Up-regulation of Heme Oxygenase-1 by Korean Red Ginseng Water Extract as a Cytoprotective Effect in Human Endothelial Cells

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Korean red ginseng (KRG) is used worldwide as a popular traditional herbal medicine. KRG has shown beneficial effects on cardiovascular diseases, such as atherosclerosis, diabetes, and hypertension. Up-regulation of a cytoprotective protein, heme oxygenase (HO)-1, is considered to augment the cellular defense against various agents that may induce cytotoxic injury. In the present study, we demonstrate that KRG water extract induces HO-1 expression in human umbilical vein endothelial cells (HUVECs) and possible involvement of the anti-oxidant transcription factor nuclear factor-erythroid 2-related factor 2 (Nrf2). KRG-induced HO-1 expression was examined by western blots, reverse transcriptase polymerase chain reaction and immunofluorescence staining. Specific silencing of Nrf2 genes with Nrf2-siRNA in HUVECs abolished HO-1 expression. In addition, the HO inhibitor zinc protoporphyrin blunted the preventive effect of KRG on H2O2-induced cell death, as demonstrated by terminal transferase dUTP nick end labeling assay. Taken together, these results suggest that KRG may exert a vasculoprotective effect through Nrf2-mediated HO-1 induction in human endothelial cell by inhibition of cell death.

Keywords: Panax ginseng, Endothelial cells, Heme oxygenase, Oxidative stress, Vascular diseases

INTRODUCTION

Endothelium is a monolayer of endothelial cells lining the entire vascular system. The vascular endothelium plays an important role in regulation of thrombotic response, modulation of vascular tone and blood flow, and regulation of immune and inflammatory responses by controlling interactions between immune cells and the vessel wall [1].

Vascular injury, including structural and functional impairment of endothelium, plays a key role in the pathogenesis of various vascular diseases, such as atherosclerosis, diabetes, and hypertension. Chronic damage to endothelium causes accumulation of lipid and increased adhesion of monocytes and platelets [2]. Furthermore, injured endothelial cells release several growth factors, leading to later migration and proliferation of smooth muscle cells [3,4]. Finally, these changes in the vessel wall could accelerate atherosclerotic plaque formation.

Korean red ginseng (KRG) is used as a well-known herbal medicine as well as a wide range of food products, including beverages, candy, jellies, and snacks. Biological properties of KRG have been extensively investigated as its consumption has been increasing. Ginseng is thought to exert various effects, including anti-oxidant, anti-thrombotic, anti-hyperlipidemic, and anti-cancer...
effects [3-6]. Recently, the interest of researchers has focused on the vascular protective effect of KRG. In endothelial cells, KRG eliminates generation of NADPH-driven superoxide [7] and increases nitric oxide synthase activity and nitric oxide concentrations, conferring a hypotensive effect [8]. Previous studies have also reported that KRG promotes endothelial proliferation and protects H$_2$O$_2$-induced cell death [9,10]. Due to its vascular protective effect, KRG is believed to be beneficial for cardiovascular diseases such as atherosclerosis, diabetes, and hypertension [11-13].

Heme oxygenase (HO) is a cytoprotective protein that catalyzes heme degradation, yielding carbon monoxide (CO), iron, and biliverdin as the final products. There are three isoforms of HO in mammals: HO-1, HO-2, and HO-3 [14,15]. In particular, HO-1 is well known as a protective enzyme with anti-oxidant, anti-apoptotic, and anti-inflammatory effects [16]. HO-1 is a stress inducible protein. Various stimuli, such as thiol scavengers, ultraviolet radiation, and oxidative stress act as inducers of HO-1 [17]. In general, these stimuli cause generation of reactive oxygen species (ROS), which could activate an adaptive response of HO-1. Up-regulation of HO-1 exerts cytoprotective effects due to activity of its products, including CO and/or bilirubin, formed from biliverdin by the activity of biliverdin reductase [18,19]. HO-1 has a beneficial role in several clinically relevant diseases. There also is increasing evidence to show that HO-1 is associated with vascular disease, such as atherosclerosis, diabetes, and hypertension [20]. HO-1, which is highly expressed in vascular tissues, protects against vasculopathy and confers a cytoprotective function in the circulation [21]. In human endothelial cells, HO-1 deficiency results in tumor necrosis factor-α and interleukin-1α mediated endothelial damage [22]. These results suggest that HO-1 may play an important role in sustaining the health of the vascular system.

