Anti-breast cancer activity of Fine Black ginseng (Panax ginseng Meyer) and ginsenoside Rg5

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A B S T R A C T

Background: Black ginseng (Ginseng Radix nigra, BG) refers to the ginseng steamed for nine times and fine roots (hairy roots) of that is called fine black ginseng (FBG). It is known that the content of saponin of FBG is higher than that of BG. Therefore, in this study, we examined antitumor effects against MCF-7 breast cancer cells to target the FBG extract and its main component, ginsenoside Rg5 (Rg5).

Methods: Action mechanism was determined by MTT assay, cell cycle assay and western blot analysis.

Results: The results from MTT assay showed that MCF-7 cell proliferation was inhibited by Rg5 treatment for 24, 48 and 72 h in a dose-dependent manner. Rg5 at different concentrations (0, 25, 50 and 100 μM), induced cell cycle arrest in G0/G1 phase through regulation of cell cycle-related proteins in MCF-7 cells. As shown in the results from western blot analysis, Rg5 increased expression of p53, p21WAF1/CIP1 and p15INK4B and decreased expression of Cyclin D1, Cyclin E2 and CDK4. Expression of apoptosis-related proteins including Bax, PARP and Cytochrome c was also regulated by Rg5. These results indicate that Rg5 stimulated cell apoptosis and cell cycle arrest at G0/G1 phase via regulation of cell cycle-associated proteins in MCF-7 cells.

Conclusion: Rg5 promotes breast cancer cell apoptosis in a multi-path manner with higher potency compared to 20(S)-ginsenoside Rg3 (Rg3) in MCF-7 (HER2+/ER+) and MDA-MB-453 (HER2+/ER−) human breast cancer cell lines, and this suggests that Rg5 might be an effective natural new material in improving breast cancer.

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1. Introduction

Ginseng (Panax ginseng Meyer) is a well characterized medicinal herb listed in the classic oriental herbal dictionary, Shin-nong-boncho-kyung. Ginseng has a sweet taste, is able to keep the body warm, and has protective effects on the five viscera (i.e., heart, lung, liver, kidney, and spleen) [1]. Ginseng can be classified by how it is processed. Red ginseng (RG; Ginseng Radix Rubra) refers to ginseng that has been steamed once. White ginseng (Ginseng Radix Alba) refers to dried ginseng. Black ginseng (BG; Ginseng Radix Nigra) is produced by repeatedly steaming fresh ginseng nine times. The fine roots (hairy roots or fibrous roots) of fresh ginseng that has been steamed nine times are called Fine Black ginseng (FBG). There are more than 30 different ginseng saponins with various physiological and pharmacological activities [2,3]. Ginsenosides are divided into two groups: protopanaxadiols and protopanaxatriols. The root of Panax ginseng reportedly has various biological effects, including anticarcinogenic effects. One study showed that ginseng extracts induce apoptosis and decrease telomerase activity and cyclooxygenase-2 (COX-2) expression in human leukemia cells [4]. In addition, ginseng extracts suppress 1,2-dimethylhydrazine-induced colon carcinogenesis by inhibiting cell proliferation [5].

Until recently, research on anticancer effects of ginseng has focused on ginsenoside Rg3 (Rg3) and ginsenoside Rh2 (Rh2). Ginsenoside Rh3 is not present in raw ginseng or White ginseng, but is synthesized during heating hydrolysis; thus, only a small amount of Rg3 is present in Red ginseng. Ginsenoside Rh3 has an anticancer effect by suppressing phorbol ester-induced COX-2 expression and decreasing activation of nuclear factor kappa-light-chain-enhancer of activated B cells (NF-kB) [6]. Its blood pressure lowering effects have been identified in rat aorta through enhancing endothelium-dependent relaxation [7]. Ginsenoside Rg3 in methanol extraction of heat-processed ginseng has antioxidative and antitumor effects [8].

Ginsenoside Rh2 is a major active anticancer saponin in ginseng extracts [9]. Ginsenoside Rh2 treatment modulates the protein expression level of p21 and cyclin D, and leads to a marked reduction in the proliferation of MCF-7 human breast cancer cells

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It also provokes apoptosis through activating p53 and inducing the proapoptotic regulator Bax in colorectal cancer cells [11]. In addition, Rh2 markedly reduces the viability of breast cancer cells (MCF-7 and MDA-MB-231) by arresting the G1 phase cell cycle via p15 INK4B and p27 KIP1-dependent inhibition of cyclin-dependent kinases [12].

