Inhibitory effects of scoparone through regulation of PI3K/Akt and MAPK on collagen-induced human platelets

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Abstract When blood vessels are damaged, a fast hemostatic response should occur to minimize blood loss and maintain normal circulation. Platelet activation and aggregation are essential in this process. However, excessive platelet aggregation or abnormal platelet aggregation may be the cause of cardiovascular diseases such as thrombosis, stroke, and atherosclerosis. Therefore, finding a substance capable of regulating platelet activation and suppressing agglutination reaction is important for the prevention and treatment of cardiovascular diseases. 6,7-Dimethoxy-2H-chromen-2-one (Scoparone), found primarily in the roots of Artemisia or Scopola plants, has been reported to have a pharmacological effect on immunosuppression and vasodilation, but studies of platelet aggregation and its mechanisms are still insufficient. This study confirmed the effect of scoparone on collagen-induced human platelet aggregation, TXA₂ production, and major regulation of intracellular granule secretion (ATP and serotonin release). In addition, the effect of scoparone on the phosphorylation of the phosphoproteins PI3K/Akt and mitogen-activated protein kinases (MAPK) involved in signal transduction in platelet aggregation was studied. As a result, scoparone significantly inhibited the phosphorylation of PI3K/Akt and MAPK, which significantly inhibited platelet aggregation through TXA₂ production and intracellular granule secretion (ATP and serotonin release). Therefore, we suggest that scoparone is an antiplatelet substance that regulates the phosphorylation of phosphoproteins such as PI3K/Akt and MAPK and is of value as a preventive and therapeutic agent for platelet-derived cardiovascular disease.

Keywords 6,7-Dimethoxy-2H-chromen-2-one · Granule secretion · Mitogen-activated protein kinases · PI3K/Akt · TXA₂

Introduction

Blood plays an absolute role in survival by providing oxygen and nutrients to each organ and tissue while also aiding in the removal of waste products. In order to do this normally, smooth circulation of blood must be achieved. Therefore, when blood vessels are damaged, a rapid hemostatic response should occur in order to minimize blood loss and maintain normal circulation. The response begins with the activation and aggregation of platelets. However, excessive or abnormal platelet aggregation may be a cause of cardiovascular diseases such as thrombosis, stroke, and atherosclerosis. Therefore, the search for a substance capable of regulating platelet activation and suppressing aggregation is important for the prevention and treatment of cardiovascular diseases [1,2]. When blood vessels are damaged, platelets are recruited to the site of injury and are activated by vascular activators (collagen, ADP, thrombin). At this time, activated phospholipase C in platelet membrane hydrolyzes phosphatidylinositol 4,5-bisphosphate into inositol 1,4,5-triphosphate (IP₃) and diacylglycerol, and the resulting IP₃ increases Ca²⁺ concentration in the cytoplasm by opening the Ca²⁺ channel of the dense tubular system in platelets [3]. During the activation of platelets, arachidonic acid hydrolyzed by cytosolic phospholipase A₂ (cPLA₂) is converted to TXA₂ due to the enzymatic action of both TXA₂ synthase and cyclooxygenase-1, and secreted out of platelets [4,5]. The released TXA₂ is known to function as a potent agonist that promotes platelet aggregation by binding to other platelet membrane...
receptors [6]. Furthermore, U46619 (9,11-dideoxy-9a,1a-methanoepoxy prostaglandin F2a), an analog of TXA₂, is known to induce platelet aggregation by phosphorylating myosin light chains and pleckstrin by increasing Ca²⁺ levels in the cytoplasm [7,8].

Mitogen-activated protein kinases (MAPK) are phosphorylase enzymes known to act on intracellular signaling, which are classified into c-Jun N-terminal kinase (JNK), extracellular signal-regulated kinase (ERK), and p38 MAPK, and their roles have been steadily studied [9]. JNK, ERK, and p38 MAPK have been reported to be detected in human platelets and have activity by phosphorylation by several agonists [10-12]. Phosphorylation of this signal transduction molecule is well known to play an important role in inducing platelet granule secretion [13,14]. In addition, MAPK phosphorylates ePLA₂ located in the cell membrane, thereby making it active, which is known to generate arachidonic acid from the phospholipids of the platelet membrane, convert it into TXA₂, and act on other platelets to trigger activation and aggregation of platelets [15,16]. In addition, the PI3K/Akt pathway has demonstrated a vital role in the regulation of platelet function, including the secretion of dense granules in platelets and platelet aggregation [17].