In this study, we examined the cardiovascular protective effect of HO-1 induced by KRG water extract in human umbilical vein endothelial cells (HUVECs). In addition, we investigated involvement of nuclear factor-erythroid 2-related factor 2 (Nrf2) in induction of HO-1 and the cytoprotective effect of HO-1 in HUVECs.

**MATERIALS AND METHODS**

**Materials**

Korean red ginseng powder was obtained from the Korea Ginseng and Tobacco Central Research Institute (Daejen, Korea). M199 medium and fetal bovine serum were purchased from Welgene Inc. (Daegu, Korea). TRIZol reagent was supplied by Invitrogen (Carlsbad, CA, USA). Zinc protoporphyrin (ZnPP) was provided from Sigma Chemical (St. Louis, MO, USA). TransPass R2 Transfection Reagent was obtained from New England Biolabs (Hercules, CA, USA). Anti-Nrf2 and anti-Lamin B were purchased from Santa Cruz Biotechnology (Santa Cruz, CA, USA). Anti-GAPDH was supplied by AbFrontier (Seoul, Korea). All other chemicals and reagents were of analytical grade.

**Preparation of red ginseng water extract**

For preparation of red ginseng water extract, we modified a method used in a previous study [23]. Korean red ginseng powder was soaked in water (1:25, W:W) for 3 h, then boiled for 40 min. Following centrifugation at 3,000 rpm for 60 min, supernatants of ginseng extract were further centrifugated at 10,000 g for 30 min and lyophilized. Ginseng extracts were dissolved in pure water immediately prior to the experiment.

**Cell culture**

HUVECs were maintained in M199 medium and supplemented with 10% fetal bovine serum, 1% penicillin and streptomycin, 10 ng/mL human fibroblast growth factor, and 18 mU/mL heparin. The cells were incubated at 37°C under 5% CO$_2$ atmosphere. HUVECs were grown to approximately 80% confluence, maintained with fresh medium described above, and subcultured every 2 to 3 d. The cells were used within passages 4 to 9 during these experiments [24].

**Western blot analysis**

We applied 20 μg of the whole cell lysate proteins to each lane and analyzed them with western blots. Western blot analysis was performed using monoclonal antibody against mouse HO-1 and monoclonal antibody against mouse glyceraldehyde-3-phosphate dehydrogenase. Horseradish peroxidase-conjugated anti-IgG antibodies were used as the secondary antibody to detect the abovementioned protein bands by enhanced chemiluminescence WESTSAVE-Up™ (Abfrontier, Seoul, Korea).

**RNA isolation and reverse transcriptase-polymerase chain reaction**

Cells were seeded in a 100 mm-diameter plate containing M199 medium. After 24 h, KRG was treated to a
final concentration of 0.5 mg/mL and the cells were incubated for 18 h. RNA extraction was achieved using 1 mL TRIzol reagent. The RNA pellets were washed in 70% ethanol, dried completely, and dissolved in diethylpyrocarbonate to inhibit RNase. Total RNA was quantified using a ND-100 spectrometer (NanoDrop Technologies, Wilmington, DE, USA). Polymerase chain reaction was performed using the synthesized cDNA as a template and using specific primers for HO-1 or β-actin as a loading control. The primer sequence for human HO-1 was 5′-ACATCTATGTGGCCCTGGAG-3′ (forward) and 5′-TGTTGGGAAGGTGAAGAAG-3′ (reverse). The amplified products were resolved by 1% agarose gel electrophoresis, stained with ethidium bromide, and photographed under ultraviolet light.

