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# Benzyl salicylate from the stems and stem barks of *Cornus walteri* as a nephroprotective agent against cisplatin-induced apoptotic cell death in LLC-PK1 cells†

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Nephrotoxicity currently plagues the therapeutic use of cisplatin. Its major side effects are predominantly associated with renal tubular cell apoptosis. In this study, we examined the potential nephroprotective constituents from *C. walteri* that can exert effects against cisplatin-induced oxidative stress and apoptosis in renal tubular epithelial (LLC-PK1) cells. Phytochemical investigation of the MeOH extract of the stems and stem barks of *C. walteri* led to the isolation of the potential nephroprotective constituents (1–15). Among the isolates, treatment with benzyl salicylate (BS) (15) (50 and 100 μM) significantly attenuated LLC-PK1 cell apoptosis and potently blocked cisplatin-induced nuclei condensation, intracellular ROS generation, and apoptosis, as indicated by Hoechst 33342 and H<sub>2</sub>DCFDA staining. Furthermore, the expression of apoptosis-related proteins such as MAPK, Bax/Bcl-2 protein ratio, and cleaved caspase-9 and -3, were significantly decreased by BS. These findings provide that benzyl salicylate can be a nephroprotective agent against cisplatin-induced oxidative stress and apoptosis.

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## Introduction

Cisplatin (*cis*-diamminedichloroplatinum(II), CDDP) is very efficacious as a first-line therapy for malignant tumors in organs such as the testis, head, and ovaries.<sup>1,2</sup> However, cisplatin treatment results in not only the damage of tumor cells but also the tissue cells.<sup>3</sup> Cisplatin is especially toxic to the proximal tubule cells of the kidney, resulting in kidney dysfunction.<sup>4</sup> Recent cisplatin-targeted nephroprotection studies have focused on the search for natural products with enormous structural and chemical diversity,<sup>5,6</sup> because reducing cisplatin-induced nephrotoxicity will not only maximize its clinical outcomes, but can be beneficial to patients receiving cisplatin chemotherapy.

*Cornus walteri* Wanger, well-known as Walter's dogwood, is a deciduous shrub belonging to the family Cornaceae. *C. walteri* is distributed in Asian countries, specifically China and Korea as an economic crop. This tree is best known as a raw material for high-grade furniture and agricultural tools. It is also cultivated for its showy, edible berries. Traditionally, its fruits and leaves are used to treat dermatologic inflammation caused by

lacquer poison, as indicated in the Chinese Materia Medica.<sup>7</sup> Its leaves are used as an antidiarrheal in Korean folk medicine.<sup>8</sup> According to previous phytochemical studies, *C. walteri* possesses a variety of bioactive secondary metabolites such as lignans, isoquercitrin, and triterpenoids.<sup>9–11</sup> The extract of *C. walteri* has also been shown to inhibit NO production in lipopolysaccharide (LPS)-activated RAW 264.7 macrophages.<sup>12</sup> The leaves of this plant also have protective effects against UVB-induced photoaging in human reconstituted skin.<sup>7</sup> However, the stem of *C. walteri* has not been under-developed and utilized.

In our continued quest to discover biologically novel natural products from a variety of Korean natural resources,<sup>13–17</sup> our group has explored the potential bioactive constituents of *C. walteri*.<sup>11,18,19</sup> Our previous studies reported that the MeOH extract of the stems and stem barks of *C. walteri* are significantly cytotoxic.<sup>11</sup> The cytotoxic effects of triterpenoids and δ-valerolactones have also been elucidated in the our studies.<sup>11,18,19</sup> These findings were our motivation to further investigate potential bioactive constituents in the MeOH extract of *C. walteri*. Hence, in this study, we performed advanced phytochemical analyses to examine the potential nephroprotective constituents of the MeOH extract of *C. walteri*, leading to the successful isolation of 15 compounds (1–15). We therefore report the isolation and structural elucidation of these compounds and the evaluation of their protective effects against cisplatin-induced nephrotoxicity in renal tubular epithelial (LLC-PK1) cells.

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## Results and discussion

*C. walteri* stems and stem barks collected from Jeju Island, Korea were extracted with 80% aqueous MeOH under reflux, and then evaporated under a vacuum to obtain a MeOH crude extract, which was subsequently solvent-fractionated with hexane, CHCl<sub>3</sub>, and *n*-BuOH, yielding soluble fractions of each solvent. Significant spots were detected in the hexane-soluble fraction after TLC analysis, and the hexane-soluble fraction was subjected to repeated column chromatography and HPLC purification, which resulted in the isolation of 15 compounds, including seven triterpenoids (1–7), five steroids (8–12), two diterpene analogs (13 and 14), and one phenolic compound (15). Based on a comparison of their spectroscopic (including <sup>1</sup>H and <sup>13</sup>C NMR) and physical data with previously reported values and LC/MS analyses, the chemical structures of all the isolates were determined and identified as: leucophyllone (1),<sup>11</sup> 3-*O*-acetylbetulic acid (2),<sup>20</sup> betulinic acid methyl ester (3),<sup>21</sup> lupe- none (4),<sup>22</sup> methyl 3-*O*-acetylbetulinate (5),<sup>23</sup> lupeol (6),<sup>24</sup> betu- lincic acid (7),<sup>25</sup> 6β-hydroxysitostenone (8),<sup>26</sup> sitostenone (9),<sup>27</sup> 5α- stigmaster-3,6-dione (10),<sup>28</sup> 3β-sitostanol (11),<sup>29</sup> 6α-hydroxy-β- sitostenone (12),<sup>30</sup> norphytan (13),<sup>31</sup> phytone (14),<sup>32</sup> and benzyl salicylate (15)<sup>33</sup> (Fig. 1A).