Many studies on BG have been performed because interest in it has increased recently. The main component of BG is reportedly Rg5 (Fig. 1) [13]. Studies demonstrate it has diverse physiological activity such as anti-inflammatory effects on lipopolysaccharide-stimulated BV2 microglial cells [14], protective effects on scopolamine-induced memory deficits in mice [15], and inhibitory effects in a mouse model with oxazolone-induced chronic dermatitis [16]. Rg5 reportedly blocks the cell cycle of SK-HEP-1 cells at the GI/S transition phase by downregulating cyclin E-dependent kinase activity [17].

Breast cancer is a very common cancer in women worldwide. In the United States, it is estimated that breast cancer is the leading cause of all cancers (29%) and the second leading cause of death (14%) [18]. In Korea, 16,015 new cases of breast cancer were reported in 2011 [19]. Anticancer activity of BG extract in the MCF-1 breast cancer cell line exhibited three-fold cytotoxicity, compared with Red ginseng extract [20]. However, ginseng fine roots contain a higher content of ginseng saponin than ginseng main roots [2]. In the present study, we therefore aimed to investigate anti-breast cancer activity (in the MCF-7 cell line) and the action mechanisms of FBG ethanol extract (EE), FBG butanol fraction (BF; primarily containing saponin), and Rg5 as the major saponin.

2. Materials and methods

2.1. Materials

Fine Black ginseng (Panax ginseng Meyer) for experiments was purchased from Kumsan Town, Chungcheongnam Province, the Republic of Korea in August 2009. All other chemicals were of an analytical reagent grade. Distilled water for high-performance liquid chromatography (HPLC) and acetonitrile were purchased from J.T. Baker SOLUSORB (Philipsburg, NJ, USA). The standards were purchased from Chromadex (Santa Ana, CA, USA) and Ambo Institute (Seoul, South Korea). Proton magnetic resonance, carbon magnetic resonance, heteronuclear multiple quantum coherence and heteronuclear multiple bond coherence spectra were measured with INOVA-500 (500 MHz) (Varian, palo alto, CA, USA).

Fig. 1. (A) High-performance liquid chromatography chromatogram of ginsenosides in the butanol (BuOH) fraction of Fine Black ginseng (FBG), compared with the chromatogram of the ginsenoside standards: a, Rg1; b, Re; c, Rf; d, Rh1; e, Rg2; f, Rh2; g, Rc; h, Rb2; i, F1; j, Rd; k, Rg6; l, F4; m, Rk3; n, Rh4; o, (20S) Rg3; p, (20R) Rg3; q, Rk1; r, Rg5. (B) The content of ginsenosides and (C) the composition of ginsenosides in BuOH fraction of Fine Black ginseng.
The mass spectrum was taken on a fast atom bombardment mass spectrometry device (JMS-700; Jeol, Seoul, Korea). For the experiments, Rg3 was purchased from Chromadex. Rosewell Park Memorial Institute (RPMI 1640) medium (Buffalo, NY, USA), fetal bovine serum (FBS), trypsin-EDTA, and penicillin/streptomycin were purchased from Welgene (Daegu, South Korea); and (3,4,5-dimethyl-2-thiazolyl)-2,5-diphenyl-2H-tetrazolium bromide (MTT) and 3,3′-dihexyloxacroxy-cyanine iodide (DiOC6) were purchased from Sigma-Aldrich (St. Louis, MO, USA). The following antibodies were used: poly (ADP-ribose) polymerase (PARP), Bid, purchased from Sigma-Aldrich (St. Louis, MO, USA), and Bcl-2, Bax, and DR4 (Santa Cruz Biotechnology, Santa Cruz, CA, USA); cytochrome C (BD Biosciences, San Jose, CA, USA); and p53, antibodies were used: poly (ADP-ribose) polymerase (PARP), Bid, purchased from Sigma-Aldrich (St. Louis, MO, USA). The following antibodies were used: poly (ADP-ribose) polymerase (PARP), Bid, purchased from Sigma-Aldrich (St. Louis, MO, USA), and Bcl-2, Bax, and DR4 (Santa Cruz Biotechnology, Santa Cruz, CA, USA).

