Proscillaridin A induces mitochondrial damage and autophagy in pancreatic cancer and reduces the stability of SMAD4 in Panc-1 cells

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Background: Pancreatic cancer (PC) is a highly metastatic and lethal cancer with a very low overall 5-year survival rate. There is an urgent need for identifying new therapeutic agents for this deadly disease. Cardiac glycosides (CGs) have been traditionally used for their potent cardiovascular activities and have also recently been reported to exhibit anti-tumor effects. Proscillaridin A (Pro A), a natural CG, has been shown to display anti-tumor effects on multiple cancer types.

Methods: The cytotoxic effect of Pro A on PC cells was determined using cell viability assay, colony formation assay and transwell assay *in vitro*. Cell apoptosis, cell cycle, reactive oxygen species (ROS) generation, intracellular Ca^{2+} levels and mitochondrial membrane potential (MMP) were assayed by flow cytometry. Panc-1-xenografted mice model was used to evaluate Pro A's effect in tumor growth. Mitochondria morphology was observed by transmission electron microscopy. LC3 aggregation was assessed by GFP-LC3 fluorescence microscopy. Gene expression was assayed by western blot or real-time quantitative polymerase chain reaction (qPCR).

Results: Pro A inhibits the proliferation, migration and invasion of Panc-1, BxPC-3 and AsPC-1 PC cells *in vitro*, and Panc-1 cells display the highest sensitivity with an IC50 at the nano-molar level. *In vivo*, Pro A treatment inhibits tumor progression in Panc-1 xenograft nude mice. Pro A treatment promotes both cell apoptosis and autophagy, and Pro A-treated PC cells display characteristics of mitochondrial damage including increased ROS generation, intracellular Ca^{2+} levels and disruption of MMP. In addition, high sensitivity towards Pro A of Panc-1 cells compared to BxPC-3 and AsPC-1 cells could be partially attributed to the loss of endogenous SMAD4 expression in the latter.

Conclusions: Our findings suggest that Pro A constitutes a promising therapeutic candidate for certain types of PC.

Keywords: Pancreatic cancer (PC); proscillaridin A (Pro A); apoptosis; autophagy; SMAD4

Introduction

Pancreatic cancer (PC) is one of the most aggressive and lethal cancers (1). Previous studies have shown that aging, environmental and behavioral changes are more closely related to the dramatic increase in the incidence of PC worldwide than genetic factors (2-5). A position paper published by the European Federation of Gastroenterology described PC as a medical emergency (6).

In the past few years, first- and second-line palliative systemic treatments and adjuvant therapies have made considerable progress in prolonging survival of PC patients. The combination therapy composed of folic acid, 5-fluorouracil, irinotecan and oxaliplatin increased the overall survival of patients with PC by 4.3 months (7,8). In addition, nab-paclitaxel combined with gemcitabine can improve the median progression-free survival rate and objective remission rate (9). However, the prognosis of PC is still very poor, and the 5-year survival rate is lower than 5% (10). There is an urgent need for new drugs and strategies against PC.

PC is characterized by complex genetic alterations. Several signaling pathways have been reported to play an important role in the pathogenesis of PC by regulating the proliferation and apoptosis of cancer cells (11). Mutations in at least 32 genes belonging to 10 cellular signaling pathways are associated with PC (12). Four genes, *KRAS*, *CDKN2A*, *TP53*, and *SMAD4*, are the most commonly mutated genes in PC (13-16), which may help to guide personalized cancer therapy. A previous study showed that the decrease in *Ras* activity could inhibit the proliferation of *KRAS* wild-type PC cells and increase the sensitivity to cetuximab (17).

Proscillaridin A (Pro A), one of the cardiac glycosides (CGs), has long been used to treat congestive heart failure, especially in the case of bradycardia, where its negative chronotropic effect is considered to be less than that of digitalis (18). In cardiomyocytes, Pro A binds and inhibits Na^+/K^+-ATPase, resulting in enhanced myocardial contraction (19). Pro A has shown anti-tumor effects on various cancers, including breast cancer (20), prostate cancer (21), hepatocellular carcinoma (22), leukemia (23) and non-small cell lung cancer (24). Pro A has also been demonstrated to inhibit PC cells at micro-molar concentrations using 3D spheroid cultures (25). The mechanism of Pro A inhibiting PC cells is not clear.