Renal tubular cell death induced by oxidative stress and apoptosis is a common feature of cisplatin-induced

nephrotoxicity.<sup>4,34</sup> The effects of cisplatin and/or compounds isolated from *C. walteri* on cell viability were determined using the Ez-Cytox (tetrazolium salt, WST-1) cell viability assay. Among the isolated compounds (1–15) tested, 6β-hydroxysitostenone (HS) (8) and benzyl salicylate (BS) (15) displayed significant protective effects against cisplatin-induced renal tubular epithelial cell damage (Fig. S1†). As shown in Fig. 1, an approximately 40% significant decrease in cell viability was observed in the 25 μM cisplatin-treated group compared with the control group (100% cell viability). Meanwhile, co-treatment with HS (25, 50, and 100 μM) and BS (10, 25, 50, and 100 μM) significantly increased cell viability to 78.5 ± 1.6% and 91.4 ± 1.2%, respectively, compared with the respective cisplatin-treated groups (Fig. 1). Interestingly, 100 μM BS was as potent as 100 μM quercetin, a frequently studied compound with excellent antioxidant properties in *in vivo* and *in vitro* models of cisplatin-induced nephrotoxicity (Fig. 1).<sup>35–38</sup>

To determine the effects of BS (50 and 100 μM) on cisplatin-induced LLC-PK1 cell morphological changes and nuclear condensation, Hoechst 33342 staining was used. As shown in Fig. 2A, no changes were observed in the control group, whereas the cisplatin-treated group showed changes in cell shape and structure such as shrinkage as well as round and blebbing morphology. Furthermore, fluorescence microscopy images of the cisplatin-treated group had a higher bright blue-fluorescence, compared with the control group (Fig. 2B). Conversely, the BS-treated (50 and 100 μM) LLC-PK1 cells revealed attenuation of the cisplatin-induced changes in their cell shape, structure, and nuclear condensation, which were changes leading to apoptotic cell death (Fig. 2).<sup>39</sup>

ROS generation is one of the potential mechanisms for cisplatin-induced renal tubular epithelial cell damage and a hallmark of the induction of apoptosis.<sup>40,41</sup> The effects of cisplatin and/or BS on ROS generation were determined using H<sub>2</sub>DCFDA (2',7'-dichlorodihydrofluorescein diacetate). As indicated in Fig. 3A, fluorescence microscopy images of the cisplatin-treated group had a higher bright green-fluorescence compared with the control group because nonfluorescent

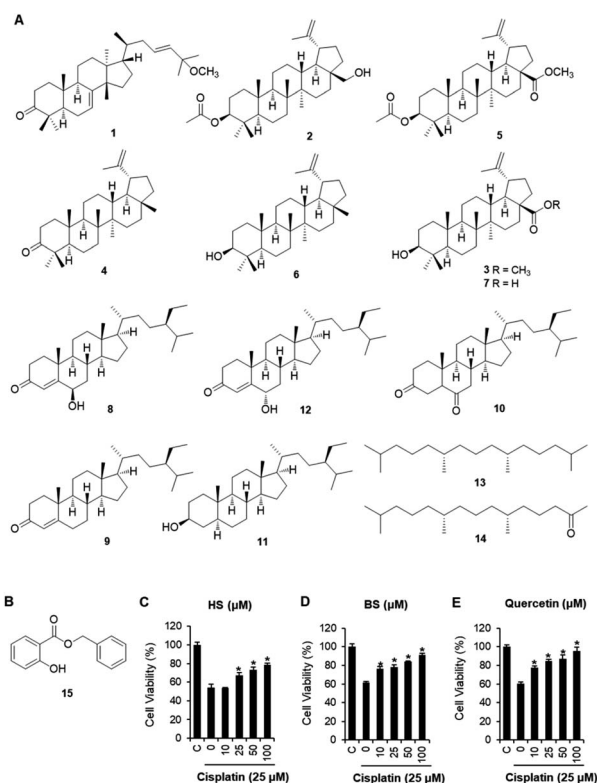


Fig. 1 Chemical structures of compounds 1–14 (A) and benzyl salicylate (BS) (15) (B). Protective effects of 6β-hydroxysitostenone (HS) (8) and BS (15), and quercetin (positive control, (E)) on the viability of cisplatin-damaged LLC-PK1 cells for 24 h, evaluated *via* the MTT assay (C and D). \**p* < 0.05 vs. the cisplatin-treated cells.

