Abstract. Glycyrrhetinic acid (GA) is a natural compound extracted from liquorice, which is often used in traditional Chinese medicine. The purpose of the present study was to investigate the antitumor effect of GA in human non-small cell lung cancer (NSCLC), and its underlying mechanisms in vitro. We have shown that GA suppressed the proliferation of A549 and NCI-H460 cells. Flow cytometric analysis showed that GA arrested cell cycle in G0/G1 phase without inducing apoptosis. Western blot analysis indicated that GA mediated G1-phase cell cycle arrest by upregulation of cyclin-dependent kinase inhibitors (CKIs) (p18, p16, p27 and p21) and inhibition of cyclins (cyclin-D1, -D3 and -E) and cyclin-dependent kinases (CDKs) (CDK4, 6 and 2). GA also maintained pRb phosphorylation status, and inhibited E2F transcription factor 1 (E2F-1) in both cell lines. GA upregulated the unfolded proteins, Bip, PERK and eIF2α. Accumulation of unfolded proteins in the endoplasmic reticulum (ER) triggered the unfolded protein response (UPR) pathway, which decreases protein synthesis and induces cell cycle arrest in the G1 phase. This study provides experimental evidence to support the development of GA as a chemotherapeutic agent for NSCLC.

Introduction

Lung cancer (LC) is an important etiology of malignant mortality worldwide with global statistics indicating >1,000,000 deaths each year (1). Non-small cell lung cancer (NSCLC) accounts for 80-85% of LC (2). The 5-year survival rate is ~6.6% for advanced stage LC (stage III or IV) in the US (3). Apart from surgery, adjuvant chemotherapy with gefitinib, erlotinib, and epidermal growth factor receptor tyrosine kinase inhibitors (EGFR-TKIs), has been widely used to clinically treat NSCLC. However, their efficacy is limited because of natural or acquired resistance (4). Therefore, there is a need to identify and develop potential anticancer drugs with increased selectivity and reduced toxicity.

Glycyrrhetinic acid (GA) is a bioactive component of glycyrrhiza (GL), which is often used in Chinese traditional medicine to treat various diseases. GA is known to possess anti-inflammatory, anti-viral and cytokine-inducing activity (5-10). Recently, the antitumor activity of GA has been extensively studied. GA has been reported to have cytotoxic effects against human ovarian cancer, hepatocellular carcinoma, breast cancer, pituitary adenoma and human bladder cancer (11-16). However, no inhibitory activity on the growth of NSCLC cell lines has been reported.

Endoplasmic reticulum (ER) stress responses are mediated by the activation of several unfolded protein response (UPR)-signaling pathways. In mammalian cells, the UPR signals increase expression of ER chaperone proteins GRP78/Bip, GRP94, and CHOP (17). The UPR coordinates the induction of ER chaperones, which decreases protein synthesis and results in growth arrest in G1 phase of the cell cycle. Previous studies have demonstrated that ER stress triggers G1-phase cell cycle arrest in various cancer cells (18). However, the molecular mechanism underlying UPR-induced G1 cell cycle arrest remains largely unknown.

In this study, we investigated the effect of GA on survival and proliferation of human NSCLC cell lines (A549 and NCI-H460), and found that GA could suppress the proliferation of both cell lines, with A549 being more sensitive than...
NCI-H460. GA arrested cells in G1 phase via inactivation of CDK4/6-cyclin-D1/D3 complex through p18/p16 activation, and inactivation of CDK2-cyclin-E2 complex through p27/p21 activation. This resulted in P RB dephosphorylation and inactivation of E2F transcription factor 1 (E2F-1) in both cell types. E2F-1 is an essential transcription factor that regulates cell cycle progression and apoptosis. Additionally, GA was found to increase the expression of Bip, protein kinase-like ER kinase (PERK) and ERP72, which are linked to ER stress.