In this study, we demonstrate that Pro A inhibits PC cells at the nano-molar concentrations by inducing apoptosis and autophagy. Endogenous SMAD4 expression is associated with the sensitivity of PC cells to Pro A, while Pro A reduces the stability of SMAD4 protein. Our findings suggest that Pro A is a promising candidate for treating certain types of PC. We present the following article in accordance with the ARRIVE reporting checklist (available at https://atm.amegroups.com/article/view/10.21037/atm-22-1085/rc).

Methods

Cell culture and drug treatment

PC cell lines (Panc-1, AsPC-1, BxPC-3) were from the Cell Bank of Shanghai Institute of Biological Sciences, China Academy of Sciences. All cell lines were cultured in RPMI 1640 medium (Corning, New York, NY, USA) at 37 ℃ with 5% CO_2. Panc-1 cell cultures were supplemented with 10% fetal bovine serum (GIBCO, New York, NY, USA), while AsPC-1 and BxPC-3 cell cultures were supplemented with 20% fetal bovine serum. A master solution (10 mM) of Pro A (Sigma Aldrich, New York, NY, USA, #R214310) was prepared by dissolving Pro A in dimethyl sulfoxide (DMSO) and stored at –80 ℃. For mouse treatment, Pro A was first dissolved in DMSO at 50 mg/mL and then diluted with phosphate buffer saline (PBS) to a final concentration of 0.5 mg/mL before use. The solution was briefly heated at 42 ℃ and sonicated till clear.

Transfection, lentivirus construction and transduction

pCDH-SMAD4 was generated by subcloning *SMAD4* cDNA amplified from pcDNA3.1-3xflag-SMAD4 (Youbio, Changsha, China) in pCDH (Addgene, New York, NY, USA). The coding DNA for the short hairpin RNA (shRNA) precursor targeting *SMAD4* was inserted into pLKO.1 (Addgene), and co-transfected with packaging plasmids psPAX2 and pMD2.G (Addgene) into HEK293T cells for production of recombinant lentivirus. Transfections were performed using TurboFect Transfection Reagent

(Thermo Fisher Scientific, Shanghai, China) following the manufacturer's instructions. The shRNA sequences are as follows: shSMAD4-1: 5'-CCGGGGTGTGCAGTTGG AATGTACTCGAGTACATTCCAACTGCACACCTT TTTG-3'; shSMAD4-2: 5'-CCGGGTAGGACTGCAC CATACACCTGCAGGTGTATGGTGCAGTCCTACT TTTTG-3'; shSMAD4-3: 5'-CCGGGTAGGACTGCA CCATACACCTGCAGGTGTATGGTGCAGTCCTAC TTTTTG-3'; shCtrl: 5'-CCGGAAAATGCAGTTGGAA TGTACTCGAGTACATTCCAACTGCATTTTTTTT TG-3'. PC cells were transduced by recombinant lentivirus in the presence of 2 µg/mL polybrene (Sigma-Aldrich). The expression of target genes was verified with western blot.

Colony formation assay

PC cells were inoculated in triplicates in 6-well plates (500 cells/well for Panc-1 and BxPC-3, 1,000 cells/well for AsPC-1), and treated with different concentrations of Pro A. Cells were left to form colonies at 37 ℃ for 14 days. The cell colonies were washed and fixed with 4% paraformaldehyde (PFA) for 15 min and stained with crystal violet for 20 min at room temperature. The colonies were recorded under a microscope and analyzed by ImageJ.

Cell migration and invasion assays

In cell migration assays using a 24-well transwell plate, PC cells were treated with different concentrations of Pro A for 24 h and then suspended in serum-free medium. The 100 µL of suspensions was added on top of the transwell membrane and 600 µL of serum-containing medium was added in the lower chamber. After 48 h, cells on the membrane were fixed with 4% PFA and stained with 0.1% crystal violet. In cell invasion assays, the extracellular matrix (Matrigel, BD Biosciences, New York, NY, USA) was added on top of the transwell membrane. The following steps were the same as in migration assays. Cells were examined under an inverted microscope and the average number of cells in five randomly selected areas were calculated.