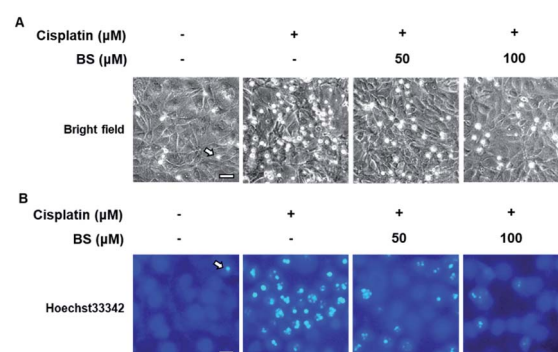


Fig. 2 Effects of benzyl salicylate (BS) on cisplatin-induced morphological changes and nuclear condensation in LLC-PK1 cells. (A) Representative phase contrast microscopy images of cells. (B) Representative fluorescence microscopy images of cells stained with Hoechst 33342. Scale bar, 50 μm.



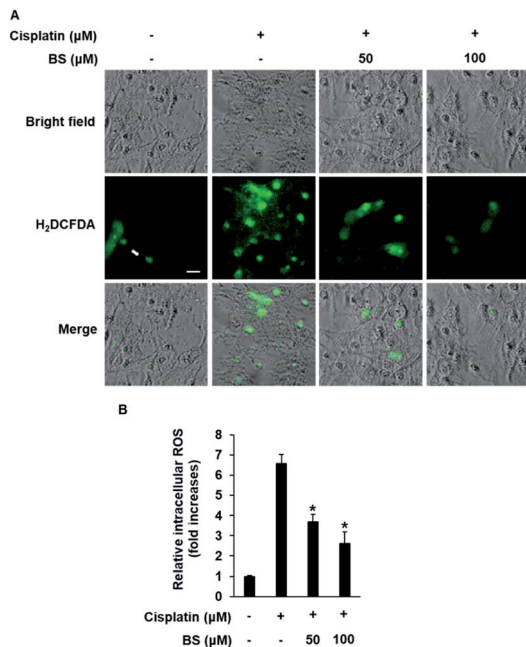


Fig. 3 Effect of BS on cisplatin-induced oxidative stress in LLC-PK1 cells. (A) Accumulation of ROS visualized with H<sub>2</sub>DCFDA. Scale bar, 50 μm. (B) Bar graph showing the fold increase in the intracellular ROS accumulation. \**p* < 0.05 vs. the cisplatin-treated cells.

H<sub>2</sub>DCFDA is converted to the fluorescent 2',7'-dichlorofluorescein (DCF) upon oxidation by ROS, whereas the BS-treated (50 and 100 μM) LLC-PK1 cells revealed attenuation of cisplatin-induced elevated intracellular ROS. As shown in Fig. 3B, quantitative analyses showed that the cisplatin-induced elevated intracellular ROS levels (7.37-fold increase) were significantly reduced by BS (50 and 100 μM) (4.20- and 2.68-fold increases, respectively). Cisplatin intensifies the generation of kidney oxidative stress, associated with the decrease of proximal tubule cell viability through mitochondria and DNA dysfunction.<sup>5</sup> Therefore, BS is a promising antioxidant agent for cisplatin-induced oxidative stress. The finding is in accordance with the study reported previously where quercetin showed the nephroprotective effect in cisplatin-treated rats because of its radical scavenging activity in the kidney tubular cells of rats.<sup>35</sup>

The protective effects of BS (50 and 100 μM) were investigated on cisplatin-induced cell death process in LLC-PK1 cells using staining with annexin V conjugated with Alexa Fluor 488 that can detect phosphatidylserine on the cell surface of apoptotic cells and propidium iodide (PI) that can detect necrotic cells by staining DNA and RNA inside of necrotic cells. Increased apoptosis (bright green color) was observed in the cisplatin-treated group compared with the control group (Fig. 4A). Conversely, the BS-treated (50 and 100 μM) LLC-PK1 cells attenuated the cisplatin-induced apoptosis. These results indicate that BS (50 and 100 μM) protects against cisplatin-induced LLC-PK1 cell death by inhibiting apoptosis. Quantitative analyses showed a significant 61.45 ± 4.78% increase in the percentage of apoptotic cells stained with annexin V in the 25 μM cisplatin-treated group compared with the control group