Materials and methods

Reagents. GA was purchased from Nanjing Zelang Medical Technology Co., Ltd. (Jiangsu, China), and dissolved in dimethyl sulfoxide (DMSO) (Sigma, St. Louis, MO, USA) to make a stock solution before use. For treatment of cells, it was diluted in culture medium at the appropriate concentrations, and the final concentration of DMSO was <0.01% (v/v). Cisplatin (Lot no. H20030675; Nanjing Pharmaceutical Factory Co., Ltd., Jiangsu, China), and insulin, propidium iodide (PI), 3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyltetrazolium bromide (MTT), and alamarBlue were from Sigma. Alexa Fluor 488 Annexin V/Dead Cell Apoptosis kit was from Invitrogen Life Technologies (Carlsbad, CA, USA). Antibodies against caspase-3, -7 and -9, p18, p16, p27, p21, cyclin-D1, -D3 and -E2, CDK6, 4 and 2, E2F-1, P RB, Bip, PERK, ERP72, β-actin, and HRP-conjugated antibodies (anti-rabbit or mouse immunoglobulin G) were obtained from Cell Signaling Technology, Inc. (Danvers, MA, USA).

BCA protein estimation kit was from Sigma. Nitrocellulose (NC) blotting membrane was from Pall Corporation (DF Mexico, Mexico). Enhanced chemiluminescence (ECL) was from Bio-Rad (Hercules, CA, USA).

Cell culture. Human NSCLC cell lines A549 and NCI-H460 were purchased from the Type Culture Collection of the Chinese Academy of Sciences (Shanghai, China). A549 cells were cultured in DMEM/F12 (Gibco-BRL, Carlsbad, CA, USA), supplemented with 10% fetal bovine serum (FBS). NCI-H460 cells were grown in RPMI-1640 medium (Gibco-BRL), supplemented with 10% FBS. All cells were cultured under 5% CO2 at 37°C.

In vitro viability assay. The effect of GA on cell viability was measured using the MTT assay. Cells were seeded in 96-well plates at 5x10^3 cells/well in 100 µl of culture medium, and treated with drug the next day for 24, 48, and 72 h. The final concentrations of GA used in the assays were 50, 25, 12.5, 6.25 and 3.125 µmol/l in triplicate, respectively. Treated cells were incubated with 20 µl of MTT (5 mg/ml) for 4 h at 37°C in the dark. Optical density of producer after incubation was measured using a microplate reader (Bio-Rad) at a wavelength of 490 nm.

Cell cycle analysis. After treatment with various concentrations of GA for different time, the cells were harvested with trypsin, washed once with PBS, and then fixed in 70% ethanol overnight at 4°C. Before flow cytometry analysis, the cells were then treated with 1 mg/ml of RNase for 30 min at 37°C, and then stained with 40 µg/ml of PI for 30 min. A total of 1x10^4 cells/sample were analyzed using a FACSCalibur flow cytometer (BD Biosciences, Heidelberg, Germany). Data were evaluated using ModFit software.