Immunofluorescence staining
HUVECs were cultured in a glass culture chamber slide (Falcon Plastics Inc., London, ON, Canada) and processed for immunofluorescence analysis. Immunofluorescence method as described previously method [25].

Nuclear factor-erythroid 2-related factor 2 silencing by siRNA
The cells were seeded on six-well plates at a density of 2×10⁵ cells/well in 2 mL of complete M199 medium. Cells were allowed to grow to 60% to 80% confluence before transfection with siRNA (SC-37049, Santa Cruz Biotechnology). For each transfection, 1,200 μL of the transfection medium was added with siRNA duplex/transfection reagent mix (TransPass R2 solution A + B), and the entire volume was added gently to the cells [26].

Measurement of intracellular reactive oxygen species generation
Intracellular ROS in H₂O₂ stimulated HUVECs is analyzed using DCF/DA staining. HUVECs were plated at a density of 4×10⁵ cells in 60 mm dishes. After 18 h incubation with KRG in the presence or absence of 1 μM ZnPP, cells were stained with 10 μM DCF/DA for 55 min, and then stimulated with 500 μM H₂O₂ for 5 min. After rinsing with PBS, the cells were examined by fluorescence microscopy.

Terminal transferase dUTP nick end labeling assay
Cells (2×10⁴ cells/300 μL/well) were seeded in an 8-well chamber slide and were washed with phosphate-buffered saline (PBS) and fixed in 4% formaldehyde for 1 h and then exposed permeabilizing solution (0.1% Triton X-100 in 0.1% sodium citrate buffer) for 2 min at 4°C. Cells were incubated with the terminal transferase dUTP nick end labeling (TUNEL) reaction mixture (Roche, Mannheim, Germany) for 1 h at 37°C in the dark. After rinsing with PBS, the cells were mounted and examined by fluorescence microscopy.

Statistical analysis
Statistical significance was estimated by student’s t-test and the results were expressed as mean±SD.

Fig. 1. Induction of heme oxygenase (HO)-1 protein by Korean red ginseng (KRG) in human umbilical vein endothelial cells (HUVECs). After treatment of HUVECs with various concentrations of KRG (A) at various time intervals (B), cell lysates were prepared and 20 μg samples of proteins were subjected to western blotting using the anti-HO-1 antibody and anti-glyceraldehyde-3-phosphate dehydrogenase (GAPDH) antibody as a loading control. Representative data from three independent experiments is shown. *p<0.05.
RESULTS

Effects of heme oxygenase-1 induction by Korean red ginseng in human umbilical vein endothelial cells

Up-regulation of HO-1 expression plays an important role in protecting cells against oxidative stress. We first demonstrated the effect of various concentrations of KRG on HO-1 induction. KRG increased HO-1 protein expression in a concentration-dependent manner (Fig. 1A). Cells were incubated with various concentrations of KRG for 18 h. Treatment of cells with KRG (0.5 mg/mL) also resulted in time-dependent increases in HO-1 protein expression in HUVECs (Fig. 1B). KRG also increased the HO-1 mRNA level, as demonstrated in Fig. 2. After incubation with various concentrations of KRG for 12 h, the HO-1 mRNA level increased in a concentration-dependent manner (Fig. 2A). KRG-treated HUVECs showed the HO-1 mRNA level in a time-dependent manner (Fig. 2B).

Immunofluorescence staining of heme oxygenase-1 in human umbilical vein endothelial cells

After KRG treatment, cells were fixed, and HO-1 localization in HUVECs was observed by immunofluorescence staining with an anti-HO-1 antibody, followed by a fluorescence-tagged secondary antibody. Immunofluorescence analysis showed that HO-1 protein levels were increased in HUVECs after treatment with KRG in a concentration-dependent manner (Fig. 3).