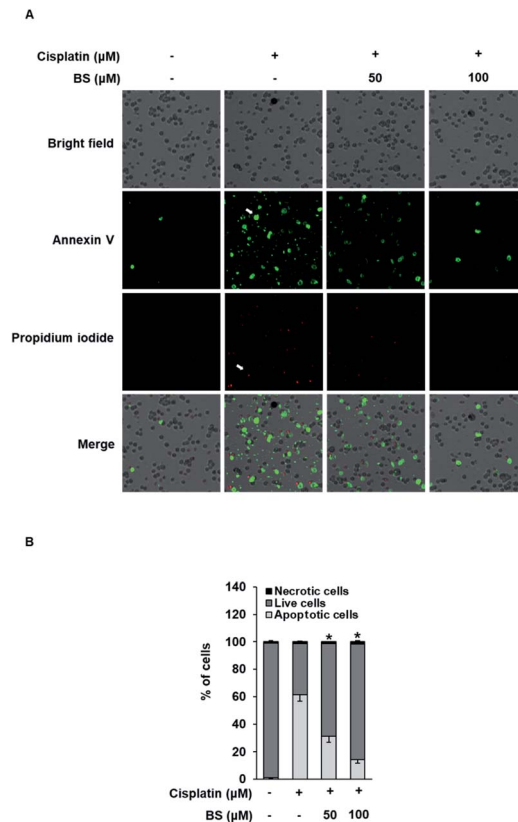


Fig. 4 Effect of BS on cisplatin-induced apoptosis in LLC-PK1 cells. (A) Visualization of apoptotic (green circles), live, and necrotic cells (red circles) cells (40× magnification). (B) Bar graph showing percentage of apoptotic, live, and necrotic cells. \**p* < 0.05 vs. cisplatin-treated cells.

(1.25 ± 0.52%; Fig. 4B). Conversely, BS treatment (50 and 100 μM) significantly decreased the percentage of apoptotic cells to 30.91 ± 4.09% and 14.20 ± 2.40%, respectively, compared with the cisplatin-treated group. In addition, quantitative analyses showed a significant decrease in the percentage of live cells in the 25 μM cisplatin-treated group (37.21 ± 0.05%) compared with the control group (97.75 ± 0.01%; Fig. 4B). Conversely, 50 and 100 μM BS treatments significantly increased the percentage of live cells to 67.75 ± 0.05% and 84.13 ± 0.03%, respectively, compared with the cisplatin-treated group. Percentage of necrotic cells stained with PI did not show significant changes. These results indicate that BS inhibited cisplatin-induced apoptosis. The finding is in agreement with earlier studies, where low concentration of cisplatin induces apoptotic cell death, whereas higher concentration of cisplatin induces necrotic cell death.<sup>42</sup>

To investigate the protective effects of BS (50 and 100 μM) on the underlying apoptotic pathway, we focused on evaluating the regulation of the mitogen-activated protein kinase (MAPK) pathways such as extracellular signal-regulated kinases (ERK), c-Jun N-terminal kinases (JNK), and p38. ERK, JNK, and p38 are important for the stress response and cell death, thus, are essential factors in cisplatin-induced apoptosis in renal tubular epithelial cells.<sup>34,43</sup> A previous study revealed that the anti-



apoptotic effect of quercetin in the cisplatin-treated rats is caused by suppression of the JNK.<sup>38</sup> BS treatment (50 and 100  $\mu\text{M}$ ) attenuated protein expression levels of MAPKs (p-JNK, p-ERK, and p-p38), compared with the levels in the cisplatin-treated group (Fig. 5A). Overall, the observed downregulation of JNK, p-ERK, and p-p38 by BS treatment (50 and 100  $\mu\text{M}$ ) may be indicative of its protective, anti-apoptotic role in cisplatin-induced LLC-PK1 cell damage.

We also investigated the effects of BS (50 and 100  $\mu\text{M}$ ) on Bax/Bcl-2 and cleaved caspase-8, -9, and -3 activation. In the apoptosis pathway, the anti-apoptotic protein Bcl-2 and pro-apoptotic protein Bax are apoptotic regulatory proteins. Down-regulation of Bcl-2 inhibits renal tubular epithelial cell death. When apoptosis is initiated, Bax is translocated to the mitochondrial membrane, leading to subsequent cisplatin-induced activation of caspases in renal tubular epithelial cells.<sup>42,44</sup> Compared with the cisplatin-treated group, BS treatment (50 and 100  $\mu\text{M}$ ) significantly inhibited the expression of Bax and increased the expression of Bcl-2, which resulted in a decreased Bax/Bcl-2 ratio (Fig. 6A). In addition, activated initiator caspases-8, and -9, and executioner caspase-3 were observed in cisplatin-induced renal tubular epithelial cell death.<sup>42,45</sup> BS treatment (50 and 100  $\mu\text{M}$ ) significantly attenuated the protein expression of cleaved caspase-9 and -3, compared with the levels in the cisplatin-treated group. However, activation of caspase-8 were not regulated significantly by treatment of BS. Overall, our results indicate that BS treatment (50 and 100  $\mu\text{M}$ ) modulated

