Indomethacin induces apoptosis in the EC109 esophageal cancer cell line by releasing second mitochondria-derived activator of caspase and activating caspase-3

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Abstract. The use of non-steroidal anti-inflammatory drugs (NSAIDs) has been associated with a reduced risk of various types of cancer, including esophageal cancer. However, the mechanisms underlying the antineoplastic effects of NSAIDs in esophageal cancer remain to be elucidated. In the present study, a significant inhibition in cell viability was observed in the EC109 cells following treatment with different concentrations of indomethacin, and these effects occurred in a dose- and time-dependent manner. This inhibition was due to the release of second mitochondria-derived activator of caspase (Smac) into the cytosol and the activation of caspase-3. Subsequently, flow cytometry was performed to investigate indomethacin-induced apoptosis following the overexpression or knockdown of Smac, and western blot analysis was performed to determine the expression of Smac and the activation of caspase-3. Overexpression of Smac was promoted apoptosis, while downregulation of Smac significantly inhibited apoptosis. Western blot analysis demonstrated that indomethacin induced apoptosis through releasing Smac into the cytosol and activating caspase-3. These results indicated that Smac is essential for the apoptosis induced by indomethacin in esophageal cancer cells.

Introduction

The second mitochondria-derived activator of caspase (Smac) protein is a novel protein, which is involved in the mitochondrial regulation of apoptosis (1,2). It has been observed that the expression of Smac is reduced in various types of malignant tissue compared with their normal tissue counterparts, which suggests that it may function as a novel tumor suppressor protein (3,4). Several previous studies have demonstrated that Smac promotes apoptosis by binding to the inhibitor of apoptosis proteins (IAPs), leading to release of caspases from the IAPs and to the activation of the caspase cascade (5-7).

Previous epidemiological, preclinical and clinical studies have suggested that indomethacin and other non-steroidal anti-inflammatory drugs (NSAIDs) may possess anticancer activities (8). A study by Kohli et al (9) observed that indomethacin promoted apoptosis in the HCT116 colon cancer cell line and this effect was attenuated by knockout of the Smac gene, which suggested that Smac is important in the apoptosis induced by NSAIDs. In addition, a study by Bank et al (10) demonstrated that Smac sensitizes NSAID-induced apoptosis by promoting the activation of caspase-3 and the release of cytochrome c. This result further elucidated the mechanism by which Smac enhances NSAID-induced apoptosis in colon carcinoma cells.

Esophageal cancer is a common type of human upper gastrointestinal carcinoma and, in 2008, a total of 400,000 individuals succumbed to mortality, worldwide and 480,000 new cases were diagnosed (11). Previous clinical trials have indicated that NSAIDs are able to significantly reduce the risk of esophageal cancer (12,13). The effects of indomethacin on the growth, apoptosis and expression of Smac in esophageal cancer cells remain to be fully elucidated. In the present study, indomethacin-mediated apoptosis was investigated in EC109 esophageal cancer cells by overexpression or knockdown of Smac, to examine the mechanism by which NSAIDs control the growth and survival of esophageal carcinoma cells.
Materials and methods

Cell culture and cell transfection. The human EC109 esophageal cancer cells were obtained from the American Type Culture Collection (Manassas, VA, USA) and maintained at 37°C and 5% CO₂ in RPMI 1640 medium (HyClone Laboratories, Inc., Logan, UT, USA) supplemented with 10% fetal bovine serum (GE Healthcare Life Sciences, Logan, UT, USA), 100 U/ml penicillin and 100 g/ml streptomycin (Sigma-Aldrich, St. Louis, MO, USA). The EC109 cells were transfected with Lipofectamine™ 2000 (Invitrogen Life Technologies, Carlsbad, CA, USA), according to the manufacturer’s instructions, during the logarithmic growth phase. Subsequent experiments were performed 48 h after transfection. The pcDNA3.1-Smac vector was constructed in our previous work (14). In brief, Smac was amplified from the mRNA of human testis tissue by reverse transcription polymerase chain reaction (RT and PCR kits, Takara Bio, Inc., Dalian, China; S1000 Thermal cyclers, Bio-Rad Laboratories, Inc., Hercules, CA, USA) using the following primers: Forward 5’-CGGGATCCCAATGCGGCTCTGTA-3’ and reverse 5’-CCCCAAGCTTGGCCCTCAATCCTACGC-3’. Then, it was digested by BamHI and HindIII (Takara Bio, Inc.) and inserted into pcDNA3.1 (Invitrogen Life Technologies) by T4 DNA ligase (Invitrogen Life Technologies). The small interference RNA (siRNA) and the control fragment for Smac RNAi were designed and synthesized by Shanghai GenePharma Co., Ltd. (Shanghai, China).