6,7-Dimethoxy-2H-chromen-2-one (Scoparone) found primarily in the roots of Scopolia or Artemisia plants, has been studied to have pharmacological properties such as immunosuppression or vasorelaxation. It has been reported that scoparone inhibits IL-1β-mediated inflammatory responses by modulating the PI3K/Akt/NF-κB pathway and thus may be a therapeutic material for joint diseases [18-20]. A previous study has shown that scoparone has the effect of inhibiting thrombus formation through the regulation of cyclic nucleotides in U46619-induced platelets [21]. However, the role of scoparone in platelet aggregation and the mechanism of scoparone on human platelets induced by collagen are currently unknown. In this study, we examined the effectiveness of scoparone on collagen-induced platelet aggregation and its important factors, TXA₂ production, platelet granule release, and the phosphorylated proteins of PI3K/Akt and MAPK. Through this, it is expected to contribute to the prevention of cardiovascular diseases caused by platelet aggregation by ingesting eggplants and wormwood, known as plants containing scoparone, as food.

Materials and Methods

Materials

Scoparone was received from Avention Corporation (Seoul, Korea) (Fig. 1). Collagen was provided by Chrono-Log Corporation (Havertown, PA, USA). TXB₂ enzyme immunoassay (EIA) kit, ATP assay kit, and serotonin EIA kit are available from Cayman Chemical Co. (Ann Arbor, MI, USA). Other reagents were purchased from Sigma Chemical Corporation (St. Louis, MO, USA). Antibodies and lysis buffers used for western blotting were acquired from Cell Signaling (Beverly, MA, USA). Enhanced chemiluminescence solution (ECL) and polyvinylidene difluoride (PVDF) membrane were purchased from General Electric Healthcare (Buckinghamshire, UK).

Preparation of washed human platelets

The Korean Red Cross Blood Center (Suwon, Korea) provided the human platelet-rich plasma (PRP). Washed platelets were prepared according to the method performed previously [22]. Platelet was obtained by centrifuging PRP at 1,300×g for 10 minutes, which was then washed two times with a wash buffer (138 mM NaCl, 12 mM NaHCO₃, 0.36 mM NaH₂PO₄, 2.7 mM KCl, 5.5 mM glucose and 1 mM Na₂EDTA, pH 6.9). Platelets were suspended with suspension buffer (138 mM NaCl, 12 mM NaHCO₃, 0.36 mM NaH₂PO₄, 2.7 mM KCl, 0.49 mM MgCl₂, 5.5 mM glucose, 0.25% gelatin, pH 7.4) resulting in the final concentration of 10⁶ cells/mL. All procedures were performed at 25°C to avoid platelet aggregation at low temperatures, and this experiment was performed with the approval of the Institutional Review Board (IRB) of Namsan University (1041479-HR-201803-003).

Measurement of platelet aggregation

In separate samples, with or without scoparone, the washed platelets (10⁶ cells/mL) were preincubated in 2 mM CaCl₂ at 37°C for a total of 3 min, and collagen (2.5 μg/mL) was then added to stimulate. To perform the platelet aggregation assay, an aggregometer was used for 5 min. The platelet aggregation rate (%) was determined by an increase in light transmission. The scoparone was dissolved using a solution of 0.1% dimethyl sulfoxide.

Measurement of cytotoxicity

Cytotoxicity was determined through the leakage of lactate dehydrogenase (LDH) from the cytosol. Human washed platelets (10⁶ cells/mL) were incubated for 2 h at room temperature with various concentrations of scoparone and then centrifuged at room temperature for 2 min at 12,000 g. The supernatant was measured by LDH assay kit at an optical density of 490 nm.

Measurement of TXB2

Incubation of the washed platelets (10⁶ cells/mL) was performed at 37°C for 3 min with the addition of various concentrations of scoparone, followed by the addition of 2 mM CaCl₂ and stimulation with collagen (2.5 μg/mL) for 5 min. Being a stable metabolite of TXA₂, the production of TXB₂ was measured by
synergy HT multi-reader (BioTek Instruments, Winooski, VT, USA) using the TXB₂ EIA kit.

