Research article

Stereospecific anticancer effects of ginsenoside Rg3 epimers isolated from heat-processed American ginseng on human gastric cancer cell

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A B S T R A C T

Background: Research has been conducted with regard to the development of methods for improving the pharmaceutical effect of ginseng by conversion of ginsenosides, which are the major active components of ginseng, via high temperature or high-pressure processing.

Methods: The present study sought to investigate the anticancer effect of heat-processed American ginseng (HAG) in human gastric cancer AGS cells with a focus on assessing the role of apoptosis as an important mechanistic element in its anticancer actions.

Results and Conclusion: HAG significantly reduced the cancer cell proliferation, and the contents of ginsenosides Rb1 and Re were markedly decreased, whereas the peaks of less-polar ginsenosides [20(S)]-Rg3, Rk1, and Rg5] were newly detected. Based on the activity-guided fractionation of HAG, ginsenoside 20(S)-Rg3 played a key role in inducing apoptosis in human gastric cancer AGS cells, and it was generated mainly from ginsenoside Rb1. Ginsenoside 20(S)-Rg3 induced apoptosis through activation of caspase-3, caspase-8, and caspase-9, as well as regulation of Bcl-2 and Bax expression. Taken together, these findings suggest that heat-processing serves as an increase in the antitumor activity of American ginseng in AGS cells, and ginsenoside 20(S)-Rg3, the active component produced by heat-processing, induces the activation of caspase-3, caspase-8, and caspase-9, which contributes to the apoptotic cell death.

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1. Introduction

Ginseng is a perennial plant belonging to the genus *Panax* and has been reported to exhibit a wide range of pharmacological and physiological actions [1]. American ginseng (AG) is a popular dietary supplement and one of the most commonly used herbal medicines in the USA, which grows as *Panax quinquefolius* L. (Araliaceae) in the USA and Canada. By contrast, *Panax ginseng* Meyer (Araliaceae) has been mainly cultivated in Asia (most notably in Korea and China), and has been used extensively in traditional Chinese medicine [2,3]. Both AG and Asian ginseng extracts have been reported to exhibit free radical scavenging activities, which, from different ginseng species and specific parts, have been thought to be related to their ginsenoside contents [4].

Ginsenosides, which are 30-carbon glycosides derived from the triterpenoid dammarane, as shown in Fig. 1, are regarded as the main active components in AG, as well as Asian ginseng. We previously identified that the structural changes in ginsenosides by heat-processing are closely associated with increased free radical-scavenging activities of AG and Asian ginseng [5,6]. Moreover, we have also recently reported the increased anticancer efficacy of ginsenosides derived from heat-processed Asian ginseng in human gastric cancer cells [7]. Although some of the studies of AG have focused on the antiproliferative activities in cancer cells [8], little is known about the effect of heat processing on the anticancer effect and mechanism of AG on gastric cancer.

Gastric cancer is the second most common cause of cancer-related deaths in the world and is estimated that over 738,000
people die from it every year [9]. Multiple therapeutic strategies are used across the world for the management of operable gastric cancer patients [10]. Various multimodality approaches using chemotherapy, radiation, or a combination of both have been evaluated in an attempt to improve the outcomes of postsurgery. Although there have been advances in the treatment of early gastric cancer, outcomes still remain poor with the majority of patients eventually dying from disease relapse [10].

The present study sought to investigate the anticancer effect of heat-processed AG in human gastric cancer cells with a focus on the possible effects of ginsenosides on caspase-dependent apoptosis, AGS cells were also pretreated with 20 μM Z-VAD-fmk for 2 hours prior to ginsenosides treatment.

2.2. Preparation of ginseng extracts

American ginseng extract was purchased from Sigma-Aldrich (St Louis, MO, USA). Heat-processed AG (HAG) was made by steaming AG extract at 120°C and 0.11 MPa for 3 h, and drying at 50°C for 3 d. Heat-processing condition was chosen according to the literature [11]. HAG extract (30 g) was suspended in water and the water-soluble polysaccharide fraction was separated by Diaion HP 20 (Mitsubishi Chemical, Tokyo, Japan) column chromatography using water as an eluting solution, followed by elution with methanol [12]. Each solution was evaporated in vacuo to give the water eluate (27 g) and methanol eluate (3 g).

