Inhibitory effect of baicalein combined with gemcitabine in human pancreatic cancer cell lines

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Received November 30, 2016; Accepted November 2, 2017

DOI: 10.3892/ol.2018.8043

Abstract. Pancreatic cancer is an aggressive disease with a particularly poor prognosis contributing to a substantial percentage of cancer-associated mortality rates. In the present study, the combination treatment of baicalein (BAI) and gemcitabine (GEM) was investigated to examine whether it inhibited the growth of the human CFPAC-1 pancreatic cancer cell line \textit{in vitro} and \textit{in vivo}. The cytotoxic interactions between BAI and GEM in human pancreatic cancer cell lines were determined using MTT assays, and the effect of the two agents on apoptosis was detected using Hoechst 33258 staining and annexin V/7-AAD. The protein levels of Bcl-2-associated X protein (Bax), B-cell lymphoma 2 (Bcl-2), caspase-3, poly ADP ribose polymerase (PARP) and survivin were detected using western blot analysis. Furthermore, the expression levels of Bax, Bcl-2, caspase-3 and survivin in tumor tissues were detected using immunohistochemistry. The results demonstrated that following GEM treatment, the growth of CFPAC-1 cells and xenografts in nude mice were inhibited, and the expression levels of Bcl-2 and survivin were downregulated, whilst the expression levels of Bax, caspase-3 and PARP were upregulated. These effects were enhanced with the use of BAI in combination with GEM. The mechanism underlying the anti-tumor effect of BAI combined with GEM may be associated with the induction of cell apoptosis and the inhibition of proliferation. To the best of our knowledge, this is the first evidence of the efficacy of BAI against pancreatic cancer and may provide the potential clinical evidence for the use of this drug combination for the treatment of patients with pancreatic cancer.

Introduction

Pancreatic cancer is a highly malignant disease, which has the fourth highest mortality rate with a 1-year survival rate of ~29% globally in 2012, in addition to the fact that the prognosis of this disease is considerably poor (1). Surgery is a potentially curative option for pancreatic cancer, however a majority of patients present symptoms when the tumor has reached an inoperable stage (2). Chemotherapy and/or radiotherapy are the standard treatments for patients with inoperable pancreatic cancer. The standard first line drug gemcitabine (GEM), which is a deoxycytidine analogue, is used to treat patients with advanced pancreatic cancer; however, its efficacy remains low with 5-year survival rates remaining at 20-25% in 2005 (3,4).

Therefore, there has been considerable interest in combining GEM with other systemic agents in order to improve patient outcomes.

Traditional Chinese herbs may represent a useful resource for drug development (5). Among them, baikaline (BAI) is a purified flavonoid extracted from the root of \textit{Scutellaria baicalensis} that possesses a wide variety of biological activities, including anti-inflammatory, anti-infection, anti-microbial, neuro-protective and anti-oxidant effects (6,7). BAI has exhibited anti-tumor effects on multiple different types of cancer by inhibiting proliferation and inducing cancer cell apoptosis, including in liver cancer, prostate cancer, lung cancer, breast cancer, bladder cancer and cervical cancer (8-13). The antitumor activity of BAI is mediated by activation of Bcl-2-associated X protein (Bax), B-cell lymphoma 2 (Bcl-2), caspase-3 and poly ADP-ribose polymerase (PARP) (14). Therefore, it is important to evaluate the beneficial effect of BAI as a single treatment or when administered in combination with other conventional anticancer agents in pancreatic cancer.

In the present study, the effects of BAI on pancreatic cancer cells were examined. It was revealed that BAI enhanced the effects of GEM treatment in the inhibition of CFPAC-1 cell viability \textit{in vitro} by inducing cell apoptosis. \textit{In vivo}, combination treatment resulted in synergistically reduced tumor volume and weight in a xenograft mouse model. Therefore, the present study identified that combined BAI and GEM treatment may be a promising potential treatment for pancreatic cancer.
Materials and methods

Cell culture. Human CFPAC-1 pancreatic ductal adenocarcinoma and PANC-1 pancreatic cancer cell lines were purchased from the Shanghai Cell Bank of the Chinese Academy of Sciences (Shanghai, China). The 2 cell lines were cultured in DMEM (Invitrogen; Thermo Fisher Scientific, Inc., Waltham, MA, USA), supplemented with 10% fetal bovine serum (Invitrogen; Thermo Fisher Scientific, Inc.), 100 U/ml penicillin and 100 µg/ml streptomycin in a 37°C humidified atmosphere containing 5% CO2/95% air.

