Salidroside Attenuates Hydrogen Peroxide-Induced Cell Damage Through a cAMP-Dependent Pathway

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Received: 25 March 2011; in revised form: 8 April 2011 / Accepted: 8 April 2011 / Published: 19 April 2011

Abstract: Salidroside, a major component of *Rhodiola rosea* L., has shown various pharmacological functions, including antioxidant effects, but the signal transduction pathway of its antioxidant effects is not very clear. In this study, we found that salidroside could attenuate hydrogen peroxide (H$_2$O$_2$)-induced HL-7702 cell damage, inhibit H$_2$O$_2$-induced cytosolic free Ca$^{2+}$ ([Ca$^{2+}$]$_i$) elevation, scavenge reactive oxygen species (ROS) and increase 3’-5’-cyclic adenosine monophosphate (cAMP) level in a dose-dependent manner, but it couldn’t influence 3’-5’-cyclic guanosine monophosphate (cGMP) levels. Therefore, these results indicated that the antioxidant effects of salidroside were associated with down-regulation of [Ca$^{2+}$]$_i$, ROS occur via a cAMP-dependent pathway.

Keywords: salidroside; antioxidant; AMP; cGMP; [Ca$^{2+}$]$_i$; ROS
1. Introduction

*Rhodiola rosea* L. has been widely used for a long time in Traditional Chinese Medicine. Salidroside (\(p\)-hydroxyphenethyl-β-D-glucoside, C\(_{14}\)H\(_{20}\)O\(_{7}\), 300.30), as shown in Figure 1, has been identified as the most potent ingredient in this medicinal plant [1]. Some studies have shown that salidroside possesses various pharmacological functions, including anti-aging [2], anti-cancer [3-5], anti-viral [6-8], anti-inflammatory [9], neuroprotective [10,11], hepatoprotective [9], cardiomyocytes protective [12,13], anti-diabetic [14] and antioxidant effects [15,16].

Oxidative damage, mediated by reactive oxygen species (ROS) has been thought as a major cause of atherosclerosis, cancer, liver disorder and the aging process [17]. In previously studies, people found that the antioxidant mechanism included \([\text{Ca}^2+]i\) and NO signaling pathways [18,19], thus it might involve the 3′-5′-cyclic adenosine monophosphate (cAMP)/3′-5′-cyclic guanosine monophosphate (cGMP)-dependent signaling pathway. To the best of our knowledge, no one has investigated the role of cAMP/cGMP in the antioxidant effects of salidroside, so the objective of the present paper was to examine the effects of salidroside on \(\text{H}_2\text{O}_2\)-induced oxidative damage, cAMP, cGMP; \([\text{Ca}^2+]i\) and ROS levels in HL-7702 cells to determine whether the cAMP/cGMP pathway participate in the antioxidant effects of salidroside. Our results would be beneficial for development of salidroside as a potential agent for treatment of oxidative stress-related diseases.

![Figure 1. Structure of salidroside.](image)

2. Results and Discussion

2.1. Protective Effect of Salidroside on \(\text{H}_2\text{O}_2\)-Induced Cell Damage

The protective effect of salidroside on \(\text{H}_2\text{O}_2\)-induced cell damage as determined by the MTT assay is shown in Figure 2. Incubating with \(\text{H}_2\text{O}_2\) (0.5 mM) for 100 min induced significant cell damage. Pretreatment with salidroside (0.03, 0.05, 0.1 μg/mL) could attenuate \(\text{H}_2\text{O}_2\)-induced cell damage in a dose-dependent manner (\(p < 0.05\)), though it couldn’t restore the damaged cells totally. Data are expressed as mean ± S.D. (n = 4). \(* p < 0.05\) and \(** p < 0.01\) compared with Control.