The directional cell migration was examined using wound-healing assay. After Pro A treatment for 12 h, cells were inoculated into 6-well plates. When the cell confluence reached 80–90%, the culture medium was replaced with serum-free medium. A pipette tip was used to make a straight scratch. At 0, 6, 12, 24 and 36 h after scratch, the active edge of the cells was observed under a microscope and photographed.

Cell proliferation assay

The inhibitory effect of Pro A on cell proliferation was determined by cell counting kit-8 (CCK-8) (Dojindo, Kumamoto, Japan). Cells were inoculated in 96-well plates (6,000 cells/well for Panc-1 and BxPC-3, 7,000 cells/well for AsPC-1). The medium was refreshed 12 h after inoculation and cells were treated with increasing concentrations of Pro A with 5 wells for each concentration and cultured for 72 h. CCK-8 reagent was added and OD450 values were measured with Epoch Microplate Spectrophotometer (Bio-Tek, New York, NY, USA). Data were plotted and half-maximal inhibitory concentration (IC50) was calculated using Graphpad 7.

Cell cycle analysis and apoptosis assays

PC cells treated with Pro A for 48-h were fixed overnight with 70% ethanol, stained for 30 min at 37 ℃ in dark with propidium iodide (PI; Beyotime Biotechnology, Shanghai, China) for cell cycle analysis or with Annexin V-FITC Apoptosis Detection Kit (Dojindo) for apoptosis assays. Cells were then analyzed using FACSCalibur flow cytometry (BD Biosciences).

Reactive oxygen species (ROS) measurement

Cells were disassociated through trypsin treatment, washed three times with PBS, and then incubated with 10 µM 2',7'-dichlorofluorescein diacetate (DCFH-DA; Yeasen, Shanghai, China) for 30 min at 37 ℃ in dark. After the incubation, cells were washed and collected. Fluorescence was measured using FACScalibur flow cytometry (BD Biosciences) at 488 nm (excitation) and 525 nm (emission).

Determination of intracellular free Ca^{2+}

Pro A-treated PC cells were washed with PBS, resuspended in serum-free medium containing 4 µM Fluo-4 AM (Yeasen), and incubated for 30 min at 37 ℃ in dark. Cells were then washed with PBS and fluorescence was measured using FACScalibur flow cytometry (BD Biosciences) at 494 nm (excitation) and 516 nm (emission).

Determination of mitochondrial membrane potential (MMP)

Pro A-treated PC cells were washed with PBS and stained with 10 µg/mL JC-1 fluorescent probe (Yeasen) for 15 min at

37 ℃ in dark. Cells were then washed with PBS, and green and red fluorescence, which represent JC-1 monomer and aggregates respectively, was measured using FACSCalibur flow cytometry (BD Biosciences). MMP was calculated by a decrease in red/green fluorescence intensity ratio.

Western blot

Protein samples were mixed with 5× SDS loading buffer (Beyotime Biotechnology) and heated at 100 ℃ in metal bath. After thorough separation, proteins were transferred onto nitrocellulose membrane (Millipore, New York, NY, USA) using Trans-blot Turbo system (Bio-Rad, Hercules, CA, USA). The membrane was blocked with PBST containing 3% bovine serum albumin (Sangon Biotech, Shanghai, China) and then incubated with a specific primary antibody overnight at 4 ℃. Antibodies for PARP, cleaved-PARP, Caspase 9, Caspase 3, cleaved-Caspase 3, P62, LC3, mTOR, Phospho-mTOR, Raptor, ATG7, SMAD4, E-cadherin and ZO-1 were from Cell Signaling Technology (Boston, MA, USA). Antibodies for MMP2 were from Servicebio (Shanghai, China). Signals were detected using ECL substrate (Millipore) and captured using ChemiDoc XRS+ (Bio-Rad).