Western blot analysis. After treatment with different concentrations of GA, the cells were lysed in RIPA buffer containing 50 mM Tris/HCl (pH 8.0), 150 mM NaCl, 1% (w/v) Nonidet P-40, 1% (w/v) sodium deoxycholate, 0.1% (w/v) SDS, 0.1 mM DTT, 0.05 mM PMSF, 0.002 mg/ml aprotinin, 0.002 mg/ml leupeptin, and 1 mM NaVO3. The protein concentrations of the supernatants were determined by the BCA Protein Assay kit. Equal amounts of the protein were loaded and separated by 10 or 12% SDS-PAGE, and then transferred onto NC membranes. The membranes were incubated overnight...
Figure 2. Glycyrrhetinic acid (GA) induces G1-phase cell cycle arrest in non-small cell lung cancer (NSCLC) cells without induction of apoptosis. Effect of GA on cell cycle was investigated using propidium iodide (PI) staining. Cells were treated with 0-50 µmol/l of GA for 24, 48 and 72 h, and then stained with PI. Green peak represents G0/G1 phase, red peak represents S phase and blue peak represents G2 phase, respectively. Upper panel shows representative of three independent experiments with similar results, and lower panel represents the bar graphs of cells in different phases. Bar graph represents mean ± SD from three independent experiments. (A) Cells were strikingly accumulated in the G1 phase after treatment with GA for 24, 48 and 72 h. (B) Representative bar graph for A549 and (C) for NCI-H460 cells. (D) Western blot analysis of caspase-3, -7 and -9 protein expression after treatment with GA in a time (24 h) and dose (0, 12.5, 25, 50 µmol/l)-dependent manner. β-actin was used as a loading control. (E) Caspase-3, -7 and -9 activity in A549 and NCI-H460 cells treated with GA in a time (24 h) and dose (0, 12.5, 25, 50 µmol/l)-dependent manner. Results shown are the mean of three independent experiments; error bars represent SD. GA-mediated G1-phase arrest is dependent on regulatory cyclin-dependent kinase inhibitors (CKIs) p18, p16, p27, p21, and GA decreases the levels of G1-phase regulatory CDKs and cyclins in both cell lines.
with primary antibodies against caspase-3, -7 and -9, p18, p16, p27, p21, cyclin-D1, -D3 and -E2, CDK6, 4 and 2, E2F-1, pRb, Bip, PERK, ERPT2, or β-actin at 4°C, and then incubated with HRP-conjugated secondary antibodies (anti-rabbit or mouse immunoglobulin G) for 1 h at room temperature. Immunoreactivity was detected by ECL (Bio-Rad). Immunoblot experiments were repeated three times. Quantitative analysis was performed using Image Lab™ Software (Bio-Rad).

Statistical analysis. All values were expressed as mean ± SD (n=3). One-way analysis of variance (ANOVA) was used to determine statistical significance, followed by post hoc multiple comparisons (Dunn's test) using SPSS 19.0. P<0.05 was considered to be statistically significant.

Results

GA suppresses the proliferation of NSCLC cells in vitro. To determine the suppression effect of GA (structure shown in Fig. 1A) on NSCLC cells, we performed a cell viability assay using A549 and NCI-H460 cell lines, respectively. After treatment for 24, 48 or 72 h, the viability of the two cell lines significantly decreased in a dose- and time-dependent manner (Fig. 1B and C).

GA arrests the cell cycle in G0/G1 phase. The results of flow cytometric analysis showed that the percentage of G0/G1 phase of both of A549 and H460 increased after treated with different concentrations of GA for 24, 48 and 72 h (Fig. 2A), respectively. Cell cycle distribution analysis showed that GA prevents the cell cycle progression by arresting the cells in the G0/G1 phase in both cell lines. In A549 cells, percentage of cells in G0/G1 phase increased from 52.03±1.42% (control group) to 71.63±6.61% for cells treated with 25 µmol/l of GA for 24 h. At 48 h, percentage of cells in G0/G1 phase increased from 54.90±5.90% (control group) to 83.00±1.41% for cells treated with 25 µmol/l of GA. While at 72 h, percentage of cells in G0/G1 phase increased from 69.70±5.38% (control group) to 81.83±2.58% for cells treated with 25 µmol/l of GA (Fig. 2A, upper panel). In NCI-H460, percentage of cells in the G0/G1 phase increased from 66.10±0.99% (control group) to 80.95±1.91% for cells treated with 25 µmol/l of GA. At 48 h, percentage of cells in G0/G1 phase increased from 84.60±2.40% (control group) to 88.95±2.19% for cells treated with 25 µmol/l of GA. While at 72 h, percentage of cells in the G0/G1 phase increased from 85.00±0.85% (control group) to 91.00±2.26% for cells treated with 25 µmol/l of GA (Fig. 2B, upper panel). No increase in S or G2/M peak was observed in either cell line.