Nuclear factor-erythroid 2-related factor 2 siRNA blocks Korean red ginseng-induced heme oxygenase-1 protein expression

HO-1 is one of the major genes encoding phase II
detoxifying and anti-oxidant enzymes and is widely distributed in mammalian tissues. It is modulated by the Nrf2/Keap1 transcription factor system [27]. Thus, we attempted to determine whether KRG could activate Nrf2 in association with HO-1 up-regulation. For this, siRNA of Nrf2 was employed. With 15 nmol of Nrf2 siRNA, the increase in HO-1 protein was abolished (Fig. 4). These results indicate a key role for Nrf2 in regulation of KRG-induced HO-1 protein expression in HUVECs.

Protection against oxidative stress provided by heme oxygenase-1 induced by Korean red ginseng

ROS acts as a secondary messenger that mediates the signal network involved in growth, survival, and death. In particular, a number of evidences support that ROS that causes oxidative stress may play an essential role in mediating cell death in endothelial cells. To clarify whether KRG-induced HO-1 plays a significant role in ROS scavenging, which attenuates endothelial cell death, we conduct DCF/DA staining that detects intracellular ROS level (Fig. 5). An increase in intracellular ROS was observed in H₂O₂-stimulated cells. ROS generation was reduced by KRG treatment and reversed by ZnPP, the specific HO-1 inhibitor. These results imply that HO-1 induction by KRG prevents ROS production and by extension, ROS-mediated cell damage and/or death.

Increase of endothelial cell death by pharmacological inhibition of heme oxygenase-1

Endothelial damage has been thought to be the initial cause of development of vascular disorders. A previous study suggested that protection in human endothelial cells by blockade of oxidative stress could be an effective strategy in treatment of atherosclerosis [28]. To determine whether KRG-induced HO-1 up-regulation is responsible for the preventive effect of endothelial cell injury, we inhibited HO-1 enzymatic activity through treatment with the specific HO-1 inhibitor, ZnPP. KRG-stimulated cells were pre-incubated with or without 1 μM ZnPP, followed by treatment with 100 μM H₂O₂ for 5 min. The protective effect of KRG on H₂O₂-induced cell death and its blockage by HO-1 inhibitor were observed by fluorescence microscopy.

Fig. 4. Inhibition of heme oxygenase (HO)-1 expression by nuclear factor-erythroid 2-related factor 2 (Nrf2) siRNA in human umbilical vein endothelial cells (HUVECs). Cells were treated with Korean red ginseng (KRG) at the indicated concentration for 18 h. Transient transfection of HUVECs with Nrf2 siRNA inhibited expression of the HO-1 protein. A volume of 15 nmol of Nrf2 siRNA abrogates KRG-induced HO-1 up-regulation. Representative data from three independent experiments is shown. GAPDH, glyceraldehyde-3-phosphate dehydrogenase. *p<0.05.

Fig. 5. Intracellular reactive oxygen species level of Korean red ginseng (KRG)-treated human umbilical vein endothelial cells (HUVECs). KRG (0.5 or 1 mg/mL) stimulated HUVECs were pre-treated for 1 h with or without 1 μM zinc protoporphyrin (ZnPP). After 18 h incubation, HUVECs were treated with DCF/DA for 55 min, followed by the treatment with 100 μM H₂O₂ for 5 min. The protective effect of KRG on H₂O₂-induced cell death and its blockage by HO-1 inhibitor were observed by fluorescence microscopy.
H_{2}O_{2} treatment significantly increased TUNEL-positive cells, which was restored by KRG. However, ZnPP pre-treatment diminished the cytoprotective effect of KRG in H_{2}O_{2}-induced cell death. These results strongly suggest that the vascular protective effect of KRG is mediated by HO-1 activity.

**DISCUSSION**

In the present study, we investigated the induction of a cytoprotective enzyme, HO-1, by KRG water extract in human endothelial cells. We show that KRG increased both mRNA and the protein level of HO-1 and its cytoprotective effect in HUVECs.

