



# Determining the effect of ellagic acid on the proliferation and migration of pancreatic cancer cell lines

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**Background:** Ellagic acid is a natural dietary compound found in several berries and fruits. It inhibits fibrosis, inflammation and carcinogenesis. We tried to find out what role ellagic acid plays in pancreatic cancer to see if it can be used as an adjuvant for chemotherapy.

**Methods:** Three pancreatic cancer cell lines, PANC-1, AsPC-1, and MIA PaCa-2, were used. Cell growth was measured by MTT assay. Activities of caspase-3 and caspase-8, and caspase-9 were evaluated to determine the apoptosis pathway. Epithelial to mesenchymal transition was identified through mRNA expression of associated genes, which were transforming growth factor-beta, matrix metalloproteinase-2 and matrix metalloproteinase-9. Quantitative real time PCR was used to verify mRNA expression. To determine the effect on migration, transwell system was used.

**Results:** Ellagic acid inhibits pancreatic cancer cell growth by stimulating apoptosis. Apoptosis induced by ellagic acid in pancreatic cancer cells, is mediated by the activation of caspase-3 and caspase-9, and not caspase-8. Ellagic acid also suppresses migration via the inhibition of epithelial mesenchymal transition (EMT) in pancreatic cancer cells. Ellagic acid decreased the expression levels of transforming growth factor-beta, matrix metalloproteinase-2 and matrix metalloproteinase-9, while it increased that of E-cadherin.

**Conclusions:** These findings suggest that ellagic acid is useful in pancreatic cancer treatment.

**Keywords:** Ellagic acid; pancreatic cancer; proliferation; apoptosis; migration

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## Introduction

Ellagic acid (2,3,7,8-tetrahydroxy[1]-benzopyranol[5,4,3-cde]benzopyran-5,10-dione) is a natural dietary polyphenolic compound found in several fruits, nuts, and plants, such as strawberries, raspberries, blackberries, grapes, walnuts, and green tea (1). Ellagic acid has multiple biological activities,

such as antioxidant, anti-inflammatory and antifibrotic properties (2-4). Proinflammatory cytokines, such as IFN- $\gamma$ , TNF- $\alpha$ , and IL-1 (5-7), were suppressed and anti-inflammatory cytokine, such as IL-10, was increased by ellagic acid. Ellagic acid also suppresses the expression of cyclooxygenase 2 and inducible nitric oxide synthase via

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the inhibition of nuclear factor-kappa B (NF- $\kappa$ B) activity in several normal and cancer cells (8,9).

The anticancer effect of ellagic acid has also been reported in several types of cancers, including liver, skin, esophageal, colon, prostate, breast, and pancreatic cancer. The antiproliferative activity of ellagic acid on several tumor cell lines has been reported (3,10-14). Ellagic acid induces apoptosis by downregulation the expression of Bcl-2, that inhibits apoptosis, and upregulation of Bax, which functions as a proapoptotic factor (15,16). In addition, ellagic acid inhibited the proliferation by downregulating cyclin D1 and cyclin E1, which required for the entry into and completion of S phase and leaded to the arrest in the G1 phase of the cell cycle (17). However, the mechanisms mediating the anticancer effect of ellagic acid are still unclear.

Pancreatic cancer has an extremely poor prognosis, with a 5-year survival rate of less than 5% (18). Most patients with pancreatic cancer die from distant metastasis (19). Tumor metastasis is a complex process. First, cancer cells grow in the primary site with avoiding apoptosis process. Second, cancer cells migrate into and invade the extracellular matrix and intravasate into the blood or lymphatic vessel. It is believed that epithelial mesenchymal transition (EMT) of cancer cells is needed in this step. Then, tumor cells must survive in the bloodstream with avoiding anoikis. Next, metastatic cells extravasate into and colonize the metastatic site (20). Most anticancer agents promote apoptosis and suppress the migration and invasion of tumor cells. Pancreatic cancer has a poor response to conventional chemotherapy and radiotherapy (21).

In this study, we aimed to determine the effect of ellagic acid on the proliferation and migration of pancreatic cancer cells. We present the following article in accordance with the MDAR checklist (available at <http://dx.doi.org/10.21037/tcr-20-2446>).

## Methods

### Cells

PANC-1, AsPC-1 and MIA PaCA-2 were obtained from Byungheon Lee in Kyungpook National University, School of Medicine. PANC-1 and MIA PaCA-2 were maintained in Dulbecco's Modified Eagle medium containing penicillin and streptomycin (100 units/mL) and 10 % FBS at 37 °C in a 5 % CO<sub>2</sub> incubator. AsPC-1 was maintained in RPMI media, containing penicillin and streptomycin (100 units/mL)

and 10 % FBS at in 37 °C in 5 % CO<sub>2</sub> incubator. The study was conducted in accordance with the Declaration of Helsinki (as revised in 2013).