2.3. Analysis and structural confirmation of ginsenosides

Analytical reversed-phase high performance liquid chromatography (HPLC) system was composed of a solvent degasser (G1322A; Agilent, Palo Alto, CA, USA), binary pump (G1329B; Agilent), and model 380 Evaporative Light Scattering Detector (ELSD; Agilent). ELSD conditions were optimized in order to achieve maximum sensitivity: temperature of the nebulizer was set for 50°C, and N2 was used as the nebulizing gas at a pressure of 2.0 bar. The Phenomenex Luna C18 column (150 mm × 4.6 mm, 5 μm, Torrance, CA, USA) was used, and the mobile phase consisted of a binary gradient of solvent A (acetonitrile:water:5% acetic acid in water = 15:80:5) and solvent B (acetonitrile:water = 80:20) at a flow rate of 1.0 mL/min. The gradient flow program was as follows: initial; 0% B, 6 min; 30% B, 18 min; 50% B, 30 min; 100% B, 37 min; 100% B, 42 min; 0% B. The amounts of ginsenosides in samples were quantified as reported previously [5]. The standard conditions containing 1–50 μg of each ginsenoside were injected into the HPLC and all calibration curves showed good linearity (R² > 0.995). The analysis was repeated twice for the verification of repeatability.

2.4. Cell culture

The human gastric cancer AGS cell line was purchased from the American Type Culture Collection (Manassas, VA, USA). The cells were grown in RPMI1640 medium (Cellgro, Manassas, VA, USA) supplemented with 10% fetal bovine serum (Gibco BRL, Carlsbad, MD, USA), 100 units/mL penicillin, and 100 μg/mL streptomycin and incubated at 37°C in a humidified atmosphere with 5% CO2.

2.5. Analysis of cell viability

AGS cells were treated with different concentrations of compounds for 24 h, and cell proliferation was measured using the Cell Counting Kit-8 (CCK-8; Dojindo Laboratories, Kumamoto, Japan) according to the manufacturer’s recommendations. Control cells were exposed to culture media containing 0.5% (v/v) DMSO. Paclitaxel was used as a positive control (data not shown). In order to examine the possible effects of ginsenosides on caspase-dependent apoptosis, AGS cells were also pretreated with 20 μM, 40 μM, and 60 μM Z-VAD-fmk for 2 hours prior to ginsenosides treatment.

2.6. Western blotting analysis

AGS cells were grown in 6-well plates and treated with the indicated concentration of compounds for 24 h. Whole-cell extracts were then prepared according to the manufacturer’s instructions using RIPA buffer (Cell Signaling Technology, Inc.) supplemented with 1 × protease inhibitor cocktail and 1 mM phenylmethylsulfonyl fluoride. Proteins (whole-cell extracts, 30 μg/lane) were separated by electrophoresis in a precast 4–15% Mini-
PROTEAN TGX gel (Bio-Rad, Hercules, CA, USA) blotted onto PVDF transfer membranes and analyzed with epitope-specific primary and secondary antibodies. Bound antibodies were visualized using ECL Advance Western Blotting Detection Reagents (GE Healthcare, Amersham, Buckinghamshire, UK) and a LAS 4000 imaging system (Fujifilm, Tokyo, Japan).

2.7. Statistical analysis

Statistical significance was determined through analysis of variance (ANOVA) followed by a multiple comparison test with a Bonferroni adjustment. A p-value of <0.05 was considered statistically significant. The analysis was performed using SPSS version 19.0 (SPSS Inc., Chicago, IL, USA).

3. Results and discussion

Many bioactive dietary agents are used alone or as adjuncts to existing chemotherapy to improve efficacy and reduce drug-induced toxicity [13]. For example, epidemiological, as well as experimental studies have shown that diets rich in vegetables and fruit are chemotherapeutically beneficial, exerting the activity to inhibit proliferation and induce apoptosis against malignancies, including gastric cancer [14–16]. In the present study, we sought to identify the anticancer effect and mechanism of HAG to examine its therapeutic potential.