MTT assay. Cell viability was assessed using a methyl-thiazol tetrazolium (MTT) assay. Exponentially growing EC109 cells were seeded into 96-well plates at a density of 1x10⁴ cells/well and incubated for 24 h at 37°C. Different concentrations (0, 50, 100, 200, 400 and 800 µM) of indomethacin (Sigma-Aldrich) were then added to the wells and incubated at 37°C for 24, 48 and 72 h. A total of 10 µl sterile MTT (5 µg/ml; Sigma-Aldrich) was added to each well. Following a further incubation at 37°C for 4 h, the reaction was stopped by the addition of 150 µl dimethyl sulfoxide. After 10 min, the formazan production was determined by measurement of the spectrometric absorbance at a wavelength of 490 nm using an enzyme immunoassay analyzer (xMark™ Microplate Absorbance spectrophotometer; Bio-Rad Laboratories, Inc.).

Flow cytometry assay. The cells were plated into 24-well plates at a density of 1x10⁵ cells/well and transfected with either the pcDNA3.1/pcDNA3.1-Smac or siRNA/control vectors using Lipofectamine™ 2000. After 48 h, indomethacin was added to a final concentration of 100 µM (overexpression group) and 400 µM (RNA interference group). The following day, the cells were trypsinized (Sigma-Aldrich) and incubated with 5 µl propidium iodide (Invitrogen Life Technologies) and 5 µl annexin V-fluorescein isothiocyanate (FITC; Invitrogen Life Technologies) for 15 min. Samples were then analyzed for apoptosis using a FACScan flow cytometer (BD Biosciences, Franklin Lakes, NJ, USA).

Western blotting. Following indomethacin treatment, the EC109 cells were transfected into precooled lysis buffer [50 mM Tris-Cl (pH 7.5), 150 mM NaCl, 1 mM EDTA (pH 8.0), 1% Triton X-100 and protease inhibitor cocktail; Roche Molecular Biochemicals, Indianapolis, IN, USA] and incubated for 30 min on ice. Proteins from different samples (30 µg protein/sample) were separated on 12% SDS-PAGE gels (Invitrogen Life Technologies) and transferred to polyvinylidene fluoride membranes (EMD Millipore, Billerica, MA, USA) by blotting. Subsequent to treatment with 5% bovine serum albumin (Shanghai Bioleaf Biotech Co., Ltd, Shanghai, China) at room temperature (25°C) for 1 h, the membranes were incubated with the primary antibody overnight at 4°C, incubated with horse-radish peroxidase-labeled goat anti-rabbit immunoglobulin G antibody (1:5,000, sc-45101; Santa Cruz Biotechnology, Inc., Dallas, TX, USA) for 1 h at 25°C, and developed using electrochemiluminescence detection (ChemiDoc MP System; Bio-Rad Laboratories, Inc.). For the western blot analysis, the following primary antibodies were used: Rabbit monoclonal anti-Smac (1:1,000; 1012-1; Epitomics, Burlingame, CA, USA), rabbit polyclonal anti-Tubulin (1:1,000; 10068-1-AP; ProteinTech Group, Inc., Chicago, IL, USA), rabbit polyclonal anti-GAPDH (1:5,000; 10494-1-AP; ProteinTech Group, Inc.), rabbit monoclonal anti-cytoschrome c oxidase subunit IV (COX IV; 1:1,000; 4850; Cell Signaling Technology, Inc., Danvers, MA, USA), rabbit polyclonal anti-caspase-3 (1:1,000; ab2302; Abcam, Cambridge, MA, USA) and rabbit monoclonal anti-pro-caspase-3 (1:1,000; ab32499; Abcam). Tubulin, COX IV and GAPDH were used as loading controls.

Results

Indomethacin reduces the viability of the EC109 cells. In order to examine the effect of indomethacin on the EC109 cells, an MTT assay was used initially to investigate the viability of EC109 cells treated with a range of concentrations of indomethacin (0, 50, 100, 200, 400 and 800 µM) for 24, 48 and 72 h. Indomethacin treatment resulted in a gradual reduction of the viability of the EC109 cells as the exposure duration and drug concentration increased, which indicated that the inhibitory effect of indomethacin on the EC109 cell survival was dose- and time-dependent (Fig. 1). No significant change was observed in the viability of the EC109 cells following indomethacin treatment, for 24 h while prolonged exposure of the drug for 72 h resulted in significant cell death. Treatment with indomethacin for 48 h resulted in a range of EC109 cell viabilities at different concentrations, between 73.58% (50 µM) and 6.57% (800 µM), with a linear correlation between the drug concentration and cell viability (Fig. 1).