MTT assay. CFPAC-1 or PANC-1 cells were cultured in 96-well plates at 1x10⁵ cells per well. After 24 h, the cells were treated with multiple concentration of BAI alone (0.1, 0.2, 0.4, 0.8, 1.6, 3.2, 6.25, 12.5, 25, 50 and 100 µM), GEM alone (0.03, 0.06, 0.125, 0.25, 0.5 and 1 µg/ml) or combination treatment (0.03, 0.06, 0.125, 0.25, 0.5 and 1 µg/ml GEM with either 1.6 or 6.25 µg BAI) for 48 h, the control group was treated with PBS (pH 7.4). Subsequently, 50 µl MTT (1 µg/ml; Sigma-Aldrich; Merck KGaA, Darmstadt, Germany) was added to the cell media and incubated for 4 h at 37°C. MTT was discarded and 150 µl DMSO was loaded in each well. The spectrophotometric absorbance of the samples was measured using a microplate reader (Bio-Rad Model 680; Bio-Rad Laboratories, Inc., Hercules, CA, USA) at 570 nm with a reference wavelength of 655 nm. The percentage of cell survival was calculated using the following formula: Cell survival=absorbance value of infected cells/absorbance value of uninfected control cells. Six reduplicate wells were measured at each concentration and every experiment was performed at ≥3 times.

Combination index (CI) analysis. CI was used to evaluate the pharmacological interactions of different combinations of BAI and GEM (15). Briefly, synergism, additivity or antagonism in the BAI and GEM combinations was calculated on the basis of the multiple drug effect equation and quantitated by CI, where CI=1 indicates that the two drugs have additive effects, CI<1 indicates synergism effects and CI>1 indicates antagonism effects. The CI was calculated based on: CI=(Dx1/((Dx1)+(Dx2)+(Dx1)(Dx2))/((Dx1)(Dx2)/(Dx1)+(Dx2))), where (Dx) is the dose of the drug, inhibiting 'x%'.

Hoechst 33258 staining. Following GEM (0.125 µg/ml), BAI (1.60 or 6.25 µM) or combination treatment, the control group was treated with PBS, the cells were fixed and stained using Hoechst 33258 for 10 min at room temperature and observed using a fluorescence microscope (Olympus Corporation, Tokyo, Japan) at x20 magnification. A total of 6 fields per treatment were assessed. The stained cells were identified as apoptotic if they were highly condensed with brightly stained nuclei, or non-apoptotic if they were stained pale blue.

Apoptosis assay by flow cytometry. CFPAC-1 or PANC-1 cells treated with 0.125 µg/ml GEM or 1.6/6.25 µM BAI alone or a combination treatment were incubated for 48 h at 37°C, the control group was treated with vehicle solution, and then cells were collected and washed twice with cold PBS and subsequently incubated with Annexin V-FITC/7-ADD double staining solution at room temperature for 15 min. Then, the stained cells were analyzed using flow cytometry (BD Biosciences, San Jose, CA, USA). The cell apoptotic rates were analyzed using ModFIT software (version 3.2; Verity Software House, Inc., Topsham, ME, USA). Apoptotic events were recorded as a combination of Annexin V+/7-AAD (early apoptotic) and Annexin V+/7-AAD+ (late apoptotic/dead) events. This experiment was conducted 3 times.