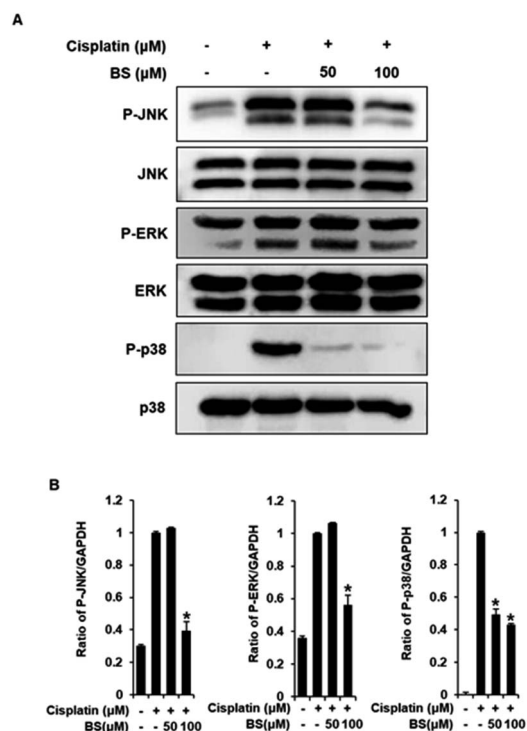


Fig. 5 Effects of BS on the levels of p-JNK, p-p38, and p-ERK in LLC-PK1 cells with cisplatin-induced damage. Immunoreactive bands of (A) p-JNK, p-p38, and p-ERK detected using western blot analyses. (B) Quantitative analyses of relative protein levels. \* $p < 0.05$  vs. the cisplatin-treated cells.

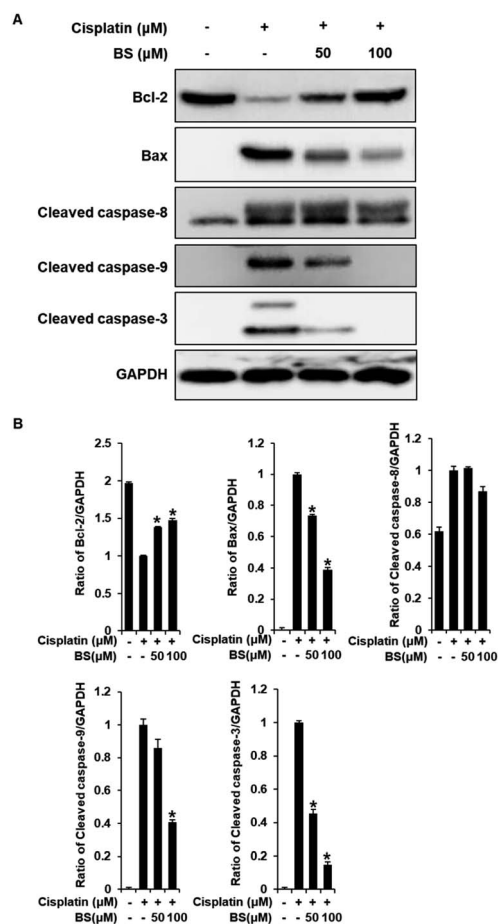


Fig. 6 Effects of BS on the levels of Bcl-2, Bax, cleaved caspase-3, -9, and -8 in LLC PK1 cells with cisplatin-induced damage. Immunoreactive bands of (A) Bcl-2, Bax, and cleaved caspase-8, -9, and -3 detected using western blot analyses. (B) Quantitative analyses of relative protein levels. \* $p < 0.05$  vs. the cisplatin-treated cells.

the Bax/Bcl-2 ratio by down-regulating the activation of caspase-9 and -3. Therefore, the protective effect of BS against cisplatin-induced renal tubular epithelial cell damage was mediated through intrinsic apoptotic pathway.

## Conclusion

In this study, we examined the potential of constituents of *C. walteri* against cisplatin-induced nephrotoxicity in LLC-PK1 cells, which led to the successful identification of the renoprotective compound benzyl salicylate (BS). BS inhibited cisplatin-induced oxidative stress and apoptosis in renal tubular epithelial cells. In addition, the anti-apoptotic mechanism employed by BS was attributed to the regulation of the protein expression of Bax, Bcl-2, cleaved caspase-9 and -3. These results suggest BS as a novel promising renoprotective agent against cisplatin-induced nephrotoxicity. Furthermore, further *in vivo* and *in vitro* studies are required to validate the mechanisms by which BS protects against cisplatin-induced renal tubular epithelial cell death. Our present study further supported the



potential of BS as a promising therapeutic candidate that benefits to nephroprotective effects.