### MTT assay

Ellagic acid was obtained from Sigma (E2250), and reconstituted with 1 M NaOH at 10 mg/mL concentration. 0 to 1,000 mM was used after diluted with culture media. Tetrazolium 3-(4,5-dimethylthiazolyl-2)-2,5-diphenyltetrazolium bromide (MTT) buffer was purchased from Sigma (CT02). 5×10<sup>4</sup>/well cells were plated on 96 well plates with the indicated concentration of ellagic acid and incubated for 72 h at 37 °C. Cell growth was assayed every 24 h, using MTT solution. Then, 10 µL of MTT buffer was added and incubated for 2 h at 37 °C. 100 µL of dimethyl sulfoxide (DMSO) solution was added and the cells were incubated for 10 min at room temperature in dark conditions. Absorbance at 550 nm was measured. To test the effect of Interferon-gamma (IFN- $\gamma$ ), from R&D systems (285-IF-100), 0.5 ng/mL of IFN- $\gamma$  was treated 16 h prior to ellagic acid treatment. Each experiment was performed in triplicate.

### Caspase activity assay

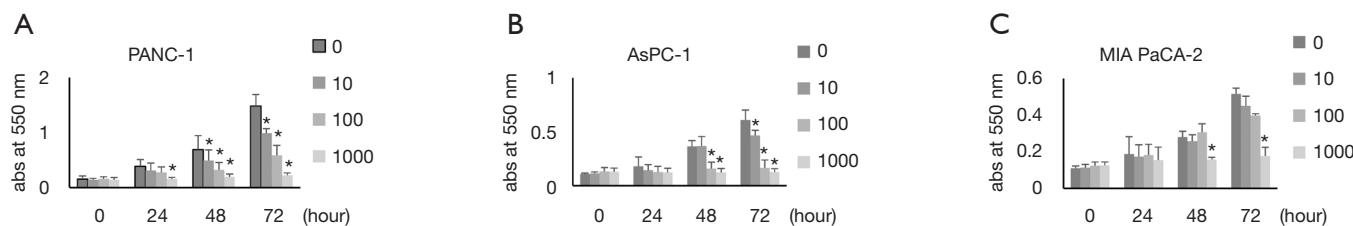
5×10<sup>4</sup>/well cells were plated on 96 well black plates with the indicated concentration of ellagic acid and incubated for 48 h at 37 °C. Caspase-3, caspase-8, and caspase-9 activities were determined using caspase multiplex kit from Abcam (ab219915) following the manufacturer's protocol. SpectraMax (Molecular Devices) was used to detect fluorescence. Activity was expressed as the fold increase in cells before ellagic acid treatment. Each experiment was performed in duplicate.

### Quantitative real time PCR

Total RNA was isolated from cells using an RNA extraction kit (Takara Bio, 9769A). cDNA was synthesized from 1 µg of total RNA using a cDNA synthesis kit (Takara Bio, 6100A) following the manufacturer's protocol. The primer pair sequences used were presented in Table 1. Quantitative real-time PCR was performed using SYBR Premix Ex Taq from Takara Bio (RR820A). Data were presented as fold changes in mRNA expression relative to GAPDH expression.

**Table 1** Primers list

Gene	Forward	Reverse
TGF- $\beta$	5'-CACTCCCGTGGCTTAGTG-3'	5'-CTGGCGAGCCTAGTTGGA-3'
E-cadherin	5'-GTCTGTAGGAAGGCACAGCC-3'	5'-TGCAACGTCGTTACGAGTCA-3'
MMP-2	5'-CAAGGACCAGGTTATTGGC-3'	5'-ATTCCTGCGAAGAACACAGC-3'
MMP-9	5'-TTGACAGCGACAAGAAGTGG-3'	5'-GCCATTACGTCGTCCTTAT-3'
GAPDH	5'-TCACCACCATGGAGAAGGC-3'	5'-GCTAAGCAGTTGGTGGTGCA-3'



**Figure 1** Ellagic acid inhibits pancreatic cancer cell proliferation. (A) PANC-1, (B) AsPC-1, (C) MIA PaCa-2. Cells were plated on microplates with ellagic acid and incubated in indicated times. 0 to 1,000  $\mu$ M ellagic acid were treated on each cell. MTT assay was performed as described in Materials and Methods. Data are presented as mean  $\pm$  SD from three individual experiments. \*, P<0.05 compared with 0  $\mu$ M ellagic acid treated group at each time.

#### Enzymes-linked immunosorbent assay (ELISA)

The concentrations of TGF- $\beta$  (R&D System, DB250), MMP-2 (R&D System, DMP2F0), and MMP-9 (R&D System, DMP900) in the culture supernatant were measured using an enzyme linked immunosorbent assay kit following the manufacturer's protocol.

#### Migration assay

$5 \times 10^4$  cells were suspended in media containing 5% FBS and were plated in the upper chamber of Transwell (8- $\mu$ m pore size; Corning, Costar) in 24-well plates. A medium containing 10% FBS was added to the lower chamber. The indicated concentration of ellagic acid was added to both chambers. Migrated cells on the Transwell membrane were fixed after 16 h of incubation. Cells were stained with crystal violet, and the numbers were calculated using MacBiophotonics Image J software.