Taken together, our data strongly suggested that GA did not induce apoptosis but caused cell cycle arrest in G0/G1 phase in NSCLC cells. Annexin assay did not show any significant changes in apoptotic/necrotic cell population for all concentrations of GA as compared to the control group in either cell line at 24, 48 and 72 h, respectively. To further validate the above data, we checked the expression levels of caspase-3, -7 and -9 in both cell lines after 24 h treatment with GA by western blot analysis. Expression of caspase-3 decreased with increase in drug concentration in NCI-H460 cells, but no significant changes in caspase-7 and -9 protein levels or activity were observed in A549 or NCI-H460 cells (Fig. 2D and E), suggesting that GA did not hinder the viability of cells.

GA downregulates the levels of cell cycle regulatory proteins and retinoblastoma (Rb) phosphorylation. To investigate the causes of cell cycle arrest, cyclin-dependent kinase inhibitors (CKIs), such as p27, p21, p18 and p16 that regulate G0/G1 phase of cell cycle progression were examined by western blot analysis (19-21). In A549 cells, the levels of p27, p21, p18 and p16 were significantly increased after 24 h treatment with GA as compared to the control cells (Fig. 3C). H460 cells also showed similar results (Fig. 3D). To further dissect the biochemical events controlling the transition of cell cycle phases, we examined the levels of several proteins, such as cyclin-D1, -D3 and -E2, CDK4, 6 and 2, which are involved in G0/G1-phase progression, and found that GA significantly decreased the expression of these proteins in both cell lines (Fig. 3E and F). GA also significantly decreased the expression levels of E2F-1, the essential transcription factor that regulates cell cycle progression and apoptosis, and pRb (Fig. 3G and H).

GA upregulated the levels of ER stress regulatory proteins. Previous studies have demonstrated that ER stress triggers G1-phase cell cycle arrest in various cancer cells (22). Therefore, we examined whether ER stress was induced by GA. Bip is the master regulator of ER function. Phosphorylation of double-stranded RNA-activated PERK is closely associated with Bip. Hence, we checked the expression of Bip, PERK and ERPT2 in both cell lines by western blot analysis, and found that the expression levels of these proteins were significantly upregulated after 24 h treatment with GA (Fig. 4A and B).

Discussion

GA is a natural active compound that is extracted from the Chinese herbal medicine *glycyrrhiza*. GA was shown to induce cell cycle arrest in G1 phase (13,15). The role of GA in NSCLC, especially its relationship with ER stress has not been reported. In the present study, we found that GA induced G0/G1 arrest in A549 and NCI-H460 cell lines, which provides a useful model system to characterize the cytotoxic effects of therapeutic agents. Furthermore, GA could have therapeutic potential in the treatment of NSCLC.

Our results have shown that GA successfully inhibited proliferation of two NSCLC cell lines, A549 and NCI-H460. Cell cycle analysis by flow cytometry showed that GA induced a modest increase in G0/G1 phase in both cell lines. However, based on expression levels of caspase-3, -7 and -9 by western blot analysis, GA did not induce apoptosis in either cell line. It is well known that eukaryotic cell cycle is regulated by the coordinated activity of protein kinase complexes, each consisting of a cyclin-dependent kinase (CDK) and cyclins. CDK complexes are formed and activated at specific stages of the cell cycle, and their activities are required for progression through distinct cell cycle phases (23). Progression through G1 phase requires the activities of cyclin-D-dependent CDK4 or 6, followed by activation of the cyclin-E- and cyclin-A-dependent kinase CDK2. The cyclin-CDK complex
formed during G1-phase catalyses phosphorylation of the dominant inhibitor of G1/S-cell cycle progression, the Rb family of tumor suppressor proteins, thereby blocking their inhibitory activity allowing the cell to progress into S phase (24-27). It is also known that these cyclin-CDK complexes often bind to CKIs including p16, p18, p21 and p27, which inhibit their
kinase activities, and prevent cell cycle progression (28). E2F-1 is an essential transcription factor that regulates cell cycle progression and cell proliferation. E2F-1 activity is regulated by the Rb protein that binds activator E2F proteins to inhibit transcription outside of G1/S in animals (29).