2.2. Salidroside Inhibited \(\text{H}_2\text{O}_2\)-Induced \([\text{Ca}^2+]i\) Elevation

We examined the effect of salidroside on \([\text{Ca}^2+]i\). We found that incubating with \(\text{H}_2\text{O}_2\) (0.5 mM) for 100 min induced a significant \([\text{Ca}^2+]i\) elevation, while pre-incubation with salidroside dose dependently suppressed the elevation of \([\text{Ca}^2+]i\) (Figure 3). Data are expressed as mean ± S.D. (n = 4). \(* p < 0.05\) and \(** p < 0.01\) compared with Control; \(# p < 0.05\) and \(## p < 0.01\) compared with \(\text{H}_2\text{O}_2\) group.
**Figure 2.** Protective effect of salidroside on H$_2$O$_2$-induced cell damage. Cells were treated with salidroside followed by H$_2$O$_2$ as described in 2.2 and cell viability was determined by MTT assay.

![Figure 2: Bar chart showing OD value at 570 nm for different conditions.](image)

**Figure 3.** Salidroside inhibited H$_2$O$_2$-induced [Ca$^{2+}$]i elevation (Recorded by inverted fluorescence microscope, Motic AE31 200×). (a). Control; (b). H$_2$O$_2$; (c). 0.03 μg/mL salidroside + H$_2$O$_2$; (d). 0.05 μg/mL salidroside + H$_2$O$_2$; (e). 0.1 μg/mL salidroside + H$_2$O$_2$; (f). Statistical analysis of fluorescence intensity data with Motic Advanced Images 3.2 software (China). Cells were treated with salidroside followed by H$_2$O$_2$ as described in 2.2. and [Ca$^{2+}$]i levels were determined with Fluo-3/AM fluorescent dye.

![Figure 3: Images showing fluorescence under different conditions.](image)
2.3. Effects of Salidroside on the Formation of cAMP and cGMP

We investigated the effects of salidroside on the level of cAMP/cGMP. As shown in Figure 4, H₂O₂ decreased the levels of cAMP (Figure 4a) and cGMP (Figure 4b) in HL-7702 cells. The decreased level of cAMP was restored by treatment with salidroside, while the decreased level of cGMP was not changed. It showed that salidroside exerts antioxidant effects by cAMP, not cGMP. Data are expressed as absorbance density (OD) and presented as mean ± S.D. (n = 4). * p < 0.05 and ** p < 0.01 compared with Control; # p < 0.05 and ## p < 0.01 compared with H₂O₂ group.

Figure 4. (a). Effect of salidroside on cAMP level in cells; (b). Effect of salidroside on cGMP level in cells. Cells were treated with salidroside followed by H₂O₂ as described in 2.2. Contents of cAMP and cGMP were measured using human cyclic adenosine monophosphate Elisa kits and human cyclic guanosine monophosphate Elisa kits respectively.

2.4. Salidroside Inhibited H₂O₂-Induced ROS Elevation

We examined the effect of salidroside on intracellular ROS. We found that incubating with H₂O₂ (0.5 mM) for 100 min induced ROS elevation significantly, while pre-incubation with salidroside dose dependently suppressed the elevation of ROS (Figure 5). Statistical analysis of fluorescence intensity data with Motic Advanced Images 3.2 software. Data were expressed as mean ± S.D. (n = 4). * p < 0.05 and ** p < 0.01 compared with Control; # p < 0.05 and ## p < 0.01 compared with H₂O₂ group.
2.5. Discussion

Today, oxidative damages caused by ROS and oxidative stress play important roles in many diseases, such as endocrine disruption [18]. \( \text{H}_2\text{O}_2 \), a major source of ROS, has been used in many studies to trigger oxidative damage [19]. In order to further studying antioxidant mechanism of salidroside, we investigated the effects of salidroside on \( \text{H}_2\text{O}_2 \)-induced oxidative damage, cAMP, cGMP; \( [\text{Ca}^{2+}]_i \) and ROS level in HL-7702 cells.

In this paper, we used \( \text{H}_2\text{O}_2 \) 0.5 mM to trigger oxidative damage in HL-7702 cells. Incubating with \( \text{H}_2\text{O}_2 \) for 100 min induced significant cell damage. Salidroside (0.03, 0.05, 0.1 \( \mu \text{g/mL} \)) could attenuate \( \text{H}_2\text{O}_2 \)-induced damage in a dose-dependent manner. It is suggested that salidroside at dose of 0.03, 0.05, 0.1 \( \mu \text{g/mL} \) could efficiently protect cells from oxidative damage.