**Measurement of ATP release**
The washed platelets (10⁸ cells/mL) were incubated at 37 °C for 3 min with the addition of various concentrations of scoparone, followed by the addition of 2 mM CaCl₂ and stimulation with collagen (2.5 μg/mL) for 5 min. After stopping the reaction with ice-cold 2 mM EDTA, the ATP released to the upper layer by centrifugation was measured using an ATP assay kit and a luminometer (BioTek Instruments, Winooski, VT, USA).

**Measurement of serotonin release**
The washed platelets (10⁸ cells/mL) had been incubated for 3 min at 37 °C with the addition of various concentrations of scoparone, followed by the addition of 2 mM CaCl₂ and stimulation with collagen (2.5 μg/mL) for 5 min. After stopping the reaction with ice-cold 2 mM EDTA, the serotonin released to the upper layer by centrifugation was measured by synergy HT multi-reader (BioTek Instruments, Winooski, VT, USA) using a serotonin EIA kit.

**Western Immunoblotting**
The addition of a 1× lysis buffer caused the termination of platelet aggregation. A BCA protein assay kit (Pierce Biotechnology, IL, USA) was used to measure and evaluate the resulting protein concentration of the platelet lysates. After proteins (20 μg) had been separated through 4-20% SDS-PAGE, they were relocated onto PVDF membranes. A dilution of 1:1,000 was set for the primary antibodies and the secondary antibodies were set at a dilution of 1:2,000. The use of ECL reagent (General Electric Healthcare, Buckinghamshire, UK) allowed protein band visualization.

**Statistical analyses**
The experimental results were expressed using mean ± standard deviation. Statistical analysis was completed using either ANOVA or unpaired Student's t-test. Scheffe's method was used to compare groups if there were significant differences between group means according to ANOVA.
Results

Effects of scoparone on collagen-induced human platelet aggregation

When platelets were treated with 2.5 μg/mL collagen to induce aggregation, the aggregation rate was 86.3±2.4%. When scoparone was added at various concentrations (10 to 300 μM), the increased aggregation by collagen was suppressed dose-dependently without cytotoxicity (Figs. 2B, C). At this time, the half-maximal inhibitory concentration (IC_{50}) of scoparone was found to be 46.52 μM (Fig. 2B). This shows that scoparone has a potent inhibitory effect on collagen-induced platelet aggregation.

Effects of scoparone on ATP release

The effect of scoparone on the release of ATP involved in platelet aggregation as an indicator of granule release in platelets was confirmed. As a result, collagen (2.5 μg/mL) increased ATP release from 0.23±0.01 μM in intact cells to 8.39±0.12 μM, which was 36.3 times higher than intact cells. However, scoparone (50, 100, 200 and 300 μM) significantly inhibited the increased ATP release by collagen at 6.53±0.21, 5.01±0.15, 3.03±0.13 and 1.34±0.02 μM, respectively (Fig. 4A).

Effects of scoparone on serotonin release

The effect of scoparone on the serotonin release involved in hemostasis through vascular contraction as an indicator of granule release in platelets was confirmed. As a result, collagen (2.5 μg/mL) increased serotonin release from 7.38±1.99 ng/10^8 cells in intact cells to 173.90±10.77 ng/10^8 cells, which was 23.6 times higher than intact cells. However, scoparone (50, 100, 200 and 300 μM) significantly inhibited the increased serotonin release by collagen to 128.83±9.07, 80.78±17.89, 48.04±3.48 and 27.84±1.00 ng/10^8 cells, respectively (Fig. 4B).

Effects of scoparone on PI3K and Akt phosphorylation

The effect of scoparone on the phosphorylation of PI3K and Akt, a phosphoprotein involved in the release of granules from platelets, was confirmed. As shown in Fig. 5, collagen significantly increased the phosphorylation of PI3K compared to intact cells. In addition, the increased phosphorylation of PI3K due to collagen was significantly decreased at concentrations with scoparone greater than 200 μM (Fig. 5). Collagen also significantly increased the phosphorylation of Akt, a PI3K target molecule, but significantly decreased at concentrations above 100 μM of scoparone (Fig. 5). This shows that scoparone inhibits the phosphorylation of PI3K and Akt promoted by collagen.