Reverse transcription-quantitative polymerase chain reaction (RT-qPCR)

Total RNA was extracted with TRIzol Reagent (Invitrogen, Los Angeles, CA, USA). cDNA was synthesized by reverse transcription using Prime Script RT reagent Kit (TaKaRa, Kyoto, Japan) according to the manufacturer's instructions. Quantitative real-time PCR was performed with SYBR Green SuperReal Premix (TIANGEN, Beijing, China) on CFX Connect Real-Time System (Bio-Rad). The expression of target genes was determined using the $2^{-\Delta\Delta Ct}$ method and normalized to that of the housekeeping gene *GAPDH*. Primer sequences are shown in Table S1.

GFP-LC3 fluorescence microscopy

Cells grown on microscope cover glasses (Jingan Biological, Shanghai, China) in a 12-well plate were transfected with pGFP-LC3 (a gift from Dr. Qiang Deng of Shanghai Medical College). Six hours after transfection, cells were treated with Pro A for 24 h. Cells were then fixed with 4% PFA, treated with 0.2% Triton X-100 and stained with 4',6-diamidino-2'-phenylindole (DAPI). Cell images were obtained using a laser scanning confocal microscope (Leica, Heidelberg, Germany).

Tumor xenograft mouse model

Four-week-old female BALB/c nude mice weighing about 16 g were from Shanghai Jihui Laboratory Animal Care and housed in Fudan experimental SPF animal center. Seven days after subcutaneous implantation of 5×10^6 Panc-1 cells into the left armpit, 12 mice were randomly divided into two groups and injected with vehicle (PBS) or Pro A (6.5 mg/kg) every two days for 3 weeks before sacrifice. Tumor volumes were measured every three days. Experiments were performed under a project license (No. 202103005S) granted by the Animal Ethics Committee of School of Basic Medical Sciences, Fudan University and were performed in accordance with the guidelines of the National Institutes of Health on the care and use of laboratory animals.

Statistical analysis

Data were analyzed using Graphpad Prism 7 and expressed as mean ± standard error of mean (SEM). Student *t*-test and one-way analysis of variance (ANOVA) were used to compare two groups and multiple groups, respectively. $P<0.05$ was considered statistically significant.

Results

Pro A inhibits proliferation, migration and invasion of PC cell lines in vitro

To investigate the anti-tumor activity of Pro A on PC, we first examined the cytotoxic effect of Pro A on three PC cell lines Panc-1, BxPC-3 and AsPC-1. A 72-h treatment of Pro A inhibited cell proliferation of all three cell lines in a dose-dependent manner. The half-maximal inhibitory concentration (IC50) values for Panc-1, BxPC-3 and AsPC-1 were 35.25, 180.3 and 370.9 nM, respectively (*Figure 1A*, *Table 1*). Cell proliferation decreased with the extension of treatment, showing that the inhibitory effect was also time-dependent (*Figure 1B*). Notably, Panc-1 cells were more sensitive to Pro A than BxPC-3 and AsPC-1 cells. The inhibitory effect of Pro A was further confirmed by colony formation assays (*Figure 1C*, *Table 1*) and cell morphology (Figure S1). These results indicated that Pro A inhibits the proliferation of PC cells and different PC cell lines display different sensitivity to Pro A.