#### Statistical analysis

The results were presented as mean  $\pm$  S.D. from at least three independent experiments. All statistical analyses were

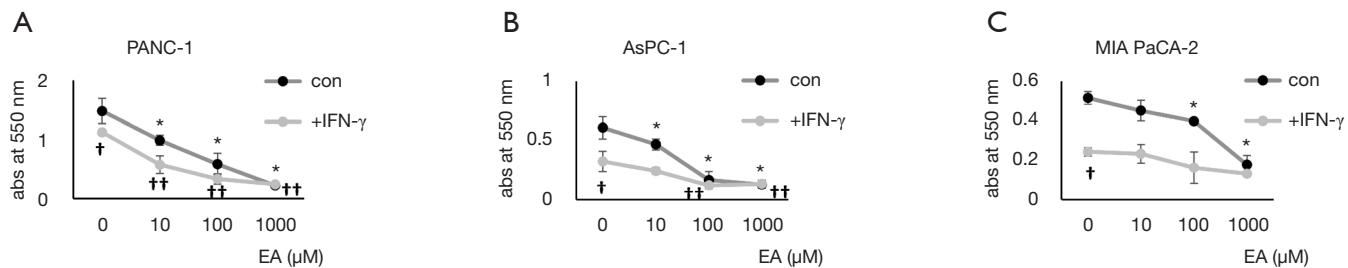
performed using the paired *t*-test. Data were considered significant if the P<0.05.

## Results

#### Effect of ellagic acid on the growth of pancreatic cancer cell lines

To evaluate the effect of ellagic acid on pancreatic cancer, three different cell lines, PANC-1, AsPC-1 and MIA PaCa-2, were used. Cell growth in presence of 10 to 1,000  $\mu$ M of ellagic acid was performed every 24 h for 72 h by MTT assay. Ellagic acid inhibited the proliferation of all three cell lines (Figure 1). However, the inhibitory concentration varied depending on cell types. Proliferation of PANC-1 was inhibited after adding 10  $\mu$ M of ellagic acid, and its inhibition was dose dependent. AsPC-1 proliferation was inhibited after adding 100  $\mu$ M of ellagic acid, while MIA-PaCa-2 proliferation was only prohibited after adding 1,000  $\mu$ M of ellagic acid.

The inhibitory effect of ellagic acid after IFN- $\gamma$  treatment was examined. IFN- $\gamma$  inhibited the expression of pancreatic cancer cells 16 h prior to ellagic acid treatment, while IFN- $\gamma$  inhibited the growth of all types of pancreatic



**Figure 2** IFN- $\gamma$  inhibits pancreatic cancer cell proliferation. (A) PANC-1, (B) AsPC-1 and (C) MIA PaCa-2. The cells were plated on microplates with ellagic acid and incubated at 48 h. Next, 0 to 1,000  $\mu$ M ellagic acid were added to each cell. MTT assay was performed as described in the Materials and Methods. Data are presented as mean  $\pm$  SD from three individual experiments. \*, P<0.05 compared with 0 h; †, P<0.05; ‡, P<0.01 compared with control group and IFN- $\gamma$  treatment group.

cancer cells. Ellagic acid further inhibited the growth of PANC-1 and AsPC-1 cells in dose-dependent manner. Further inhibitory effects of ellagic acid were not observed in MIA PaCa-2 cells (Figure 2).

#### Effect of ellagic acid on caspase-3 and caspase-9 activity in pancreatic cancer cells

Caspase-3, caspase-8, and caspase-9 activities were measured using a multiplex activity assay kit as described in Materials and Methods. Pancreatic cells were treated with IFN- $\gamma$  and ellagic acid for 48 h. The caspase-3 and caspase-9 activities increased after treatment with ellagic acid in a dose-dependent manner; however, the caspase-8 activity in PANC-1 cells and AsPC-1 cells remained unchanged. IFN- $\gamma$  did not affect the caspase activity in PANC-1 cells but affected that in AsPC-1 cells (Figure 3A,B,C,D,E,F). The caspase-3, caspase-8, and caspase-9 activities in MIA PaCa-2 cells remained unchanged after ellagic acid treatment without IFN- $\gamma$  treatment. After IFN- $\gamma$  treatment, only caspase-3 activity was increased dose-dependently by ellagic acid (Figure 3G,H,I).

#### Ellagic acid promoting migration and EMT in pancreatic cancer cells

TGF- $\beta$  and E-cadherin mRNA expression was examined by q-PCR. TGF- $\beta$  expression was inhibited, and E-cadherin expression was increased by ellagic acid in PANC-1 and AsPC-1 cells in a dose- and time-dependent manner (Figure 4A,B,C,D). TGF- $\beta$  and E-cadherin expression in MIA PaCa-2 cells was remained unchanged by ellagic acid

(Figure 4E,F).