The HPLC chromatogram of AG extract is illustrated in Fig. 2. AG include the typical ginsenosides Re, Rg1, Rb1, Rc, Rb2, and Rd (Fig. 2A). After heat processing at 120 °C, the ginsenosides Rb1 and Re were markedly decreased, whereas the peaks of less polar ginsenosides (20(S,R)-Rg3, Rk1, and Rg5) were newly detected at about 23 min and 27 min (Fig. 2B, Table 1). Therefore, ginsenosides Rb1 and Re were more efficiently deglycosylated and transformed into less polar ginsenosides during heat processing than other ginsenosides contained in AG.

Fig. 3A shows the morphological changes of human gastric cancer AGS cell treated with AG or HAG. The AGS cell line has been shown to grow in athymic mice and to have the same histochemical and cytological characteristics as the specimen taken from the patient [15], and recently, this cell line has been widely used as a model system for evaluating cancer cell apoptosis [17,18]. As a result, HAG was found to induce apoptotic body formation stronger than AG (Fig. 3A), and HAG significantly suppressed AGS cell proliferation from a lower concentration, whereas AG did not suppress cell proliferation at any concentration (Fig. 3B). Therefore, ginsenosides Rb1 and Re were more efficiently deglycosylated and transformed into less polar ginsenosides during heat processing than other ginsenosides contained in AG.

We next examined the antiproliferating efficacies of ginsenosides Rb1 and Re with or without heat-processing, because the amounts of these ginsenosides were decreased markedly after heat-processing in AG than in other ginsenosides (Fig. 2A). Both ginsenosides Re and Rb1 showed no efficacy in cancer cell proliferation (Fig. 4) without heat-processing. However, heat processed-Rb1 significantly suppressed cell proliferation from a lower concentration (Fig. 4A), and this result was similarly observed in the case of methanol eluate (Fig. 3D). By contrast, ginsenoside Re showed a weak efficacy even at a high concentration of 100 μg/mL (Fig. 4B). Therefore, the major active component of HAG was considered to be related to heat-processed ginsenoside Rb1. From the HPLC analysis of ginsenoside Rb1, prior to and after heat-processing, ginsenoside Rb1 was transformed into ginsenosides 20(S,R)-Rg3, Rk1, and Rg5 after heat-processing at 120 °C (Fig. 4C and D) as shown in AG (Fig. 2). We previously reported that ginsenoside Re gradually transformed into less polar ginsenosides Rg2, Rg6, and F4 upon heat-processing [7]. Considering these, less polar ginsenosides 20(S,R)-Rg3, Rk1, and Rg5 in HAG were generated mainly from ginsenoside Rb1 by heat processing at 120 °C.

We next examined the antiproliferative effects of 20(S,R)-Rg3 or Rk1/Rg5 mixtures, which were collected using preparative HPLC. As shown in Fig. 5A, 5B, 20(S,R)-Rg3 reduced cancer cell viability stronger than Rk1/Rg5 mixture, and each IC50 value were 23.6 μg/mL and 42.9 μg/mL, respectively. Interestingly, the efficacy of 20(S,R)-Rg3 was similar with that of the methanol eluate, as well as of heat-processed Rb1 (Figs. 3D and 4A). To further confirm the main active component, anticancer effects of 20(S)-Rg3 and 20(R)-Rg3 were individually examined. Subsequently, ginsenoside 20(S)-Rg3 was obviously identified as the main active component of HAG.
while there was no effect in ginsenoside 20(R)-Rg3 (Fig. 5C and D). Thus, anticancer efficacy of HAG was thought to be mainly related to ginsenoside 20(S)-Rg3, which was transformed from ginsenoside Rb1 during heat processing.