Indomethacin treatment releases Smac from mitochondria into the cytosol and activated caspase-3. The EC109 cells were treated with 200 µM indomethacin for 0, 24 or 48 h and the expression levels of Smac in the mitochondria and cytosol were analyzed by western blotting. In the absence of indomethacin, Smac was localized to the mitochondria, however, following indomethacin stimulation, Smac was released into the cytosol. The increase in cytosolic levels
of Smac correlated with the activation of caspase-3 during treatment of EC109 cells with indomethacin (Fig. 2). In addition, the release of Smac and the activation of caspase-3 were prominent following treatment of indomethacin for 48 h. Therefore, a treatment duration of 48 h was selected for the subsequent experiments.

**Smac knockdown inhibits indomethacin-induced apoptosis.**

To further detect the function of Smac in the EC109 cells treated with indomethacin, siRNA of Smac or its control segment was transfected into the EC109 cells for 48 h. Western blotting confirmed that the expression of Smac in the Smac-knockdown cells was reduced compared with the control cells (Fig. 3A). The Smac-knockdown EC109 cells and control cells were then treated with different concentrations of indomethacin for 48 h and the cell viability was assessed by MTT assay. The results identified that the viability of the Smac-knockdown cells was significantly higher compared with the control cells following treatment with 100, 200 or 400 µM indomethacin (P<0.05, Fig. 3B).

Following treatment with 400 µM indomethacin for 48 h, the Smac-knockdown and control cells were stained with propidium iodide and annexin V-FITC to assess the induction of apoptosis by flow cytometry. In the indomethacin-treated cultures, the number of apoptotic EC109 cells in the Smac-knockdown group was significantly lower compared with the control group (P<0.05, Fig. 4A and B). Furthermore, western blotting demonstrated that the expression of Smac in the cytosol and the activity of caspase-3 were reduced in the Smac-knockdown cells treated with indomethacin, which indicated that knockdown of Smac inhibited the activation of caspase-3, resulting in a reduction in indomethacin-induced apoptosis (Fig. 4C).

**Overexpression of Smac enhances indomethacin-induced apoptosis.**

The EC109 cells were transfected with either pcDNA3.1 or pcDNA3.1-Smac for 48 h and western blotting confirmed that the overexpression of Smac was successful (Fig. 5A). The cell viability was detected by MTT following treatment with different concentrations of indomethacin. Overexpression of Smac was found to reduce the viability of EC109 following treatment with 50, 100, 200 or 400 µM indomethacin (P<0.05, Fig. 5B). Furthermore, flow cytometric analysis indicated that the apoptotic rates of the Smac-overexpressed EC109 cells were significantly higher compared with the control group following treatment with
Figure 4. Smac-knockdown inhibits indomethacin-induced apoptosis. (A) EC109 cells were either untreated or exposed for 48 h to 400 µM indomethacin following transfection with Smac or control siRNA for 48 h. Annexin V/PI staining was used for the detection of apoptosis. (B) Apoptotic rates are presented as the mean ± standard deviation (*P<0.05, vs. control). (C) Expression levels of Smac in the cytosol and caspase-3 were detected by western blot analysis. Smac, second mitochondria-derived activator of caspase; COX IV, cytochrome c oxidase subunit IV; PI, propidium iodide; siRNA, short interference RNA.

Figure 5. Overexpression of Smac promoted the reduction of cell viability by indomethacin. (A) Smac was overexpressed in the EC109 cells. Protein expression was evaluated by western blotting in the EC109 cells following transfection with the pcDNA3.1-Smac or pcDNA3.1 for 48 h. (B) Cell viability was detected by MTT assay following treatment of the Smac overexpressed EC109 cells with different concentrations of indomethacin for 48 h. (*P<0.05, vs. control). Smac, second mitochondria-derived activator of caspase.
100 µM indomethacin (P<0.05, Fig. 6A and B). Finally, western blotting demonstrated that the increased apoptotic rate in the Smac-overexpressing EC109 cells was due to increased levels of cytosolic Smac and caspase-3 activation (Fig. 6C).