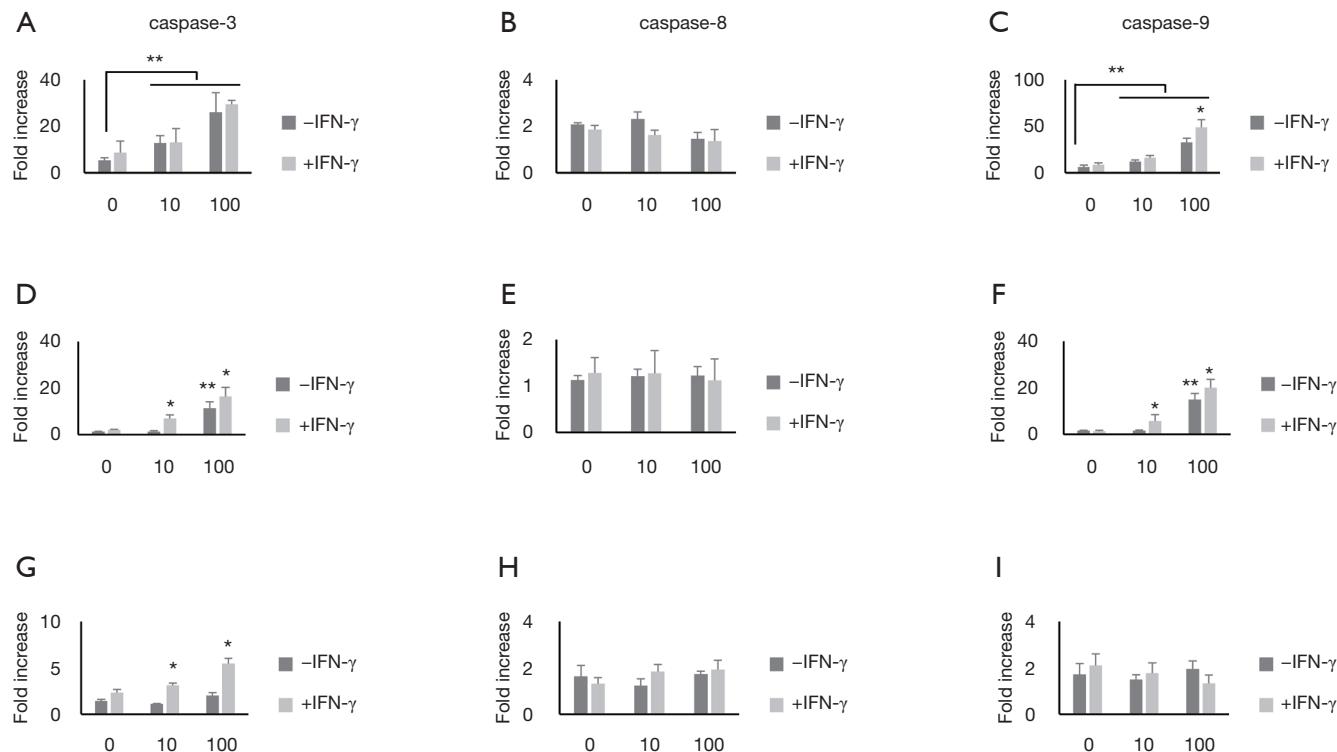
MMP-2 and MMP-9 mRNA expression was also evaluated. Both MMP-2 and MMP-9 expression was inhibited by ellagic acid in PANC-1 and AsPC-1 cells in a dose- and time-dependent manner (Figure 5A,B,C,D). Ellagic acid did not affect the MMP-2 and MMP-9 expression in MIA PaCa-2 cells (Figure 5E,F).

The Inhibitory effect of ellagic acid on TGF- $\beta$ , MMP-2, and MMP-9 expression was confirmed by ELISA. The expression levels of TGF- $\beta$ , MMP-2, and MMP-9 proteins in PANC-1 and AsPC-1 cells were decreased by ellagic acid in a dose-dependent manner (Figure 6A,B,C,D,E,F). TGF- $\beta$ , MMP-2, and MMP-9 protein expression in MIA PaCa-2 cells remained unchanged after ellagic acid treatment (Figure 6G,H,I).

The effect of ellagic acid on pancreatic cell migration was examined using Boyden chambers. Pancreatic cells were plated with 2% FBS on upper wells, and migration was induced by 10 % FBS in lower wells. The migrated cells were counted. PANC-1 and AsPC-1 cell migration was inhibited when ellagic acid was present; however the migration of MIA PaCa-2 cell was not inhibited (Figure 7A,B,C).

#### Discussion

Ellagic acid has been shown to have an antiproliferative effect on pancreatic cancer cells both *in vitro* and *in vivo* used by PANC-1 (14,22). In this report, two other pancreatic cancer cells were used. Proliferation of all three pancreatic cell lines, PANC-1, AsPC-1, and MIA PaCa-2, was inhibited by ellagic acid; however, the affected dose and