Ginseng has been used for treatment of various diseases, including cardiovascular disorders, in East Asia for over 2,000 yr and has become the most famous medicinal plant in the world. KRG is prepared from fresh ginseng steamed and sun-dried. Under high pressure and high temperature, several components in ginseng could be chemically transformed, inducing better pharmacological activity, such as the anti-oxidant, anti-carcinogenic, and ameliorative effect on blood circulation, than white ginseng [29,30]. These changed constituents are mostly triterpene glycosides, called ginsenosides, which are major active ingredients of ginseng. Recently, the diverse effects of ginsenosides on endothelial cells have been extensively studied. Ginsenoside Rg_{3} has anti-inflammatory and anti-atherosclerotic activities by reduction of cell adhesion molecules and pro-inflammatory cytokines in human endothelial cells [31]. It also prevents endothelial cell death via inhibition of the mitochondrial apoptotic signaling pathway [32]. Another ginsenoside Rb_{1}, is also one of the major constituents of ginseng. Its cytoprotective effect in oxidized low-density lipoprotein-injuring endothelial cells has been previously studied *in vitro* [33]. Ginsenoside Rg_{3} induces vascular protein expression in human endothelial cells, which regulates proliferation and migration of endothelial cells [34]. Ginsenoside protopanaxatriol has been reported to prevent H_{2}O_{2}-induced endothelial cell death by modulation of intracellular redox status [10]. Other constituents of red ginseng include brown reaction products. Browning pigments in ginseng extract are increased when ginseng is steamed at 100°C, almost all of which are water soluble [35]. Water soluble browning reaction products isolated from KRG have been reported to have anti-oxidative effects due to their free radical scavenging activity [36]. It is assumed that the protective effect of KRG is attributed to the anti-oxidant and anti-apoptotic activity of these active components.

HO-1 has a key role in the cellular defense mechanism against oxidative stress generated by ROS. Increase of HO-1 activity protects against oxidative damage-induced cell death via the mechanism associated with free heme catabolism by HO-1 [37]. Numerous studies have supported the suggestion that anti-cell death activity of HO-1 represents its cytoprotective roles in the vascular system. A study of angiogenesis revealed that overexpression of HO-1 enhanced endothelial cell proliferation [38]. The beneficial effect of HO-1 is mediated by the actions of its byproducts, CO and bilirubin. CO changes blood fluidity and blood flow by modulation of vasomotor tone, vascular smooth muscle cell proliferation, and platelet agglutination [21]. Bilirubin exerts a vascular protective effect by preservation of EC integrity, prevention of EC death, and increase of vascular reactivity [39].

Previous evidence has indicated that HO-1 expression...
may be controlled at the transcriptional level by KRG. Hwang and Jeong [40] revealed that a key ginsenoside, Rb₁, induced HO-1 expression via Nrf2 and ARE pathways in 6-OHDA-treated human dopaminergic cells. Nrf2-mediated HO-1 induction by red ginseng extract in rat pheochromocytoma cells has also been reported [41]. However, there have been no reports revealing the mechanism underlying KRG induced HO-1 induction in human endothelial cells. We have determined that the major transcription factor of anti-oxidant gene Nrf2 is involved in this induction of HO-1 in HUVECs, confirming RNAi-mediated gene silencing. As mentioned above, Nrf2 promotes HO-1 expression induced by several ginseng constituents in various cell lines. Therefore, the transcriptional activation of Nrf2 following KRG water extract treatment may be associated with a noticeable increase in HO-1 expression in HUVECs.

In this study, we suggest that KRG water extract may exert a cytoprotective effect through HO-1 induction and that this up-regulation of HO-1 in HUVECs is mediated by the transcription factor Nrf2. This study supports a possible therapeutic mechanism of KRG in cardiovascular diseases.

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