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2.2. Preparation of ethanol extract and butanol fraction

Fine Black ginseng (10 kg) was selected, dried, and powdered. Exactly 2 kg of powdered samples were refluxed two times with 10 L of 95% ethyl alcohol for 2 h in a water bath. The extracts were filtered through filter paper (Nylon membrane filters 7404-004; Whatman, Dassel, Germany) and concentrated by a vacuum evaporator (yield: 18.35%). Ethyl alcohol extract (150 g) was dissolved in 1500 mL of water and extracted with 1500 mL of diethyl ether. The aqueous layer was extracted three times with 1500 mL of water-saturated n-butanol (n-BuOH). The n-BuOH fraction (84.50 g) was evaporated.

2.3. HPLC analysis

The ginsenoside composition of the concentrate was analyzed by HPLC, as suggested by Ko and colleagues [13,21]. The total ginsenoside content and composition of each sample were analyzed three times. The 99% pure ginsenoside standards used in this experiment were purchased from Chromadex and the Ambo Institute. For the experiment, the Waters 1525 binary HPLC system (Waters, Milford, MA, USA) and the Eurospher 100-5 C 18 column (3 × 250 mm; Knauer, Berlin, Germany) were used. The mobile phase was a mixture of acetonitrile (HPLC grade) and distilled water (HPLC grade). The content of acetonitrile was sequentially increased from 17% to 30% (35 min), from 30% to 40% (60 min), from 40% to 60% (100 min), from 60% to 80% (110 min), from 80% to 80% (120 min), from 80% to 100% (125 min), from 100% to 100% (135 min), and finally from 100% back to 17% (140 min, lasting for 5 min). The operating temperature was at room temperature and the flow rate was 0.8 mL/min. The elution profile on the chromatogram was obtained by using a UV/VIS detector at 203 nm (Waters 2487 dual λ absorbance detector; Waters) (Fig. 1A).

2.4. Isolation and identification of ginsenoside Rg5

The n-BuOH fraction (60 g) was chromatographed on a silica gel column (1 kg) with eluting solvents of CHCl3-MeOH-H2O (70:30:4) to obtain six subfractions (F1–F5). The F4 fraction (2.59 g) was further subjected to octadecylsilane (ODS) (C-18) column chromatography (500 g, 60% acetonitrile (ACN)) to provide Rg5 (0.19 g) (Fig. 1B).

In the FAB-MS spectrum of compound I [M-H]− peak appeared at m/z 765. 1H-NMR spectrum (pyridine-d5) showed eight methyl signals (CH3-18, 30, 29, 28, 27, 26, 21) at δ 0.835, 0.979, 1.040, 1.130, 1.313, 1.644, 1.840, H-3 and H-23 signals at δ 3.288, δ 2.791. On the other hand, proton signal resulting from the binding to C-3 glucose, and it was confirmed from δ 3.937 to δ 4.597. And the H-1′ and H-1″ signal from the binding to C-3 glucose was confirmed in δ 4.950 and δ 5.400. 13C-NMR spectrum (pyridine-d5) showed C-1, C-2, C-3, C-4, C-5, C-6, C-7, C-8, C-9, C-10, C-11, C-12, C-13, C-14, C-15, C-16, C-17, C-18, C-19, C-20, C-21, C-22, C-23, C-24, C-25, C-26, C-27, C-28, C-29, C-30, 3-Glc C-1′, 3-Glc C-2′, 3-Glc C-3′, 3-Glc C-4′, 3-Glc C-5′, 3-Glc C-6′, 3-Glc C-1″, 3-Glc C-2″, 3-Glc C-3″, 3-Glc C-4″, 3-Glc C-5″, 3-Glc C-6″ signals at δ 39.76, 28.60, 89.42, 40.75, 76.89, 18.93, 35.84, 40.21, 51.26, 37.51, 32.72, 73.08, 50.94, 51.52, 33.13, 27.25, 51.40, 16.94, 17.09, 140.66, 13.66, 123.82, 27.95, 123.92, 131.74, 26.18, 18.22, 28.60, 16.31, 17.52, 105.62, 83.95, 78.45, 72.15, 78.76, 63.34, 106.55, 77.64, 78.84, 72.12, 78.62, 63.19 (Fig. 2) [22].