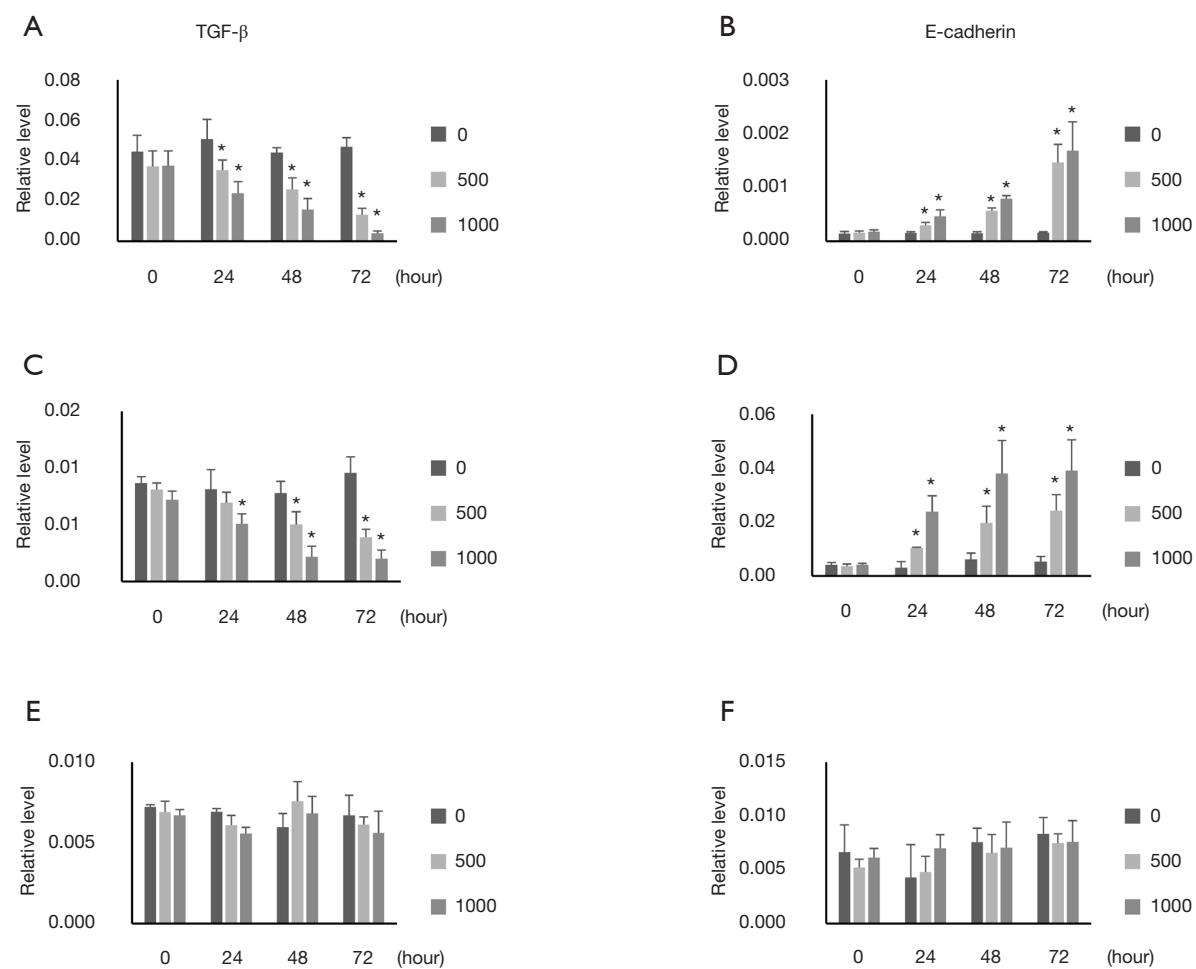
**Figure 3** Ellagic acid inhibits caspase-3 and caspase-9 activity. (A,B,C) PANC-1, (D,E,F) AsPC-1, (G,H,I) MIA PaCA-2. Cells were plated on microplates with ellagic acid and incubated for 48 h. 0 to 1,000  $\mu$ M ellagic acid were treated on each cell with or without IFN- $\gamma$ . Caspase activity was measured by caspase multiplex kit as following manufacture's protocol. Activity was calculated as the fold increase from cells before ellagic acid treatment. Data are presented as mean  $\pm$  SD from three individual experiments. \*\*,  $P<0.05$  compared with control group and IFN- $\gamma$  treatment group. \*,  $P<0.05$  compared with 0 h.

time were different for each cell (*Figure 1*). The reduction in cell growth of PANC-1 was observed when 10  $\mu$ M ellagic acid was added, but 1,000  $\mu$ M of ellagic acid was needed to reduce MIA PaCA-2 cell growth within 48 h. AsPC-1 growth decreased within 48 h after adding 100  $\mu$ M ellagic acid. Taken together, PANC-1 is most sensitive, while MIA PaCA-2 is most resistant to ellagic acid. The effect of ellagic acid on a xenograft model of pancreatic cancer cells was not evaluated in this study. Further studies using different pancreatic cancer cell lines or patient- derived xenograft models are needed to verify the different effects of ellagic acid.

To determine the cause of growth reduction by ellagic acid, the activities of caspase-3, caspase-8, and caspase-9 were measured. The caspase-3 and caspase-9 activities in PANC-1 and AsPC-1 cells increased after ellagic acid treatment, but the caspase-8 activity did not affect cell

growth. The caspase activity did not increase after ellagic acid treatment in the MIA PaCA-2 cells (*Figure 3*). Caspase activity was not detected in MIA PaCA-2 cells, which are not affected by ellagic acid. Caspase signaling is mediated by apoptosis, a type of programmed cell death, which is conducted in two pathways: the extrinsic pathway and the intrinsic pathway. The extrinsic pathway is mediated by caspase-8, and the intrinsic pathway can be initiated by caspase-9. Caspase-3 is activated in both pathways and triggers apoptosis (23,24). This finding suggests that the inhibition of pancreatic cancer cell growth by ellagic acid is caused by apoptosis and is triggered by the intrinsic pathway.