Flow cytometric analysis of A549 and NCI-H460 cells treated with GA showed that GA inhibits cell cycle progression by blocking the transition from G1 to S phase. To further investigate this result, western blot analysis was used to examine proteins associated with the cell cycle, e.g., cyclin-D1/D3, which is expressed in G1 phase and binds to CDK4 and 6 to activate them, followed by activation of the cyclin-E-dependent kinase CDK2. These protein kinase complexes were inhibited by GA. GA also significantly decreased the expression levels of E2F-1 and pRb in both cell lines. Our results indicated that GA induced growth inhibition mainly via regulation of p16, p18, p21 and p27 status in A549 and NCI-H460 cells.

In the present study, the analysis of DNA content versus light scatter of the GA-treated A549 and NCI-H460 cells indicated no apoptosis. Similarly, the expression of caspase-3, -7 and -9 measured by western blot analysis indicated that GA could not induce apoptosis in these cells. However, GA induced expression of ER proteins GRP78/Bip, PERK and ERP72, which are associated with ER stress. This result suggested that GA inhibited proliferation of A549 and NCI-H460 cells and caused G0/G1-phase cell cycle arrest via ER stress rather than apoptosis.

GRP78/Bip is a major cellular target of the UPR, an ER chaperone that not only binds to unfolded proteins but also regulates the activation of ER stress transducers such as IRE1, PERK, and ATF6 (30-32). GRP78/Bip is ubiquitously expressed at very low levels in growing cells, but it is highly expressed in response to numerous cellular stresses. ERP72, a member of the protein disulfide isomerases (PDI) family, is localized in the ER, and plays a major role in quality control and folding (33). Dysregulation of ER chaperone/folding enzymes ERP72 and GRP78/Bip occurred early after ablation of PERK function suggesting that changes in ER secretory functions could reduce insulin gene expression and cell proliferation (34,35).

Previous studies have found that CKIs and cyclins play important roles in ER stress and cell cycle arrest. p27 was reported to be a critical mediator of ER stress-induced G1 cell cycle arrest in melanoma cells (36). p21 integrates the DNA damage response with ER stress signaling, which

Figure 4. Glycyrrhetinic acid (GA) induces endoplasmic reticulum (ER) stress. (A) A549 and NCI-H460 cells treated with various concentrations of GA (12.5, 25 and 50 µmol/l) for 24 h were analyzed by immunoblotting with antibodies against Bip, protein kinase-like ER kinase (PERK) and ERP72. β-actin was used as a loading control. (B) Densitometry analysis of Bip, PERK and ERP72 levels relative to β-actin were performed using three independent experiments. Error bars represent SD; *p<0.01, #p<0.05.
then regulates mitochondrial death pathways during chronic genotoxic stress (37). Translational regulation of cyclin-D1 in response to ER stress is a mechanism for checkpoint control that prevents cell cycle progression (17). PERK has been shown to mediate cell cycle arrest by blocking cyclin-D1 translation during UPR (17,38). Similarly, our study has shown that induction of members of the INK4 (p16, p18) or Kip/Cip (p21, p27) families of cell cycle kinase inhibitors causes ER stress, and accumulation of unfolded proteins in the ER triggers UPR, which is a stress signaling pathway. The UPR coordinates the induction of ER chaperones, decreases protein synthesis, and causes growth arrest in G1 phase of the cell cycle.

We have convincingly shown that GA inhibits proliferation of NSCLC cell lines by causing cell cycle arrest in G0/G1 phase in a time- and dose-dependent manner, without inducing apoptosis. We have elucidated a new mechanism of action of GA against NSCLC by inducing G1-phase cell cycle arrest through ER stress pathway. Since GA synergizes the effect of anticancer drugs, it provides new insight into the therapeutic index of NSCLC treatment.

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