Western blot analysis. Whole cell lysates from CFPAC-1 cells treated with vehicle control or drugs for 48 h, then were washed using PBS once and lysed using RIPA lysis buffer (Beiyotime Institute of Biotechnology, Haimen, China), containing protease inhibitor cocktail (Roche Diagnostics, Basel, Switzerland) and phosphatase inhibitor cocktail (Roche Diagnostics), and sonicated for 15 min on ice. Protein concentration was measured using a BCA protein assay (Bio-Rad Laboratories, Inc.). A total of 30 µg samples were separated by SDS-PAGE on a 10% gel and transferred to polyvinylidene difluoride membranes using the Bio-Rad electrotransfer system (Bio-Rad Laboratories, Inc.). The membrane was blocked using 5% fat-free milk for 1 h at room temperature and then incubated with appropriate primary antibody diluted in 3% BSA solution at 4°C overnight. For immunodetection, the following antibodies were used for our analysis: anti-Bax (cat no. 5023), -Bcl-2 (cat no. 15071), -Caspase-3 (cat no. 9662), -PARP (cat no. 9532) and -Survivin (cat no. 2808) antibodies (1:1,000; Cell Signaling Technology, Inc., Danvers, MA, USA), followed by incubation with DyLight dye-conjugated secondary antibodies (cat nos. 35502 and 35568; 1:10,000; Thermo Fisher Scientific, Inc.) for 1 h at room temperature. Blots were scanned using the Odyssey Western Detection system (LI-COR Biosciences, Lincoln, NE, USA), followed by quantification using ImageStudio software (version 5.2.5; LI-COR Biosciences). Anti-β-actin (cat no. sc-77778; Santa Cruz Biotechnology, Inc.; 1:1,000 dilution) was used as a loading control.

In vivo model. A total of 24 female BALB/c nude mice (4-5 week-old, 15-16 g) were purchased from the Shanghai Laboratory Animal Center (Shanghai, China). Mice were allowed free access to sterilized water and food and were housed in individual ventilated cages at 23±5°C under a 12-h light/dark cycle. CFPAC-1 cells were resuspended in serum-free DMEM at a concentration of 1x10⁶ cells/ml. A total volume of 0.2 ml cell suspension (total 2x10⁶ cells) was then injected subcutaneously into the right anterior armpit of nude mice to establish a xenograft model. The 24 injected mice were randomly divided into 4 groups: control, BAI, GEM and combination treated groups (n=6 per group). Control group was treated with vehicle solution. The mice were sacrificed at day 28 or when the tumor grew to a maximum of 1,500 µM³. The Animal Care and Use Committee of Nanjing University of Chinese Medicine ethically approved the experimental protocol.

Immunohistochemical assay. Tumor xenografts were fixed in formalin solution overnight at room temperature and embedded in paraffin blocks, which were sliced into 4-µm thick sections for immunohistochemical staining. Following
deparaffinization, the sections were incubated at 4°C overnight with antibodies of Bcl-2 (cat no. 15071), Bax (cat no. 5023), caspase-3 (cat no. 9662) and survivin (cat no. 2808) (1:500; Cell Signaling Technology, Inc.). The sections were washed 3 times with PBS with 0.1% Tween-20 (PBST), and incubated for 1 h at room temperature with a horseradish peroxidase-conjugated goat anti-mouse immunoglobulin G antibody (1:200; cat no. sc-2031; Santa Cruz Biotechnology, Inc.). The slices were washed with PBST 3 times, and incubated in diaminobenzidine (Sangon Biotech Co., Ltd., Shanghai, China) for 5 min at room temperature and counterstained with hematoxylin (Sangon Biotech Co., Ltd.) for 1 min at room temperature. Images were captured using a light microscope (Olympus Corporation) at a magnification of x20. Five sections per tumor tissue were examined.

**Statistical analysis.** All experiments were repeated at ≥3 times and all data are presented as the mean ± the standard error of the mean. Graph Pad Prism 6 (GraphPad Software Inc., La Jolla, CA, USA) was use to perform statistical analysis. Comparisons among multiple groups were determined using a one-way or two-way analysis of variance test (two-way analysis of variance was used for tumor volume analysis) followed by Dunnett’s or Bonferroni's post hoc test. P<0.05 was considered to indicate a statistically significant difference.