The effects of Pro A on migration and invasion capacities of PC cell lines were tested using transwell and wound-healing assays. Migration and invasion were significantly reduced after Pro A treatment (*Figure 2A,2B*, *Table 1*).
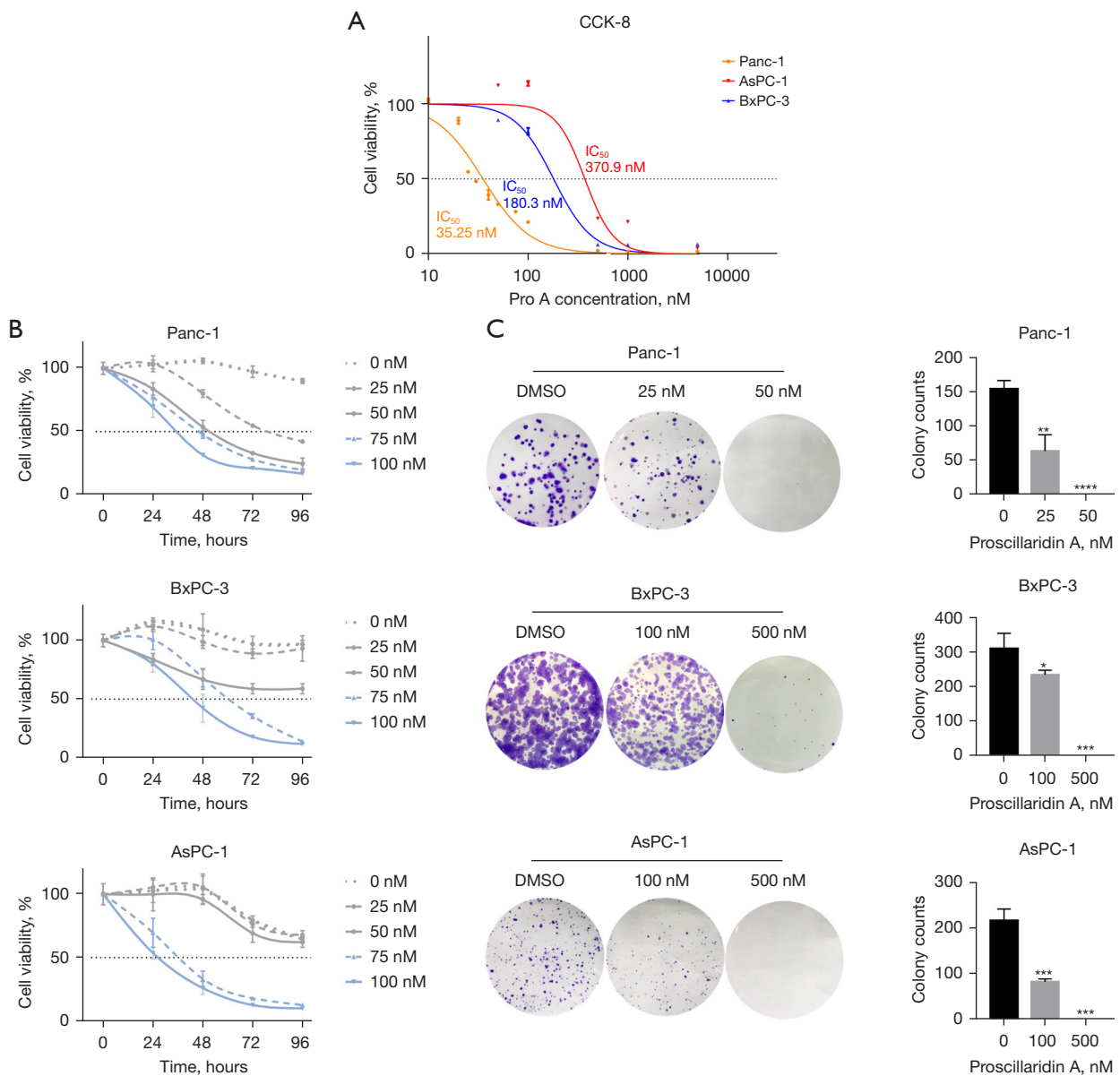
Figure 1 Pro A inhibits proliferation of PC cell lines. (A) PC cell lines were treated with different concentrations of Pro A for 72 h and cell viability was measured using CCK-8 assay. Half-maximal inhibitory concentration (IC50) values were calculated and listed on the right. (B) PC cells were treated with indicated concentrations of Pro A for indicated times and cell viability was measured using CCK-8 assay. (C) Colony formation assay was performed using PC cells treated with indicated concentrations of Pro A for 2 weeks. Each experiment was repeated three times. The colonies were fixed with 4% PFA and stained with 0.1% crystal violet. Magnification: ×1. Error bars represent SEM. *, $P<0.05$; **, $P<0.01$; ***, $P<0.001$; ****, $P<0.0001$. CCK-8, cell counting kit-8; PC, pancreatic cancer; Pro A, proscillaridin A; DMSO, dimethyl sulfoxide; PFA, paraformaldehyde; SEM, standard error of the mean.

