract was diluted with the culture medium to achieve a concentration of 1 mcg/mL and 25 mcg/mL. Vitamin C powder was suspended in double-distilled water and diluted with culture medium to obtain a concentration of 10 mcg/mL and 250 mcg/mL.

Subject recruitment and sample collection
The blood sample was obtained from eight patients with SCAD in Dr. Soetomo General Hospital with inclusion criteria as follows: male, aged 40-59, stable angina, and coronary angiography showed >50% stenosis of left main coronary artery or >70% of other coronary arteries. Subjects with a history of percutaneous coronary intervention, coronary artery bypass grafting, acute myocardial infarct, diabetes, smoking, and anemia were excluded. The study protocol was approved by the Health Research Ethics Committee of Dr. Soetomo General Hospital, Surabaya (No.292/Panke.KKE/IV/2016). Each subject has signed informed consent before subject recruitment.

EPCs isolation and culture
Peripheral blood mononuclear cells (PBMCs) were isolated from the blood sample by Ficoll Histopaque 1077 (Sigma-Aldrich, USA). To isolate EPCs from PBMCs, a standard protocol was conducted as described previously. Briefly, 5x10⁵ cells/mL PBMCs were cultured in the fibronectin-coated 6-well plate with basal hematopoietic stem cell expansion medium (Sigma-Aldrich, USA) supplemented with 15% fetal bovine serum and 40 ng/mL vascular endothelial growth factor. The culture was maintained at 37°C with 5% CO₂ in a humidified atmosphere. Two days after, non-adherent cells were discarded, and fresh medium was added. Two weeks after, cultured cells were stained FITC-labeled anti-human CD34 antibody clone 581 (Biolgend, USA) and documented with an inverted immunofluorescence microscope. EPCs were confirmed from CD34 expression.

EPCs proliferation assay
MTT cell proliferation assay kit (Sigma-Aldrich, USA) was used to measure EPCs proliferation as described previously. Treated EPCs were added with MTT reagent and incubated in a 37°C incubator with 5% CO₂ for 4 hours. Proliferation was determined from the reduction of tetrazolium (MTT) into insoluble formazan product by viable EPCs and documented with an image analyzer.

Statistical analysis
Statistical analyses were carried out using IBM SPSS Statistics 25.0 (IBM Corp, USA). Data were compared using the ANOVA test and considered to be significantly different if p < 0.05 or p < 0.001.
Oktaviono, et al.: Extracellular-Signal Regulated Kinase Signalling Pathway Mediates the Increased Proliferation of EPCs Treated with Garlic (*Allium sativum*) Extract, Purple Sweet Potato (*Ipomoea batatas*) Extract, and Vitamin C

**Figure 1:** a) EPCs colony under 100x magnification in the inverted microscope; b) Immunofluorescence view of FITC-labelled CD34 expression which confirms EPCs; White bar represents 100 µm.

**Figure 2:** Garlic extract, purple sweet potato extract, and vitamin C improve EPCs proliferation in a dose-dependent manner. EPCs proliferation after treated with a) 10 mcg/mL and 100 mcg/mL garlic extract; b) 1 mcg/mL and 25 mcg/mL purple sweet potato extract; c) 10 mcg/mL and 250 mcg/mL Vitamin C for 48 h. EPCs proliferation was measured using MTT proliferation assay and statistically analyzed, as described in Materials and Methods. Sextuplicate was performed for each group. *: significant difference at \( p < 0.05 \); **: significant difference at \( p < 0.001 \).

**Figure 3:** Garlic extract, purple sweet potato extract, and vitamin C increase ERK expression with higher doses. ERK phosphorylation in the electrophoresis gel after treated with a) 10 mcg/mL garlic extract; b) 100 mcg/mL garlic extract; c) 1 mcg/mL purple sweet potato extract; d) 25 mcg/mL purple sweet potato extract; e) 10 mcg/mL vitamin C; and f) 250 mcg/mL vitamin C for 48 h.

Improve EPCs proliferation and migration.\(^{13}\) It is also suggested that garlic extract improves EPCs’ neovasculogenesis capability through modulation of the PI3/Akt pathway.\(^{14}\)

Purple sweet potato contains a high amount of anthocyanins, which was proven to improve impaired EPCs proliferation and migration *in vivo*.\(^{15}\) High-level of anthocyanin also has been proven to increase EPCs proliferation capability in a dose-dependent manner through the reduction of intracellular ROS.\(^{9}\) This suggested that the benefit of purple sweet potato extract on the EPCs proliferation might involve the ROS pathway. Interestingly, purple sweet potato showed an inhibitory effect on breast cancer, gastric cancer, bladder cancer cell, and colon adenocarcinoma proliferation.\(^{1,17-19}\) This suggests that purple sweet potato might both stimulate or inhibit cell proliferation depending on the type of cells.