**Results**

**Synergistic cytotoxicity of the combined treatment of GEM and BAI in pancreatic cancer cell lines.** To test the potential of the cytotoxicity of BAI, the human pancreatic cancer cell line CFPAC-1 was incubated with a range concentrations (0.1-100 μM) of BAI for 48 h and then the number of cells in each group was determined. Treatment with higher concentrations of BAI (3.2-100.0 μM) significantly inhibited the viability of CFPAC-1 cells compared with the control group (P<0.001; Fig. 1A). For the cells treated with 0.125 µg/ml GEM, the number of viable cells decreased by 32% compared with the control group (Fig. 1B). Treatment with BAI in addition to GEM or GEM alone revealed that the viability of CFPAC-1 was inhibited in a concentration-dependent manner. Subsequently, to assess the synergistic effect of GEM in combination with BAI on cell proliferation, CFPAC-1 cells were incubated with a range of concentrations of GEM, in combination with 1.6 or 6.25 µg BAI for 48 h. MTT assays revealed that combined treatment of either concentration of BAI with 0.125 µg/ml GEM was significantly more cytotoxic in CFPAC-1 cells compared with the same concentration of GEM alone (CI (0.125 µg/ml GEM and 1.6 µM BAI)=0.181; CI (0.125 µg/ml GEM and 6.25 µM BAI)=0.465; Fig. 1B). A similar effect was identified in the PANC-1 cell line, where 0.125 µg/ml GEM in combination with 6.25 µM BAI significantly decreased the cell viability compared with 0.125 µg/ml GEM treatment alone (Fig. IC).

**BAI and GEM cooperatively induce apoptosis in pancreatic cancer cells.** To determine if GEM and BAI treatment induces apoptosis, Hoechst 33258 staining and flow cytometric analysis were used. As indicated in Fig. 2, apoptosis analysis revealed that treatment with GEM or BAI alone induced minimal apoptosis; however, the combined treatment of GEM with 1.6 or 6.25 µM BAI resulted in a significant increase of the number of apoptotic cells was observed in CFPAC-1 and PANC-1 cell
**Figure 2.** BAI augments apoptosis induced by GEM in human CFPAC-1 and PANC-1 pancreatic cancer cell lines. Morphological analysis of nuclear chromatin was performed using Hoechst 33258 staining in CFPAC-1 cells treated with (A) 1.6 µM or (B) 6.25 µM BAI, in addition to 0.125 µg/ml GEM alone or in combination for 48 h. (C) PANC-1 cells were treated with 6.25 µM BAI and 0.125 µg/ml GEM alone or in combination for 48 h using Hoechst 33258 staining. Early and late apoptotic events in the cells were determined using Annexin V/7-AAD staining and flow cytometry staining in CFPAC-1 cells treated with (D) 1.6 µM or (E) 6.25 µM BAI, in addition to 0.125 µg/ml GEM alone or in combination for 48 h. (F) PANC-1 cells were treated with 6.25 µM BAI and 0.125 µg/ml GEM alone or in combination for 48 h using Annexin V/7-AAD staining. Images were captured using a light microscope at a magnification of x20. **P<0.001 and ***P<0.0001 with comparisons indicated by lines. GEM, gemcitabine; BAI, baicalein.

**Figure 3.** Effects of BAI alone, GEM alone or combined treatment on the protein expression levels of Bax, Bcl-2, caspase-3, PARP and survivin in the human CFPAC-1 pancreatic cancer cell line. (A) Representative images of Bax, Bcl-2, caspase-3, PARP and survivin protein expression, generated using a western blot analysis. β-actin was used as a loading control. Quantification of the intensity of (B) Bax, (C) Bcl-2, (D) caspase-3, (E) PARP and (F) survivin. **P<0.01 and ***P<0.001 with comparisons indicated by lines. GEM, gemcitabine; BAI, baicalein; Bax, Bcl-2 associated protein X; Bcl-2, B-cell lymphoma 2; PARP, poly ADP ribose polymerase.
lines compared with the control (P<0.0001; Fig. 2). In addition, there was a significant increase in apoptosis in cells treated with GEM in combination with 6.25 µM BAI compared with GEM alone in CFPAC-1 and PANC-1 cell lines (P<0.001; Fig. 2). Furthermore, western blot analysis demonstrated that the expression levels of Bax, caspase-3 and PARP were significantly increased, and Bcl-2 and survivin were significantly decreased, in the combination BAI (1.6 or 6.25 µM) and GEM (0.125 µg/ml) treated group compared with the control group (P<0.001; Fig. 3), and compared with the BAI-alone treated group (P<0.01; Fig. 3). These results provide evidence that the synergistic cytotoxic effects of GEM and BAI against pancreatic cancer cells may be attributed to the induction of apoptosis.