IFN- $\gamma$  has been reported to inhibit the proliferation of pancreatic cancer cells, including AsPC-1, Capan-1, Capan-2, and Dan-G cells, via caspase-1 dependent induction of apoptosis (25). IFN- $\gamma$  also promotes EMT in

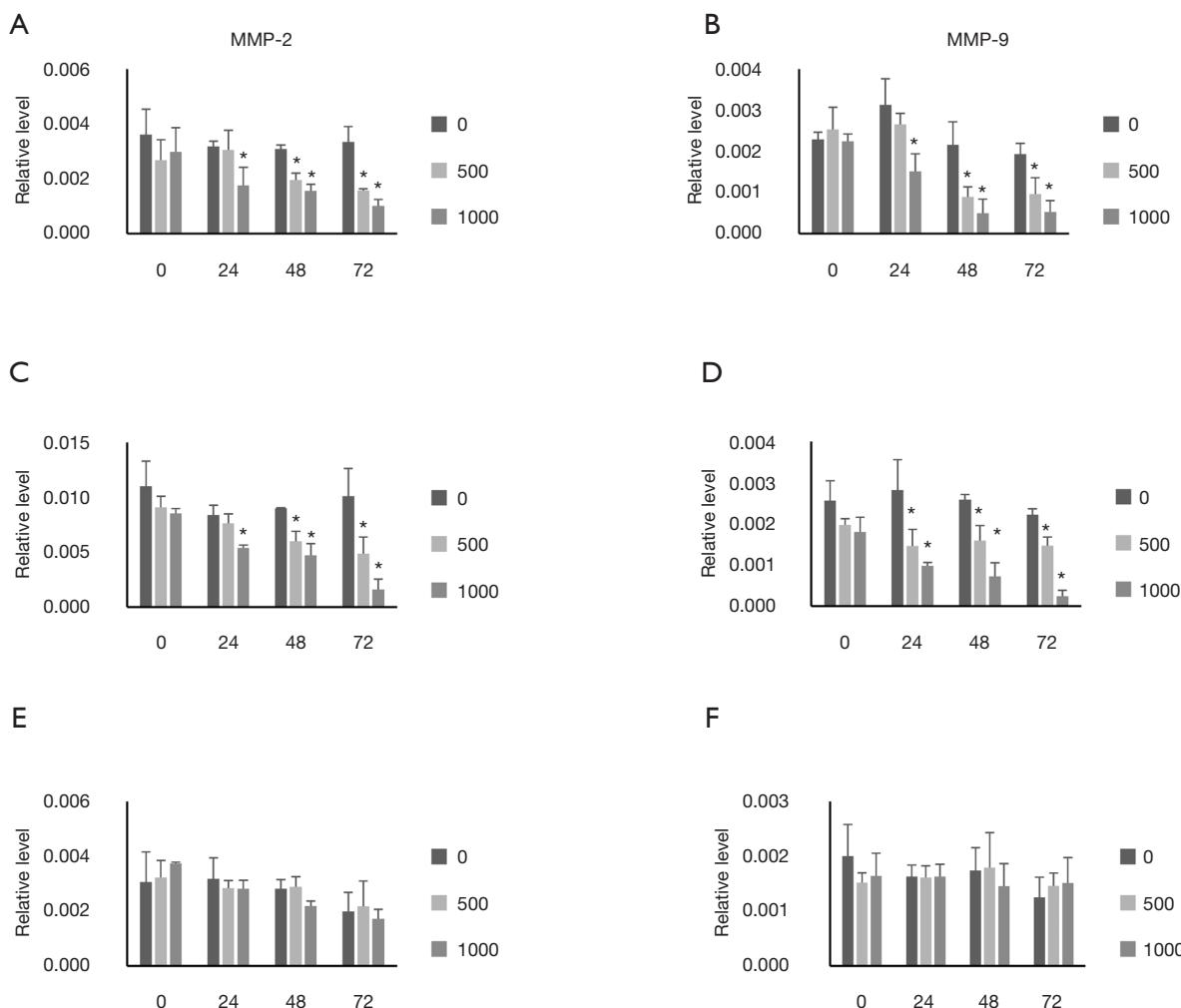


**Figure 4** Ellagic acid inhibits TGF- $\beta$  mRNA expression and promotes E-cadherin mRNA expression. (A,B) PANC-1, (C,D) AsPC-1, (E,F) MIA PaCa-2. Cells were incubated with each indicated concentration ellagic acid for 72 h. During incubation, cells were collected every 24 h. RNA was isolated from cells and cDNA was synthesized from RNA. q-PCR was performed and relative expression of GAPDH were presented. (A,C,E) relative TGF- $\beta$  expression of GAPDH. (B,D,F) relative E-cadherin expression of GAPDH. All data are presented as mean  $\pm$  SD from a minimum of three individual experiments. \*, P<0.05 compared with control.

pancreatic cells (26). In this study, the effect of ellagic acid on cell growth after IFN- $\gamma$  treatment was also examined. IFN- $\gamma$  inhibits the growth of all three different pancreatic cells, PANC-1, AsPC-1, and MIA PaCa-2. Ellagic acid further inhibited the growth of PANC-1 and AsPC-1, but not the growth of MIA PaCa-2 after IFN- $\gamma$  treatment (*Figure 2*). However, caspase-3 and caspase-9 activities were not further increased by ellagic acid and IFN- $\gamma$  treatment in PANC-1 cells. AsPC-1 and MIA PaCa-2 cells have further increased the activity of caspase-3 by ellagic acid and IFN- $\gamma$  treatment, but this effect was small. The absence of changes in proliferation after IFN- $\gamma$  treatment did not increase the

severity of apoptosis.