## Experimental section

### General experimental procedures

Optical rotations were measured using a Jasco P-1020 polarimeter (Jasco, Easton, MD, USA). IR spectra were recorded with a Bruker IFS-66/S FT-IR spectrometer (Bruker, Karlsruhe, Germany). UV spectra were obtained using an Agilent 8453 UV-visible spectrophotometer (Agilent Technologies, Santa Clara, CA, USA). Nuclear magnetic resonance (NMR) spectra were recorded with a Bruker AVANCE III 700 NMR spectrometer operating at 500 MHz ( $^1\text{H}$ ) and 125 MHz ( $^{13}\text{C}$ ) (Bruker, Karlsruhe, Germany). Semi-preparative HPLC was performed using a Gilson 306 pump with a Shodex refractive index detector at a flow rate of  $2\text{ mL min}^{-1}$ . LC/MS analyses were performed on an Agilent 1200 Series HPLC system equipped with a diode array detector and 6130 Series ESI mass spectrometer, using an analytical Kinetex C18 100 Å column ( $100 \times 2.1\text{ mm i.d.}, 5\ \mu\text{m}$ ; Phenomenex, Torrance, CA, USA). Silica gel 60 (70–230 mesh and 230–400 mesh; Merck, Darmstadt, Germany) and RP-C<sub>18</sub> silica gel (Merck, 40–63  $\mu\text{m}$ ) were used for column chromatography. Merck precoated silica gel F<sub>254</sub> plates and RP-18 F<sub>254s</sub> plates were used for TLC. Spots were detected after TLC under UV light or by heating after spraying with anisaldehyde-sulfuric acid.

### Plant material

*C. walteri* stems and stem barks were collected from Jeju Island, Korea, in October 2005, and identified by one of the authors (K. H. Kim). A voucher specimen (SKKU 2005-10a) was deposited at the herbarium of the School of Pharmacy, Sungkyunkwan University, Suwon, Korea.

### Extraction and isolation

*C. walteri* stems and stem barks (2.5 kg) were dried, chopped, and crude extracts were prepared using 80% aqueous MeOH (2  $\times$  6 h) under reflux, and then filtered. The filtrate was evaporated under a vacuum to obtain a MeOH extract (220 g), which was suspended in distilled H<sub>2</sub>O (7.2 L) and then successively partitioned with hexane, CHCl<sub>3</sub>, and *n*-BuOH, yielding 9.5, 25.0, and 43.0 g of residue, respectively. The hexane-soluble fraction (9.5 g) was chromatographed on a silica gel (230–400 mesh, 300 g) column eluted with hexane–EtOAc (3 : 1 to 1 : 1, gradient system), to yield fractions H1–H5. Fraction H1 (4.0 g) was separated using an RP-C<sub>18</sub> silica gel column using 100% MeOH, to acquire five fractions (H11–H15). Fraction H12 (800 mg) was subjected to a Sephadex LH-20 column using a solvent system of dichloromethane–MeOH (1 : 1), to give subfractions H121–H122. To obtain compound 15 (12 mg), fraction H122 (50 mg) was subjected to a silica gel Waters Sep-Pak Vac 6 cc with hexane–EtOAc (30 : 1). Fraction H13 (800 mg) was further separated using silica gel column chromatography eluting a gradient solvent system of hexane–EtOAc (7 : 1 to 1 : 1, v/v), to provide 6 subfractions (H131–H136). Compound 12 (6 mg) was