Vitamin C treatment has been proven to improve the proliferation of adipocyte stem cells, cardiac progenitor cells, and intestinal stem cells.\(^{13,20,21}\) Similar to this research, vitamin C at the dose of 10 mcg/mL was shown to prevent the impairment of EPCs proliferation.
caused by TNF-α through P38 inhibition. Interestsingly, vitamin C at the dose of 100 mg/dl can reduce the proliferation of the EPCs and vasculogenesis. This suggests that dose-dependent effect of vitamin C to the EPCs proliferation.

In this research, the higher dose of garlic extract (100 mcg/mL), purple sweet potato extract (25 mcg/mL), and vitamin C (100 mcg/mL) were proven to increase phosphorylated ERK of the EPCs compared to the lower dose. While the exact mechanism of impaired proliferation in the EPCs in CHD patients remains unclear, it is suggested that oxidative stress may be involved in the EPCs impairment. Several antioxidants such as vitamin E, resveratrol, and L-arginine have proven to improve EPCs proliferation in-vivo and in vitro. Oxidative stress downregulates ERK signaling pathway in the embryonic stem cell. While ERK signal transduction pathway is responsible for promoting cell proliferation, nutrient uptake, and cell survival. Reduced ERK signaling pathway also has proven to reduce EPCs proliferation in patients with stable angina. Hence, It is speculated that the beneficial effect of the garlic extract, purple sweet extract, and vitamin C extract to improve EPCs proliferation may involve its antioxidant capability which increases ERK phosphorylation.

CONCLUSION
Treatment with garlic extract, purple sweet potato extract and vitamin C increase EPC proliferation which might involve ERK signaling pathway.

ACKNOWLEDGMENT
The authors are grateful to the staff of the Surabaya Regenerative and Stem Cell Centre who assist the laboratory works.

CONFLICTS OF INTEREST
The authors declare no conflict of interest.

REFERENCES
Oktaviono, et al.: Extracellular-Signal Regulated Kinase Signalling Pathway Mediates the Increased Proliferation of EPCs Treated with Garlic (*Allium sativum*) Extract, Purple Sweet Potato (*Ipomoea batatas*) Extract, and Vitamin C

**GRAPHICAL ABSTRACT**

**ABOUT AUTHORS**

Yudi Her Oktaviono work as an interventional cardiologist in the Department of Cardiology and Vascular Medicine Soetomo General Hospital and Faculty of Medicine, University of Airlangga. Previously, he led the Indonesian Cardiologist association in Surabaya and currently he is the vice president of the Indonesian Cardiologist association. His research interests include complex interventional cardiology, endothelial progenitor stem cells, usage of natural compound for cardiac disease and detection of early cardiac markers.

Alisia Yuana Putri work as a cardiologist in the Department of Cardiology and Vascular Medicine Soetomo General Hospital and Faculty of Medicine, University of Airlangga and Manyar Medical Center. Her research interests endothelial progenitor stem cells and cardiology risk in surgery.

Yesita Rizky Firmansyah Putri is a medical doctor which currently work as Trust Grade Doctor of Neurosciences at University Hospital Coventry and Warwickshire, UK. She had postgraduate degree in Biomedicine from Brawijaya University. Her research interests are Clinical Cardiology and Neurology.

Makhyan Jibril Al-Farabi is a cardiology resident in the Department of Cardiology and Vascular Medicine Soetomo General Hospital and Faculty of Medicine, University of Airlangga. He also have postgraduate degree from University College London, Quantics School of Business and Technology and Brawijaya University. His research interests includes stem cells, clinical cardiology, cardiometabolic syndrome, healthcare entrepreneurship and AI in cardiology.
Oktaviono, et al.: Extracellular-Signal Regulated Kinase Signalling Pathway Mediates the Increased Proliferation of EPCs Treated with Garlic (*Allium sativum*) Extract, Purple Sweet Potato (*Ipomoea batatas*) Extract, and Vitamin C

**Ferry Sandra** is a professor in the Department of Biochemistry and Molecular Biology Dental Faculty, Trisakti University Jakarta. He is also responsible to led the the BioCORE Laboratory. Previously, he was the director of Stem Cells and Cancer Institute (SCI). His research interests includes stem cells, cancer, inflammation and dentistry.

**Cite this article:** Oktaviono YH, Putri AY, Al-Farabi MJ, Firnansyah YR, Sandra F. Extracellular-Signal Regulated Kinase Signalling Pathway Mediates the Increased Proliferation of EPCs Treated with Garlic (*Allium sativum*) Extract, Purple Sweet Potato (*Ipomoea batatas*) Extract, and Vitamin C. Pharmacog J. 2020;12(3):