Ellagic acid blocks the effects of proinflammatory agents and mainly inhibits the expression of cyclooxygenase 2 (COX-2) and nuclear factor-kappa B (NF- $\kappa$ B) in several types of cells, including pancreatic cancer cells (1,14,22). E-cadherin expression increases because of the presence of ellagic acid in pancreatic cancer cells (22). The expression of MMP-2, MMP-9, and TGF- $\alpha$  decreased because of the presence of ellagic acid in hepatocytes and breast cancer cells (27,28). Thus, we hypothesized that ellagic acid may inhibit migration of pancreatic cancer cells. To confirm this, the levels of TGF- $\beta$ , E-cadherin, MMP-2, MMP-9

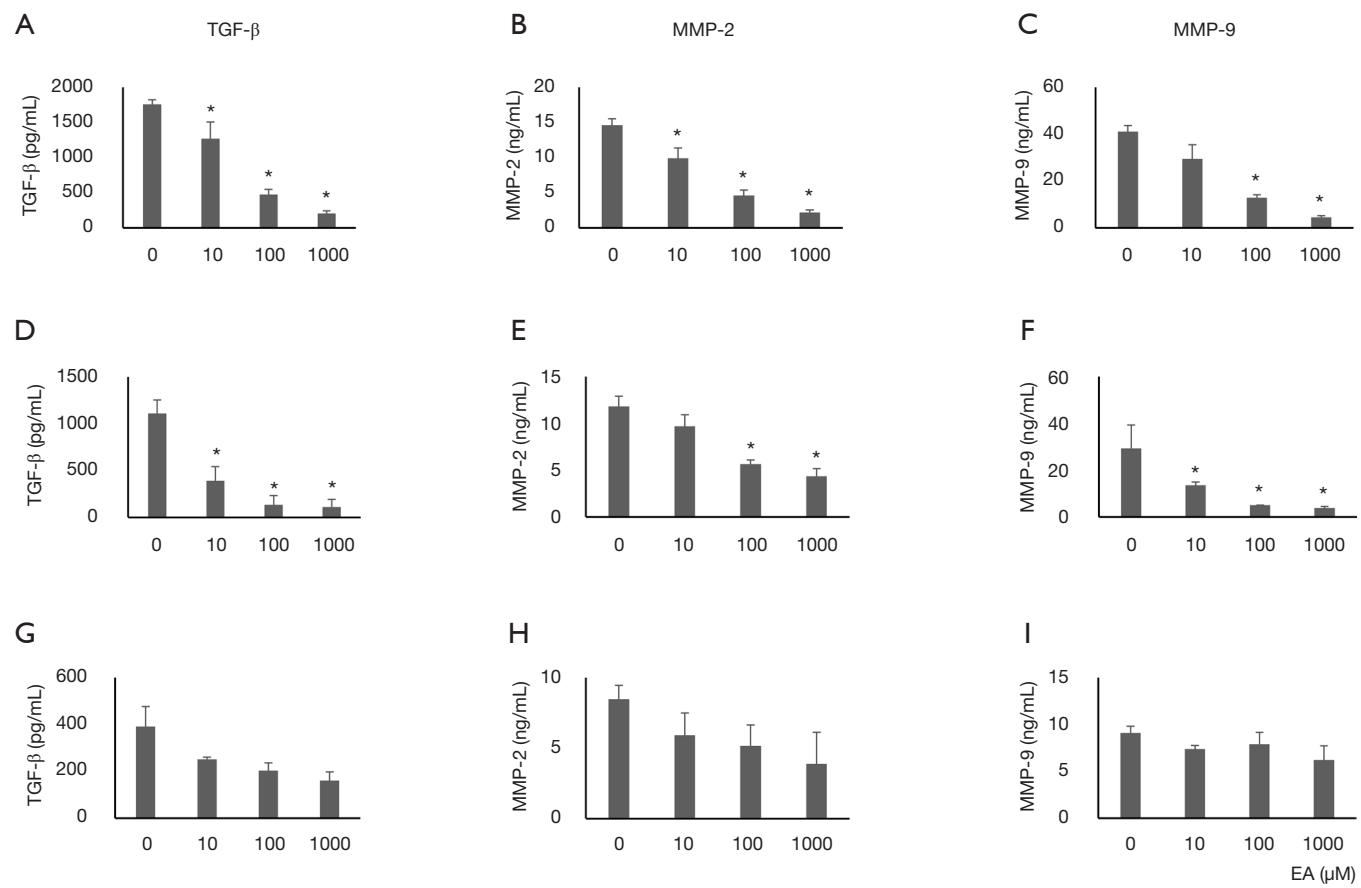


**Figure 5** Ellagic acid inhibits MMP-2 and MMP-9 mRNA expression. (A,B) PANC-1, (C,D) AsPC-1, (E,F) MIA PaCa-2. Cells were incubated with each indicated concentration ellagic acid for 72 h. During incubation, cells were collected every 24 h. RNA was isolated from cells and cDNA was synthesized from RNA. q-PCR was performed and relative expression of GAPDH were presented. (A,C,E) relative MMP-2 expression of GAPDH. (B,D,F) relative MMP-9 expression of GAPDH. All data are presented as mean  $\pm$  SD from a minimum of three individual experiments. \*, P<0.05 compared with control.