purified from subfraction H131 (100 mg) using semi-preparative normal-phase HPLC (Apollo Silica column,  $250 \times 10.0\text{ mm}$ ,  $5\ \mu\text{m}$ , flow rate:  $2\text{ mL min}^{-1}$ ) and eluted with an isocratic solvent system of hexane–EtOAc (18 : 1, v/v). Five subfractions (H141–H145) were acquired from fraction H14 (300 mg) using silica gel column chromatography with a gradient solvent system of hexane–EtOAc (16 : 1 to 1 : 1, v/v). To obtain compound 2 (40 mg), subfraction H142 (50 mg) was isolated using semi-preparative normal-phase HPLC (Apollo Silica column,  $250 \times 10.0\text{ mm}$ ,  $5\ \mu\text{m}$ , flow rate:  $2\text{ mL min}^{-1}$ ) with an isocratic solvent system of hexane–EtOAc (28 : 1, v/v). Compound 13 (30 mg) was obtained from subfraction H144 (50 mg) using semi-preparative normal-phase HPLC (Apollo Silica column,  $250 \times 10.0\text{ mm}$ ,  $5\ \mu\text{m}$ , flow rate:  $2\text{ mL min}^{-1}$ ) eluted with an isocratic solvent system of hexane–EtOAc (12 : 1, v/v). To obtain compound 3 (7 mg), fraction H15 (100 mg) was separated using a C18 Sep-Pak column eluting an isocratic solvent system of hexane–EtOAc (10 : 1, v/v) and further isolated using semi-preparative normal-phase HPLC (Apollo Silica column,  $250 \times 10.0\text{ mm}$ ,  $5\ \mu\text{m}$ , flow rate:  $2\text{ mL min}^{-1}$ ) with an isocratic solvent system of hexane–EtOAc (20 : 1, v/v). Fraction H2 (2.5 g) was loaded onto an RP-C<sub>18</sub> silica gel column and separated by eluting with 100% MeOH to obtain eight fractions (H21–H28). Compound 8 (60 mg) was isolated from fraction H27 (150 mg) using semi-preparative normal-phase HPLC (Apollo Silica column,  $250 \times 10.0\text{ mm}$ ,  $5\ \mu\text{m}$ , flow rate:  $2\text{ mL min}^{-1}$ ) with an isocratic solvent system of hexane–EtOAc (8 : 1, v/v). Fraction H28 (100 mg) was isolated using semi-preparative normal-phase HPLC (Apollo Silica column,  $250 \times 10.0\text{ mm}$ ,  $5\ \mu\text{m}$ , flow rate:  $2\text{ mL min}^{-1}$ ) with an isocratic solvent system of hexane–EtOAc (8 : 1, v/v) to obtain compound 7 (25 mg). Fraction H3 (1.7 g) was loaded onto an RP-C<sub>18</sub> silica gel column and fractionated by eluting with 100% MeOH to obtain six fractions (H31–H36). Fraction H32 (200 mg) was separated by reverse-phase Lobar column with an isocratic solvent system of hexane–EtOAc (4 : 1, v/v) to obtain four subfractions (H321–H324). Compound 5 (4 mg) was obtained from subfraction H321 (150 mg) using semi-preparative reverse-phase HPLC (Econosil C18 column,  $250 \times 10.0\text{ mm}$ ,  $5\ \mu\text{m}$ , flow rate:  $2\text{ mL min}^{-1}$ ) eluting with an isocratic solvent system of 95% MeOH. Fraction H33 (400 mg) was fractionated using a reverse-phase Lobar column with an isocratic solvent system of hexane–EtOAc (4 : 1, v/v) to obtain five subfractions (H331–H335). Fraction H331 (50 mg) was purified using semi-preparative reverse-phase HPLC (Econosil C18 column,  $250 \times 10.0\text{ mm}$ ,  $5\ \mu\text{m}$ , flow rate:  $2\text{ mL min}^{-1}$ ) with an isocratic solvent system of 100% MeOH to give compound 1 (7 mg). Compound 14 (30 mg) was obtained from fraction H34 (120 mg) using semi-preparative normal-phase HPLC (Apollo Silica column,  $250 \times 10.0\text{ mm}$ ,  $5\ \mu\text{m}$ , flow rate:  $2\text{ mL min}^{-1}$ ) with an isocratic solvent system of hexane–EtOAc (3 : 1, v/v). Compounds 9 (5 mg) and 10 (8 mg) were purified from fraction H36 (120 mg) using semi-preparative normal-phase HPLC (Apollo Silica column,  $250 \times 10.0\text{ mm}$ ,  $5\ \mu\text{m}$ , flow rate:  $2\text{ mL min}^{-1}$ ) with an isocratic solvent system of hexane–EtOAc (4 : 1, v/v). Fraction H4 (1.3 g) was loaded onto an RP-C<sub>18</sub> silica gel column and fractionated by eluting with 100% MeOH to give six fractions (H41–H46).



Fraction H44 (120 mg) was isolated using semi-preparative normal-phase HPLC (Apollo Silica column, 250 × 10.0 mm, 5 μm, flow rate: 2 mL min<sup>-1</sup>) with an isocratic solvent system of hexane–EtOAc (2 : 1, v/v) to obtain compounds **4** (5 mg) and **6** (25 mg). Fraction H5 (500 mg) was fractionated using an RP-C18 silica gel column with an isocratic solvent system of 85% MeOH to obtain five fractions (H51–H55). Fraction H51 (150 mg) was passed over the Sephadex-LH20 column with CHCl<sub>3</sub>–MeOH (1 : 1, v/v) and further purified using semi-preparative reverse-phase HPLC (Econosil C18 column, 250 × 10.0 mm, 5 μm, flow rate: 2 mL min<sup>-1</sup>) eluting with an isocratic solvent system of 100% MeOH to obtain compound **11** (6 mg).

### Cell culture

The pig kidney epithelial cell line (LLC-PK1) was used to investigate the nephroprotective effect against cisplatin-induced renal cell damage. LLC-PK1 cells were purchased (American Type Culture Collection, Rockville, MD, USA) and grown to 80% confluence with DMEM containing 10% FBS, 1% penicillin/streptomycin, and 4 mM L-glutamine at 37 °C with 5% carbon dioxide and 95% relative humidity.

### Renoprotective effect against cisplatin-induced kidney cell damage

LLC-PK1 cells were plated onto 96-well culture plates (1 × 10<sup>4</sup> cells per well). After 24 h, they were exposed to various concentrations of the isolated compounds in the presence of 25 μM cisplatin for 24 h. The relative percent cell viability was assessed with the Ez-Cytox reagent kit (Daeil Lab Service, Seoul, Korea), following the manufacturer's instructions. The optical densities (OD) were measured at 450 nm using a microplate reader (PowerWave XS; Bio-Tek Instruments, Winooski, VT, USA).