**Growth inhibitory effect of BAI and GEM in a mouse model.** In the CFPAC-1 xenograft nude mouse model, all mice survived the duration of the experiment with no observable toxic effect. Mice were sacrificed after 28 days and the weights and volume of the xenografts were measured. Compared with the control group, xenografts from the mice treated with BAI or GEM therapy were decreased in volume and weight (Fig. 4A and B). Furthermore, the combination therapy consisting of 6.25 µM BAI and 0.125 µg/ml GEM induced a significant decrease in tumor volume and weight compared with the control and with BAI treatment alone (P<0.0001; Fig. 4C and D). Immunohistochemical assays additionally revealed that the protein levels of Bcl-2, survivin, Bax and caspase-3 followed a similar pattern to that of the in vitro experiment, since Bcl-2 and survivin were significantly decreased and Bax and caspase-3 were significantly increased in the combined treatment group compared with the control group (P<0.0001; Fig. 4E-H). Furthermore, survivin, Bax and caspase-3 levels were significantly altered in the combination therapy group compared with the GEM-alone treated group (P<0.01; Fig. 4E-H).
Discussion

In the present study, the effect of BAI alone and in combination with GEM on the apoptosis and cell viability of the human CFPAC-1 pancreatic ductal adenocarcinoma cell line was investigated in vitro and in vivo. In addition to this, the effects of the combination treatment of BAI and GEM in the Panc-1 cell line in vitro was examined. The mechanism underlying the induction of apoptosis in cancer cells by BAI may be associated with the altered expression levels of pro-apoptotic and anti-apoptotic molecules including Bcl-2, Bax, survivin, PARP and caspase-3. Furthermore, the combination treatment of BAI with GEM significantly reduced the tumor weight and volume relative to either treatment as a monotherapy in the CFPAC-1 xenograft mouse model. In future studies, the growth inhibitory effect of BAI and GEM in a Panc-1 xenograft nude mouse model will be examined. Therefore, the present study investigated for the first time, to the best of our knowledge, the in vitro and in vivo effects of BAI in pancreatic cell lines and in a xenograft model.

BAI, the major bioactive flavone extracted from S. baicalensis, is more potent than baicalin, which is hydrolyzed by microflora to BAI (16). So far BAI, at doses that are toxic to malignant cells, have been demonstrated to exert no or little toxicity on normal cells (17). Additionally, BAI has been revealed to inhibit the proliferation of multiple human cancer cell lines (8,9). The anticancer effect of BAI is the induction of apoptotic cell death and the mechanisms include cyclin-dependent kinase modulation, the activation of PARP, and caspase-3, caspase-7 and cytochrome c release (18,19). These results indicate that BAI possesses therapeutic potential against cancer.

Apoptosis is an important process for homeostasis by maintaining the balance between cell death and cell survival. The ratio of pro- and anti-apoptotic molecules regulates cell death. Bax is a member of the Bcl-2 protein family associated with apoptosis (20). Bax mediates the permeabilization of the outer mitochondrial membrane and the release of cytochrome c into the cytoplasm, which activates caspase-3 to induce chromosomal cleavage and apoptosis (21). Survivin belongs to the inhibitor of apoptosis protein family and is considered a node protein, inhibiting apoptosis and regulating cell mitosis (22). Promising cancer treatment strategies that target apoptotic inhibitors, including Bcl-2 family proteins and inhibitor of apoptosis proteins are presently under investigation (23).

To conclude, in the present study, BAI was revealed to inhibit the proliferation of pancreatic cancer cells due to the induction of apoptosis in vitro and in vivo. Furthermore, evidence was provided that the efficacy of BAI was significantly improved when administered in combination with GEM. Therefore, the present study may provide an alternative strategy for the treatment of pancreatic cancer.

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