There is a mutual relationship between calcium and ROS signaling, which affects the local \( \text{Ca}^{2+} \) homeostasis [20]. \( \text{H}_2\text{O}_2 \) induced an \( [\text{Ca}^{2+}]_i \) rise which could still be evoked upon \( [\text{Ca}^{2+}]_i \) withdrawal and mitochondrial uncoupling [21]. In our research, 0.5 mM \( \text{H}_2\text{O}_2 \) successfully increased the level of \( [\text{Ca}^{2+}]_i \) in HL-7702 cells. \( [\text{Ca}^{2+}]_i \) levels are critical to cell life and changes in \( [\text{Ca}^{2+}]_i \) will affect matrix synthesis, as well as other functions [22]. In this experiment, salidroside inhibited \( \text{H}_2\text{O}_2 \)-induced \( [\text{Ca}^{2+}]_i \) and ROS level and might be used in \( \text{Ca}^{2+} \) overloading disease.

cAMP is one of the most important second messengers in regulating a wide variety of cellular events and processes, such as metabolism, gene expression, cell division, migration, exocytosis and secretion. In non-excitable cells the fluctuations of \( [\text{Ca}^{2+}]_i \) and [cAMP]\( i \) are often interrelated and

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**Figure 5.** Salidroside inhibited \( \text{H}_2\text{O}_2 \)-induced ROS elevation (Recorded by inverted fluorescence microscope, Motic AE31 200×). (a). Control; (b). \( \text{H}_2\text{O}_2 \); (c). 0.03 \( \mu \text{g/mL} \) salidroside + \( \text{H}_2\text{O}_2 \); (d). 0.05 \( \mu \text{g/mL} \) salidroside + \( \text{H}_2\text{O}_2 \); (e). 0.1 \( \mu \text{g/mL} \) salidroside + \( \text{H}_2\text{O}_2 \); (f). Statistical analysis of fluorescence intensity data with Motic Advanced Images 3.2 software. Cells were treated with salidroside followed by \( \text{H}_2\text{O}_2 \) as described in 2.2.
linked via adenylate cyclase or phosphodiesterase [23]. Elevation of the cAMP level will decrease the 
$[Ca^{2+}]_i$, and our result is consistent with this. The increased level of cAMP participates in activating 
PKA and consequently this enzyme phosphorylizes its substrate proteins. Thus, the increased level of 
cAMP is linked to the activation of PKA [24]. Salidroside influenced H$_2$O$_2$-induced cAMP level, so it 
might be linked PKA-mediated phosphorylation.

cGMP is an important secondary messenger synthesized by the guanylyl cyclases and is also 
included in many cellular events and processes. It is believed that cGMP is produced via the activation 
of guanylate cyclase in the presence or absence of NO. NO is produced by the activation of NOS and 
affect the formation of cGMP. cGMP could affect $[Ca^{2+}]_i$ through direct inhibition of L-type Ca$^{2+}$ 
channels [15]. cAMP and cGMP both are regulators of NO [25]. Salidroside couldn’t influence cGMP 
level, so the change of NO may be mediated via cAMP-depend pathway.

In the experiment, we found that salidroside attenuated H$_2$O$_2$-induced cell damage, inhibited 
H$_2$O$_2$-induced $[Ca^{2+}]_i$ elevation, scavenged ROS at low dose through cAMP-dependent pathway. These 
findings suggested that antioxidant protection of salidroside was associated with the cAMP-dependent 
pathway and decrease of $[Ca^{2+}]_i$ and ROS levels.