2.5. Cell culture and MTT assay

MCF-7 (HER2+/ER−) and MDA-MB-453 (HER2+/ER−) human breast cancer cell lines were maintained using RPMI 1640 medium supplemented with 10% (vol/vol) FBS (Welgene, Daegu, South Korea) plus 100 units/mL penicillin and streptomycin in a 5% carbon dioxide air incubator at 37°C. Cell cytotoxicity was measured by MTT assay. Cells were seeded in 96-well tissue culture plates at the density of 0.2 × 104 cells per well with 100 μL medium, and were allowed to become attached for 24 h. One hundred microliters of the medium with different concentrations of Rg5 (e.g., 0μM, 25μM, 50μM, and 100μM) were added to each well. At indicated times, 30 μL MTT stock solution (3 mg/mL) were added to each well. After culturing the cells at 37°C for 2 h, dimethyl sulfoxide (DMSO) was added to dissolve the formazan crystals. The absorbance was read at the wavelength of 540 nm with a microplate reader (EL800, Biotek Instruments Inc., Winooski, VT, USA).

Fig. 2. (A) The HMQC (500 MHz, in pyridine) and (B) the fast atom bombardment mass spectroscopy spectrum of ginsenoside Rg5.
2.6. Western blot analysis

After treatment, the pellet of cells was rinsed with ice-cold phosphate buffered saline (PBS) and lysed in radio-immunoprecipitation assay buffer (0.1% sodium dodecyl sulfate, 0.5% sodium deoxycholate, 50mM Tris-HCl and 0.1% NP-40, pH 8.0 with 150mM sodium chloride) for 1 h at 4°C. The cell lysate was cleared by centrifugation at 17,000 rpm for 10 min at 4°C. Each supernatant sample was separated by 10% sodium dodecyl sulfate—polyacrylamide gel electrophoresis and the separated protein was transferred to polyvinylidene fluoride (PVDF) membranes. After blocking with 5% nonfat dry milk in TBS-T (25mM Tris and 0.1% Tween 20, 137mM sodium chloride) at room temperature (RT) for 2 h, the membranes were incubated with primary antibodies overnight at 4°C and treated with horseradish peroxidase-conjugated secondary antibodies for 2 h. The signals were detected with the ECL Advance Detection Kit (GE Healthcare Bio-Sciences Corp., Piscataway, NJ, USA) by LAS-3000 luminescent image analysis.

2.7. Annexin V/fluorescein isothiocyanate/propidium iodide assay and analysis of surface expression of DR4 and DR5

Apoptosis was evaluated by annexin V/fluorescein isothiocyanate/propidium iodide (annexin V-FITC/PI) dual staining. Treated
cells were harvested and resuspended in 1 × binding buffer. A combination of annexin V/FITC solution and PI solution were added to each tube. The stained cells were incubated at room temperature for 30 min in the dark. Samples were analyzed by the FACSCanto II Flow Cytometer (BD Biosciences, San Jose, CA, USA).

Cells were analyzed for their cell surface expression of DR4 and DR5 by indirect staining with primary mouse anti-human DR4 and DR5, followed by phycoerythrin (PE)-conjugated goat anti-mouse immunoglobulin G1 (IgG1). The treated cells were harvested and washed with PBS containing 1% bovine serum albumin. Cells were incubated with anti-DR4 or anti-DR5 antibody for 30 min at 4°C in the dark. After incubation, cells were washed twice and reacted with PE-labeled secondary antibody for 30 min at 4°C in the dark. Isotype-matched nonbinding antibodies (Iso) were the negative control cells. Samples were measured by flow cytometry.

2.8. Analysis of cell cycle and microscopic images of DAPI staining

Analysis of the cell cycle was performed by staining with PI. Cells were seeded into a 100-mm dish, which contained 1 × 10⁶ cells per plate. After 24 h, the media were changed to RPMI 1640 medium supplemented with indicated concentrations of Rg5. After 48 h of incubation, the cells were trypsinized and washed with ice-cold PBS, fixed with ice-cold 90% ethanol, and then incubated at −20°C until analysis. For cell cycle analysis, the cells were resuspended in 300 mL of PBS containing 30 μL RNase A solution (10 mg/mL; Sigma-Aldrich) and 1.5 μL PI solution (1 mg/mL; Molecular Probes). After incubation at 37°C for 30 min, cells were determined using the FACSCanto II Flow Cytometer (BD Biosciences, San Jose, CA, USA). The cell cycle distribution was analyzed by FlowJo software (Tree Star, Inc., Ashland, OR, USA).