Apoptosis is recognized as an essential mechanism of physiological cell death, and caspases play pivotal roles in cell apoptosis. In line with this notion, we investigated whether ginsenoside 20(S)-Rg3-induced cell death is involved in apoptosis. A Western blot analysis was first used to evaluate the expression of proteins involved in the apoptotic response to determine if apoptosis occurs via the intrinsic or extrinsic pathway (Fig. 6A–C). Exposure to ginsenoside 20(S)-Rg3 for 24 h induced the cleavage of PARP, as well as that of caspase-3, caspase-8, and caspase-9, in a dose-dependent manner. In addition, ginsenoside 20(S)-Rg3 significantly triggered the downregulation of Bcl-2 and upregulation of Bax in a dose-dependent manner. Next, we examined the effect of the pan-caspase inhibitor Z-VAD-fmk on cell proliferation to confirm the role played by caspases in ginsenoside 20(S)-Rg3-induced apoptosis. As shown in Fig. 6D, pretreatment with 60 μM Z-VAD-fmk abrogated apoptotic cell death induced by the ginsenoside 20(S)-Rg3, although the recovery was weak at the high concentration of 50 μg/mL. These findings demonstrate that ginsenoside 20(S)-Rg3 induces the activation of caspase-3, caspase-8, and caspase-9, which contributes to apoptotic cell death.

Ginsenosides 20(S)-Rg3 and 20(R)-Rg3 are epimers of each other depending on the position of the hydroxyl group (OH) on carbon-20 (Fig. 1), and this epimerization is known to be produced by the selective attack of the OH group after the elimination of...
Fig. 4. Changes in effects of ginsenosides Rb1 and Re upon heat-processing on AGS cell proliferation. (A) Cells were treated with ginsenoside Rb1 or heat-processed ginsenoside Rb1 at different concentrations (12.5 μg/mL, 25 μg/mL, 50 μg/mL, and 100 μg/mL) for 24 h. (B) Cells were treated with ginsenoside Re or heat-processed ginsenoside Re at different concentrations (12.5 μg/mL, 25 μg/mL, 50 μg/mL, and 100 μg/mL) for 24 h. Relative cell proliferation was measured by the CCK-8 assay. Each value represents the mean ± standard deviation of three independent experiments. (C) HPLC chromatogram of ginsenoside Rb1 prior to heat-processing. (D) HPLC chromatogram of ginsenoside Rb1 after heat-processing. *p < 0.05 compared with vehicle control. CCK-8, Cell Counting Kit-8; HPLC, high performance liquid chromatography.

Fig. 5. Anticancer effects of ginsenosides on AGS cell proliferation. (A) Cells were treated with ginsenoside 20(S,R)-Rg3 at different concentrations (12.5 μg/mL, 25 μg/mL, 50 μg/mL, and 100 μg/mL) for 24 h. (B) Cells were treated with ginsenoside Rg5/Rk1 at different concentrations (12.5 μg/mL, 25 μg/mL, 50 μg/mL, and 100 μg/mL) for 24 h. (C) Cells were treated with ginsenoside 20(S)-Rg3 at different concentrations (6.25 μg/mL, 12.5 μg/mL, 25 μg/mL, 50 μg/mL, and 100 μg/mL) for 24 h. (D) Cells were treated with ginsenoside 20(R)-Rg3 at different concentrations (6.25 μg/mL, 12.5 μg/mL, 25 μg/mL, 50 μg/mL, and 100 μg/mL) for 24 h. Relative cell proliferation was measured by the CCK-8 assay. Each value represents the mean ± standard deviation of three independent experiments. *p < 0.05 compared with the vehicle control. CCK-8, Cell Counting Kit-8.
Counting Kit-8; PARP, poly(ADP-ribose) polymerase; Z-VAD-fmk, Benzyloxycarbonyl-
for 24 h. Relative cell proliferation was measured by the CCK-8 assay. Each value
actin. (D) Cells were pretreated with the different concentrations (20
m
(85 kDa) in AGS cells treated with ginsenoside 20(S)-Rg3. 
37 kDa), cleaved caspase-3 (17 kDa, 19 kDa), PARP (116 kDa), and cleaved PARP
results for (A) extrinsic and (B) intrinsic apoptosis in AGS cells showing the levels of
paradoxical effect of ginsenoside 20(S)-Rg3. Therefore, stereospecificity in the medicinal
efficacy of these ginsenosides has been reported by several researchers. The OH group of ginsenoside 20(S)-Rg3
channel regulation [21]. Ginsenoside 20(S)-Rg3 serves as a major component through activation of caspase-3,
caspase-8, and caspase-9 in the event.

Conflicts of interest

The authors declare no conflict of interest.

Acknowledgments

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