Discussion

NSAIDs are a group of drugs with antipyretic, analgesic, anti-inflammatory and anti-rheumatic properties (15). In addition, previous epidemiological surveys and preliminary clinical studies have indicated that the long-term use of NSAIDs may reduce the risk of colon, stomach, breast, prostate and lung cancer and other types of malignant tumor (16-20). Several studies have demonstrated that NSAIDs may inhibit tumor growth by inhibiting the expression and activity of cyclooxygenase (COX), inducing apoptosis and suppressing angiogenesis (21,22). Kohli et al (9) reported that the NSAID-induced apoptosis in colon cancer cells was dependent on the pro-apoptotic factor, Smac, as knockout or RNA interference of the Smac gene in colon cancer cells significantly reduced the apoptosis induced by NSAIDs. Increasing evidence suggests that the use of NSAIDs also significantly reduces the risk of esophageal cancer, although the mechanism by which NSAIDs inhibit esophageal cancer growth remain to be elucidated (23,24).
Indomethacin is a commonly used NSAID. It has been observed to induce the apoptosis of lung, head and neck cancer cells (25,26). In the present study, different concentrations of indomethacin were used to treat the EC109 cells. The cell viability and cell growth were significantly inhibited by indomethacin in a dose- and time-dependent manner. Subsequently, western blot analysis revealed that Smac was released into the cytosol and caspase-3 was activated in the EC109 cells treated with indomethacin. Thus, it was hypothesized that indomethacin induced apoptosis in the EC109 cells by activating the Smac-dependent apoptotic pathway. A previous study demonstrated that the release of Smac and cytochrome c occurred during sulindac-induced apoptosis (10). Our previous study described a similar function of Smac in cisplatin-induced apoptosis, in which Smac led to the activation of caspase-3 and -9 in lung cancer cells (27).

Smac is an important protein in the mitochondrial apoptotic pathway. Previous studies have revealed a reduction or loss in the expression of Smac in various tumor tissues compared with their normal tissue counterparts, suggesting that Smac functions as a tumor suppressor gene and its normally low level of expression is associated with tumorigenesis (1,3,4). In addition, Smac also inhibits tumor growth by promoting apoptosis induced by the tumor necrosis factor-related apoptosis-inducing ligand, enhancing antitumor immune response, inhibiting the cell cycle and sensitizing tumor cells to radiotherapy or chemotherapy (28,29). Reduced expression levels of Smac have been observed in esophageal tumor cell lines and in tumor specimens from patients with esophageal cancer (30). A lower levels of Smac expression has been associated with a reduced sensitivity to chemotherapy in the treatment of esophageal cancer (30). It has also been reported that the overexpression of Smac sensitizes esophageal cancer cells to cisplatin chemotherapy (31).

In the present study, the expression of Smac was increased and decreased in order to further investigate the role of Smac in the indomethacin-induced apoptosis of EC109 cells. The results revealed that knockdown of Smac inhibited the indomethacin-induced apoptosis in the EC109 cells, which were transfected with Smac siRNA. Western blot analysis demonstrated that indomethacin treatment of Smac-knockdown EC109 cells reduced the cytosolic expression of Smac and the activation of caspase-3. These results suggested that the translocation of Smac from the mitochondria to the cytosol was important in promoting the activation of caspase-3 and the indomethacin-induced apoptosis. To further investigate the effects of indomethacin-induced apoptosis, the apoptotic rates and activation of caspase-3 were measured following overexpression of Smac in the EC109 cells. The results were consistent with those of the Smac-knockdown experiments, indicating that indomethacin-induced apoptosis is dependent on and regulated by Smac and caspase-3. These observations regarding the role of Smac in indomethacin-induced apoptosis in EC109 cells provides a novel treatment strategy for esophageal cancer.

At present, Smac mimetics, which are composed of the last four-eight N-terminal residues of Smac, have been successfully applied in cancer treatment in the laboratory, and are being assessed in early clinical studies (32,33). Smac mimetics have also been combined with several chemotherapeutic drugs to enhance the efficacy of chemotherapy and to overcome drug resistance (34-36). However, at present, few studies have been performed on the combined use of Smac peptides and NSAIDs. NSAIDs are inexpensive and relatively non-toxic compared with conventional chemotherapeutic drugs (37). Whether the combination of NSAIDs and Smac peptides is more effective compared with either alone in inducing apoptosis in esophageal cancer cells, with or without low dose chemotherapeutic agents, remains to be elucidated and is of interest.

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References