Fig. 4. The effects of the (A and B) ginsenoside Rg3 and (C and D) ginsenoside Rg5 on cell viability in MDA-MB-453 (HER2+/ER−) and MCF-7 (HER2−/ER+) breast cancer cell lines. Cells were treated with the indicated concentration of ginsenosides for 24 h, 48 h, and 72 h. Cell viability was assessed by 3-(4,5-dimethyl-2-thiazolyl)-2,5-diphenyl-2H-tetrazolium bromide (MTT) assay. * Indicates a significant difference at p < 0.05, versus the control. ** Indicates a significant difference at p < 0.001, versus the control.
Cells were plated at 0.3 × 10^6 cells in six-well plates. After treatment, the cells were fixed in DMSO/methanol (1:4) solution for 12 h at 4°C, stained with 4',6-diamidino-2-phenylindole (DAPI) for 20 min, and observed by fluorescence microscopy.

2.9. Statistical analysis

Statistical significance was performed by Turkey's multiple comparison tests (Sigma Plot version 10.0; Systat Software, San Jose, CA). All experiments were repeated at least three times. Data were analyzed by one-way analysis of variance (ANOVA), and each value was presented as the mean ± the standard deviation.

3. Results

3.1. Isolation and analysis of ginsenoside Rg5

The yield of ginsenosides from ginseng hairy root (i.e., fine root) was higher than the yield from the main root [2], and the saponin content of FBG was higher than that of BG [23]. First of all, the HPLC results showed Rg5 was the main constituent among the ginsenosides in FBG (Fig. 1A). Rg5 was separated from FBG BF using column chromatography (silica gel, ODS) (Figs. 1B, 1C), and the chemical structure was confirmed by spectroscopic methods [e.g., NMR, mass spectroscopy (MS)] (Fig. 2).

3.2. Cytotoxicity of the FBG extract and its main component, ginsenoside Rg5

The effects of FBG EE and FBG BF on cell viability were evaluated in MCF-7 and MDA-MB-453 breast cancer cell lines by MTT assay. The results showed that EE reduced MCF-7 cell viability after 48 h of treatment and it decreased cell viability of MDA-MB-453 cells after 72 h (Figs. 3A, 3B). Increased cell viability was detected in MCF-7 cells when it was treated with 50 μg/ml (at 24 h, 48 h, and 72 h) and 100 μg/ml (24 h) of BF, but at higher concentrations (150 μg/ml and 200 μg/ml) the cell viability was decreased in a dose-dependent manner (Figs. 3C, 3D). As Figs. 4C and 4D show, Rg5, the main component of FBG, exerted significant cytotoxicity in a dose- and time-dependent manner. The cytotoxic effect of 20(S)-Rg3 in MCF-7 cells unexpectedly showed no significant difference. These results were consistent when Rg3 was treated in MDA-MB-453 cells (Figs. 4A, 4B).

3.3. Effects of ginsenoside Rg5 on cell cycle arrest

The results from flow cytometric analysis [i.e., fluorescence-activated cell sorting (FACS)] indicated that Rg5 significantly induced cell cycle arrest (Figs. 5A, 5B). This was further confirmed by the cell cycle assay with the data representing suppressed cell proliferation in MCF-7 cells after Rg5 treatment. Rg5 increased the number of cells in the G0/G1 phase and decreased the number of cells in the S phase. Based on these results, Rg5 may induce cell cycle arrest at the G0/G1 phase. Protein expression of cyclin D1, cyclin E2 and CDK4 was decreased, whereas the expression of p15INK4B, p53 and p21WAF1/CIP1 was increased (Figs. 6A, 6B).