### Determination of intracellular ROS

The cells were plated onto 6-well culture plates (1 × 10<sup>4</sup> cells per well). After 24 h, they were exposed to 50 or 100 μM benzyl salicylate (BS) in the presence of 25 μM cisplatin for 24 h. The LLC-PK1 cells were then stained with 2',7'-dichlorofluorescein diacetate (H<sub>2</sub>DCFDA; Sigma Aldrich, St. Louis, MO, USA) for 30 min at 37 °C and then washed with PBS. The cells were visualized under the IX50 fluorescent microscope (Olympus, Tokyo, Japan) equipped with a CCD camera, and fluorescence intensities of DCF at 495 nm excitation and 517 nm emission were measured using a fluorescence microplate reader (SPARK 10M; Tecan, Männedorf, Switzerland).

### Image-based cytometric assay

To detect the apoptotic cell population, LLC-PK1 cells were plated onto 6-well culture plates (4 × 10<sup>5</sup> cells per well). After 24 h, LLC-PK1 cells were treated with 50 or 100 μM BS in the presence of 25 μM cisplatin. The LLC-PK1 cells were then incubated with Annexin V-Alexa Fluor 488 and PI, following the manufacturer's instructions. The relative percentage of

apoptotic, live, and necrotic cells was assessed with the Tali image-based cytometer (Invitrogen, Carlsbad, CA, USA).

### Western blotting

The cells were plated into 6-well culture plates (4 × 10<sup>5</sup> cells per well). After 24 h, they were treated with 50 or 100 μM BS in the presence of 25 μM cisplatin. Following the manufacturer's instructions, the LLC-PK1 cells were then lysed with RIPA buffer (Cell Signaling Technology, MA, USA) supplemented with 1 mM phenylmethylsulfonyl fluoride for 20 min at 4 °C to obtain the cell lysates.

The protein concentrations of cell lysates were quantified using the bicinchoninic acid (BCA) protein assay kit (Thermo Fisher Scientific, Waltham, MA, USA). Equal amounts of proteins (20 μg of protein/lane) were separated using electrophoresis in a precast 4–15% Mini-PROTEAN TGX gel (Bio-Rad, Hercules, CA, USA) and transferred to a polyvinylidene difluoride membrane (Merck Millipore, Darmstadt, Germany). Membranes were incubated overnight at 4 °C with rabbit anti-p44/42 MAPK (Erk1/2) antibody (1 : 1000; Cat. no. 4695S; Cell Signaling Technology, Inc., Beverly, MA, USA), rabbit anti-phospho-p44/42 MAPK (Erk1/2) (Thr202/Tyr204) antibody (1 : 1000; Cat. no. 8544S; Cell Signaling Technology, Inc.), rabbit anti-SAPK/JNK antibody (1 : 1000; Cat. no. 9252S; Cell Signaling Technology, Inc.), rabbit anti-phospho-SAPK/JNK (Thr183/Tyr185) antibody (1 : 1000; Cat. no. 9251S; Cell Signaling Technology, Inc.), rabbit anti-p38 MAPK antibody (1 : 1000; Cat. no. 8690S; Cell Signaling Technology, Inc.), rabbit anti-phospho-p38 MAPK (Thr180/Tyr182) antibody (1 : 1000; Cat. no. 4511S; Cell Signaling Technology, Inc.), rabbit anti-Bcl-2 antibody (1 : 1000; Cat. no. 3498S; Cell Signaling Technology, Inc.), rabbit anti-Bax antibody (1 : 1000; Cat. no. 14796S; Cell Signaling Technology, Inc.), rabbit anti-cleaved caspase-8 antibody (1 : 1000; Cat. no. 8592S; Cell Signaling Technology, Inc.), rabbit anti-cleaved caspase-9 antibody (1 : 1000; Cat. no. 20750S; Cell Signaling Technology, Inc.), rabbit anti-cleaved caspase-3 antibody (1 : 1000; Cat. no. 9664S; Cell Signaling Technology, Inc.), rabbit anti-GAPDH antibody (1 : 1000; Cat. no. 2118S; Cell Signaling Technology, Inc.), and anti-rabbit HRP-conjugated IgG (1 : 2000; Cat. no. 7074S; Cell Signaling Technology, Inc.). Protein bands were visualized with ECL Advance western blotting Detection Reagents (GE Healthcare, Cambridge, UK) and a FUSION Solo Chemiluminescence System (PEQLAB Biotechnologie GmbH, Erlangen, Germany).

### Statistical analyses

Statistical significance was determined using analysis of variance (ANOVA), followed by a multiple comparison test with Bonferroni's adjustment. A *p*-value <0.05 was considered statistically significant.

### Conflicts of interest

The authors declare no competing financial interests.



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