Effects of Scoparone on MAPK phosphorylation

The effect of scoparone on the phosphorylation of phosphoprotein MAPK (ERK, JNK, and p38) involved in platelet granule release and TXA_{2} production was identified. As shown in Fig. 6, Collagen significantly increased the phosphorylation of JNK and p38 compared to intact cells but did not significantly affect the...
phosphorylation of ERK. In addition, the phosphorylation of JNK or p38 increased by collagen was significantly inhibited at concentrations above 100 or 300 μM of scoparone, respectively (Fig. 6). In addition, scoparone significantly inhibited ERK phosphorylation at 200 μM (Fig. 6). This shows that scoparone regulates the signaling process of platelet aggregation by inhibiting phosphorylation of ERK, JNK, and p38, known as MAPK.

Discussion

Although scoparone has been reported to have an effect on platelet activation, the effects of scoparone on platelet aggregation and various signaling substances in the process, in particular phosphoproteins, have not been investigated, therefore the purpose of this study is to clarify the effects of scoparone in collagen-induced platelet aggregation.

PI3K/Akt pathway is well known as a phosphoprotein that functions in platelet signaling transduction, and its phosphorylation has been reported to play a major role in the regulation of platelet function, including the secretion of platelet dense granules and platelet aggregation [17]. In addition, MAPK are well known as phosphorylation enzymes, including ERK, JNK, and p38 MAPK, which are involved in platelet activation and aggregation [9]. MAPK is detected in human platelets and is reported to be activated through phosphorylation when platelets are activated by several agonists [10-12]. According to Mei-Chi et al., phosphorylation of MAPK such as p38 is critical for the release of arachidonic acid, a precursor of TXA₂, and the generation of TXA₂, leading to platelet aggregation. An important indicator in evaluating components or substances for platelet inhibitory activity is the generation of TXA₂ because TXA₂ acts as a powerful autacoid that additionally activates and aggregates other platelets [23]. Therefore, substances that inhibit the production of TXA₂ are usefully used as antiplatelet substances, and for example, substances such as aspirin and ozagrel are known [24,25].

In this study, scoparone had a significant concentration-dependent inhibitory effect on collagen-induced platelet aggregation, with a strong IC₅₀ of 14.05 μM (Fig. 2). The effects of scoparone on the production of TXA₂, an important marker of platelet aggregation, and the secretion of platelet granules (ATP and serotonin release) were measured. In addition, we tried to elucidate the relationship between scoparone and phosphorylation of PI3K/Akt and MAPKs. As the results, it was confirmed that scoparone suppressed
the production of strongly increased TXA$_2$ by collagen, and ATP and serotonin release, which are indicators of intracellular granule secretion, were strongly reduced by scoparone (Figs. 4B, C). In addition, it was confirmed that the phosphorylation of PI3K/Akt and MAPK, phosphoproteins known to regulate them as signaling molecules, was significantly inhibited by scoparone. In particular, in the case of ERK in MAPK, phosphorylation was not significantly caused by collagen, and the inhibitory effect by scoparone was also weak. There is a result that the degree of inhibiting phosphorylation of ERK in substances inhibited platelet aggregation is weaker than that of JNK or p38 [26]. It has also been reported that the inhibition of ERK pathway has no effect on agonist-induced aggregation of human platelets [27]. In this regard, it seems that inhibition of ERK pathway has no effect on agonist-induced aggregation of human platelets [27]. In this regard, it seems that inhibition of ERK pathway has no effect on agonist-induced aggregation of human platelets [27]. In any case, these results suggest that scoparone inhibits the phosphorylation of phosphoproteins such as PI3K/Akt and MAPK, thereby inhibiting platelet aggregation by reducing TXA$_2$ production and intracellular granule secretion (ATP and serotonin release).

In conclusion, we found that scoparone is valuable as an antiplatelet agent to regulate phosphorylation of phosphoproteins in signaling processes such as PI3K/Akt and MAPK. Therefore, scoparone is of great value as an effective therapeutic and prophylactic agent in cardiovascular diseases caused by platelet aggregation.

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Conflict of interest The authors declare no conflict of interest.

References