3.4. Effects of ginsenoside Rg5 on apoptosis-related proteins

As Fig. 7A shows, treatment with Rg5 induced caspase-8 and caspase-9, caspase-7, caspase-6. The full-length Bid consequently disappeared in a dose-dependent manner. Poly (ADP-ribose) polymerase (PARP) cleavage was detected in Rg5-treated MCF-7 cells, which indicated that Rg5 reduced cell viability by inducing apoptosis. Promotion of mitochondria-mediated intrinsic apoptotic pathway by Rg5 was evidenced by Bax/Bcl-2 dysregulation,
activation of caspase-9, and release of cytochrome C (Fig. 7A). Apoptosis was evaluated by annexin V/FITC/PI dual staining. After 48 h, Rg5 significantly increased apoptosis at 25 μM and 50 μM and reduced apoptotic cells at 100 μM, whereas necrotic cells were increased (Fig. 7B). The increased expression of DR4 and DR5 on the cell surface was obvious when cells were treated at the 100 μM concentration of Rg5 (Fig. 8A). Activation of p38 mitogen-activated protein kinases (MAPKs) is necessary for apoptosis induced by exposure to ultraviolet radiation, cytokines, chemotherapy, ceramide, and serum deprivation [24]. When cells were treated with Rg5 (50 μM and 100 μM), p38 MAPKs were activated with the generation of reactive oxygen species (data not shown) (Fig. 8C). Survivin, an inhibitor of apoptotic proteins, is highly expressed in most types of cancer and is a regulator of mitosis; survivin-targeting cancer treatment is validated with great efficacy and no serious toxicity [25]. The expression of survivin was suppressed at high

Fig. 6. Effects of ginsenoside Rg5 on G1-related proteins in MCF-7 cells. Cells were treated with DMSO (0.01%) or ginsenoside Rg5 (25, 50 and 100 μM) for 48 h. Protein expressions were determined using western blotting. *p < 0.05, **p < 0.001 versus control are significant different.

Fig. 7. Effects of ginsenoside Rg5 on MCF-7 cell apoptosis. (A) Effects of ginsenoside Rg5 on apoptosis-related proteins in MCF-7 cells. Cells were treated with DMSO (0.01%) or ginsenoside Rg5 (25, 50 and 100 μM) for 48 h. (B) Apoptosis was evaluated by Annexin V-FITC/PI dual staining. The value represents mean ± SD (n = 3). Statistical significance at *p < 0.05, **p < 0.001.
concentrations of Rg5 (Fig. 8D). Apoptotic cells were visualized with DAPI as fluorescent probes. When cells were incubated for 48 h with Rg5 at indicated concentrations (i.e., 0 μM, 50 μM, and 100 μM), the cells displayed the typical apoptosis morphology such as fragmented and condensed nuclei with cellular shrinkage (Fig. 9B). Cells treated with Rg5 at the 100 μM concentration showed a necrosis-like morphology (Fig. 9C).

4. Discussion

Red ginseng is fresh ginseng that is dry-steamed once using water vapor. Black ginseng refers to ginseng that is steamed nine times. Fine Black ginseng refers to the fine roots (i.e., hairy roots) of BG steamed nine times.

As Fig. 1C shows, FBG BF contained more than 40% of the total saponins. The amount of total saponin in the FBG BF was 17 times higher than in BG EE, and was 26 times higher than in RG EE [26]. Fine Black ginseng contained the highest content of Rg5 (9.831%) (Fig. 1C). The amount of Rg5 in FBG BF was 34 times higher than in BG EE, and was 110 times higher than in RG EE [26]. Rg5, the main component of FBG BF, was isolated using column (silica gel, ODS) chromatography, and the chemical structure was confirmed by spectroscopic analysis (i.e., NMR, MS) (Fig. 2).

The difference in chemical structure between Rg5 and Rg3 is the polar hydroxyl group of C-20 in Rg3. When C-20 is induced
dehydrated reaction [43], that is applied to the high-pressure steam, Rg3 is converted to Rk1 and Rg5. Dehydration of the C-20 of the ginsenoside structure increases its bioactivity [27]. Rg5 (i.e., Rg3 that has been dehydrated at C-20) reportedly has cytostatic activity of human hepatoma SK-HEP-1 cells that is approximately four times stronger than that of Rg3 [17].