3. Experimental

3.1. Materials

Salidroside (standard material, purity >98%) was purchased from the National Institute for the 
Control of Pharmaceutical and Biological Products. 3-[4,5-Dimethyl-2-thiazolyl]-2,5-diphenyl-2-
tetrazolium bromide (MTT) was purchased from Sigma (USA). Fluo-3/AM was obtained from Dojin 
Laboratories (Japan). 5-(and-6)-Chloromethyl-2’,7’-dichlorodihydrofluorescein diacetate, acetyl ester 
(CM-H$_2$DCFDA) was obtained from Invitrogen (Molecular Probe), USA. RPMI 1640 was obtained 
from Gibco (USA). Human cyclic guanosine monophosphate ELISA kit and Human cyclic adenosine 
onophosphate ELISA kit were obtained from Blue Gene (Shanghai, China). FBS was obtained from 
Hangzhou (China). All other reagents were from commercial suppliers and of standard 
biochemical quality.

3.2. Protective Effect of Salidroside on H$_2$O$_2$-Induced Cell Damage

The human hepatocyte HL-7702 was obtained from SGST (China). The protective effect of 
salidroside was assessed by the MTT assay. Briefly, cells seeded on 96-well culture plates at 5 × 10$^4$/ well 
were incubated with salidroside for 24 h. Following exposure to H$_2$O$_2$ (0.5 mM) for 100 min, the MTT 
solution (10 μL, 5 mg/mL) was added into each well and made the final concentration 0.5 mg/mL, and 
then the plates were incubated for an additional 2 h. After the medium was removed, DMSO (100 μL) was 
added into each well before reading the microplates at 570 nm. Cell viability was expressed as absorbance 
density (OD).
3.3. Determination of [Ca$^{2+}$]i

The [Ca$^{2+}$]i was determined using Fluo-3/acetoxymethyl ester (Fluo-3/AM), described previously with slight modification [26]. Briefly, cells were loaded for 20 min at 37 °C with 20 μM Fluo-3/AM containing 1 μM pluronic acid F-127 for proper dispersal and 0.25 mM sulfinpyrazone, to inhibit the leakage of the Fluo-3 dye. Shortly before use, a sample of loaded cells was washed with Krebs-Ringer-Hepes (KRH) buffer (130 mM NaCl, 1.3 mM KCl, 2.2 mM CaCl$_2$, 1.2 mM MgSO$_4$, 1.2 mM KH$_2$PO$_4$, 10 mM Hepes, 10 mM glucose, pH 7.4), to remove nonhydrolyzed Fluo-3/AM. Fluorescence measurements were performed using an inverted fluorescence microscope (Motic AE31invert, Xiamen, China).

3.4. Measurement of cAMP and cGMP

Cells (5 × 10$^4$ /mL) were pre-incubated with various concentrations of salidroside for 48 h at 37 °C in 24-well culture plates, then incubated with H$_2$O$_2$ for 100 min. cAMP and cGMP were measured using the ELISA kits according to the manufacturer’s instructions. The levels of cAMP and cGMP were expressed by absorbance density (OD).

3.5. Measurement of ROS

The intracellular ROS levels of cells were determined using CM-H2DCFDA assay [26]. For these experiments, cells were incubated with phosphate-buffered saline (PBS) and 5 mM CM-H2DCFDA. As a control, cells were incubated with PBS only. As a positive control, cells were incubated with PBS, 5 mM CM-H2DCFDA, and 0.5 mM H$_2$O$_2$. After a 100 min incubation period with 0.5 mM H$_2$O$_2$, cells were trypsinized, washed, and resuspended in PBS. The levels of fluorescence intensity were immediately detected using an inverted fluorescent microscope (Motic AE31invert, Xiamen, China).

3.6. Statistical Analysis

All data were shown as means ± S.D. (standard deviation of the mean) Statistical analyses were performed using Student’s t-test and one-way analysis of variance (ANOVA). p < 0.05 was considered as statistically significant.

4. Conclusions

The data reported here firstly suggested that the antioxidant effects of salidroside were associated with down-regulation of [Ca$^{2+}$]i, ROS and at least partly via cAMP-dependent pathway.

Acknowledgements

This study was financed by the Chinese postdoctoral station of Jilin University (Number 801091800604).
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Sample Availability: Samples of the salidroside are available from the authors.

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