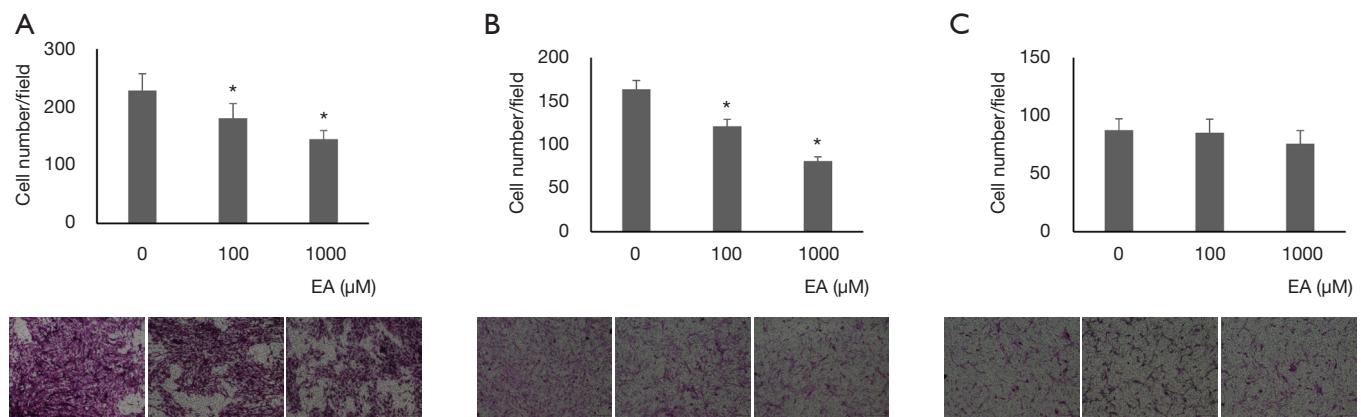
mRNA, and protein expression were measured. All of these decreased in PANC-1 and AsPC-1 cells, but not in MIA PaCa-2 cells. The concentration of ellagic acid decreased in a dose- and time-dependent manner (Figures 4-6). Ellagic acid also inhibits the migration of PANC-1 and AsPC-1 cells, but not the migration of MIA PaCa-2 cells. Taken together, ellagic acid may suppress EMT in pancreatic cells.

The effect of ellagic acid on proliferation and migration of pancreatic cancer cells was examined in this study. Ellagic acid inhibits proliferation and migration of

pancreatic cancer cells. Inhibition of proliferation is caused by increased apoptosis. Ellagic acid-induced apoptosis occurs in the caspase-3 and caspase-9 pathways, but not in the caspase-8 pathway. Inhibition of migration occurs with the decrease in E-cadherin and decrease in TGF- $\beta$ , MMP-2, and MMP-9 levels. This effect of ellagic acid is only observed in PANC-1 and AsPC-2 cells, but not in MIA PaCa-2 cells. These results suggest that ellagic acid may inhibit pancreatic cancer, but it is dependent on the characteristics of cancer cells.



**Figure 6** Ellagic acid inhibits TGF- $\beta$ , MMP-2 and MMP-9 protein expression. (A,B,C) PANC-1, (D,E,F) AsPC-1, (G,H,I) MIA PaCa-2. Cells were incubated with each indicated concentration ellagic acid for 48 h. Cell supernatant was collected and ELISA assay was performed. (A,D,G) TGF- $\beta$ . (B,E,H) MMP-2. (C,F,I) MMP-9. All data are presented as mean  $\pm$  SD from a minimum of three individual experiments. \*, P<0.05 compared with control.



**Figure 7** Ellagic acid inhibits migration of pancreatic cancer cells. (A) PANC-1, (B) AsPC-1, (C) MIA PaCa-2. Cells were plated on upper chamber, and incubated for 16 h. indicated times. After remove unmigrated cells on upper chamber, migrated cells were fixed and stained with crystal violet. Cell number in high-power field were counted and presented. Data are presented as mean  $\pm$  SD from three individual experiments. \*, P<0.05 compared with control. Microscopic images were also presented. Scale bar: 200  $\mu$ m.

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## Footnote

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**Data Sharing Statement:** Available at <http://dx.doi.org/10.21037/tcr-20-2446>

**Conflicts of Interest:** All authors have completed the ICMJE uniform disclosure form (available at <http://dx.doi.org/10.21037/tcr-20-2446>). The authors have no conflicts of interest to declare.

**Ethical Statement:** The authors are accountable for all aspects of the work in ensuring that questions related to the accuracy or integrity of any part of the work are appropriately investigated and resolved. The study was conducted in accordance with the Declaration of Helsinki (as revised in 2013).

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