Therefore, the purpose of this study was to elucidate anti-breast cancer activity of FBG extract and Rg5 in MCF-7 cells. The FBG extract and Rg5 showed significant cytotoxic activity. In previous studies, the BG extract in comparison to RG extract exhibited stronger cytotoxic activity in vitro on the MCF-1 breast cancer cell line, HT-1080 fibrosarcoma cell line and Hepa1C1C7 murine hepato-cell line [20].

The anticancer properties of Rg3 are associated with inducing apoptosis [28], regulating cell cycle [29], blocking angiogenesis [30], and inhibiting proliferation. Rg3 exhibits anticancer activity in various cell lines such as human hepatocellular carcinoma cells (Hep3B) [31], the PC-3M prostate cancer cell line [32], VX2 liver tumors [33], and the U87MG human glioblastoma cell line [28]. However, the cytotoxic effect of 20(S)-Rg3 in MCF-7 cells showed no significant difference, and the results were consistent when MDA-MB-453 cells were treated by Rg3 (Figs. 4A, 4B).

Cell cycle arrest and apoptosis were performed to determine the mechanism of action for the anticancer effects of Rg5. As a result, Rg5 induced significant G0/G1 cell cycle arrest. The results of western blot analysis showed increased Bax (i.e., pro-apoptotic regulator), caspase-6 and caspase-7 (i.e., effector caspases), DR4, and DR5. These results were evident even when Rh2 induced apoptosis in colorectal cancer cells through activation of p53 [34]. The tumor suppressor p53 induces cell self-destruction through the endogenous mitochondrial pathway and exogenous death receptor pathway. This is called p53-dependent apoptosis (i.e., p53-induced apoptosis). In particular, p53-dependent apoptosis is used to induce the expression of proapoptotic members. Bax also is expressed by the activation of p53 [35,36]. When the cells undergo DNA damage, p53 stops the cell cycle through p21 or it induces apoptosis. In response to cellular stress or DNA damage, p53 is stabilized by post-transcriptional modifications and the level of p53 increases [37]. Stabilization and activation of p53 is responsible for cellular anti-proliferative mechanisms such as apoptosis, growth arrest, and cell senescence [38]. This study confirmed the influence of Rg5 on the activity of Bax and p53.

The data showed that the expression of DR4 and DR5 was upregulated by Rg5 in a dose-dependent manner. The tumor nerosis factor-related apoptosis-inducing ligand (TRAIL) is a promising agent for cancer treatment because it selectively induces apoptosis in various cancer cells, but not in normal cells [39]. Many tumor cells are resistant to TRAIL-induced apoptosis. Therefore, it is important to develop combination therapies to overcome this resistance [40]. Rg5 did not increase TRAIL-induced apoptosis, which suggests that Rg5 does not increase the susceptibility of TRAIL-resistant MCF-7 cells. Therefore, Rg5 was unsuitable for combination therapy.

To examine whether Rg5 reduced cell viability via apoptosis, cells were analyzed by using annexin V–FITC/PI staining assay. Rg5 at 0μM, 25μM, and 50μM concentrations increased apoptosis in a dose-dependent manner. However, at 100μM concentration of Rg5, apoptotic cells were reduced, whereas necrotic cells were increased. There are many natural substances similar to this situation. Procyanidin, a polyphenol compound with strong bioactivity and pharmacologic activity, exists widely in grape seeds, hawthorn, and pine bark. Procyanidin induces apoptosis and necrosis of prostate cancer cell line PC-3 in a mitochondrion-dependent manner. With extended procyanidin treatment, the apoptosis rate decreased, whereas the necrosis rate increased. This change was associated with cytotoxic properties that were related to alterations in cell membrane properties [41,42].

Rg5 induces cancer cell apoptosis in a multipath mechanism, and is therefore a promising candidate for antitumor drug development. The antitumor role of Rg5 would be useful in therapeutic approaches (e.g., in combination therapy with other cancer chemotherapy drugs). In this study, we elucidated the effects of Rg5 in MCF-7 and MDA-MB-453 human breast cancer cell lines, which demonstrated that Rg5 may be an effective chemotherapeutic agent for breast cancer. However, further studies are needed to identify the precise mechanism of Rg5. There is also a need for in vivo experiments to confirm the anticancer activity of Rg5.

Conflicts of interest

The authors have no conflicts of interest to declare.

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