Epithelial-mesenchymal transition (EMT), a process in which epithelial cells acquire mesenchymal features, is associated with tumor initiation, invasion, metastasis, and resistance to therapy. We detected the key molecular markers of EMT by western blot to determine whether Pro A affects the EMT process. Pro A treatment of PC cells resulted in decreased expression of cell invasion and metastasis markers Vimentin and MMP2, and increased

Table 1 Inhibitory effects of Pro A on PC cell lines

Pro A on cell line	Panc-1 [IC50 (CCK-8) =35.25 nM]		BxPC-3 [IC50 (CCK-8) =180.3 nM]		AsPC-1 [IC50 (CCK-8) =370.9 nM]	
	25 nM	50 nM	100 nM	500 nM	100 nM	500 nM
Colony formation	41.1%**	0%****	75.2%*	0%***	38.3%***	0%***
Migration	60.3%****	9.5%****	13.3%****	0%****	44.5%****	0%****
Invasion	17.5%****	0%****	80.7%	15.6%****	81.1%***	25.6%****
Apoptosis	122.0%	213.3%***	134.3%	524.3%****	93.5%	192.1%**
ROS	140.6%	175.3%*	202.3%****	285.2%****	195.1%	577.2%****
Ca^{2+}	100.4%	130.5%**	227.6%***	421.4%****	79.3%*	204.4%****
MMP	42.0%*	30.9%*	127.7%	22.0%*	94.4%	12.2%***

Results presented in figures are summarized here. Percentages are calculated by normalizing data from Pro A-treated cells against DMSO-treated cells. Statistical significance was calculated using Student t-test. *, P<0.05; **, P<0.01; ***, P<0.001; ****, P<0.0001. Pro A, proscillaridin A; PC, pancreatic cancer; CCK-8, cell counting kit-8; ROS, reactive oxygen species; MMP, mitochondrial membrane potential; DMSO, dimethyl sulfoxide.

expression of markers of suppression of cell invasion E-cadherin and ZO-1 (*Figure 2C*). These data suggested that Pro A reduces the metastatic potential of the PC cells.

Pro A inhibits tumor growth in Panc-1-xenografted mice

To evaluate the anti-tumor effect of Pro A on tumor growth *in vivo*, Panc-1 cells were subcutaneously injected into the left armpit of nude mice (5×10⁶ cells/mouse). Mice were randomly divided into two groups, receiving either 6.5 mg/kg of Pro A or the same volume of PBS every 2 days. Tumor volumes were measured every 3 days. The results showed that starting from day 15 post-treatment, the mean tumor volume of Pro A-treated mice was significantly smaller than the PBS group (*Figure 3A*). On day 24, mice were sacrificed and the mean tumor weight of Pro A-treated mice was markedly lower than PBS group (0.11±0.02 *vs.* 0.24±0.05 g, P<0.0001, n=6) (*Figure 3B,3C*). The assays of TUNEL, Ki67 and cleaved-Caspase 3 can indicate cell proliferation and apoptosis, respectively. Immunohistochemical results showed that compared with PBS treated mice, tumors in Pro A-treated mice contained fewer Ki67 positive cells and more cleaved-Caspase 3 positive cells (*Figure 3D*). These results indicate that Pro A inhibits PC progression in xenografted mice.

Pro A induces apoptosis and blocked cell cycle progression

Compared with control cells, PC cells treated with Pro A for 48 h showed significantly increased apoptosis as assayed by an annexin V/PI staining. Similarly, Panc-1 cells were more sensitive to Pro A induced apoptosis than BxPC-3 and AsPC-1 cells (*Figure 4A*, *Table 1*). In Panc-1 cells treated with Pro A for 48 h, molecular markers of apoptosis (decreased levels of caspase 9 and increased levels of cleaved caspase 3 and PARP) could be easily observed (*Figure 4B*). We further examined the effect of Pro A on PC cell cycle progression. The results showed that Pro A treatment resulted in the increase of G2 phase PC cells and the decrease of S phase PC cells (*Figure 4C*). These results suggest that Pro A can induce PC cell apoptosis and block the progression of cell cycle.

Pro A induces mitochondrial damage in PC cells

Mitochondria are the site of adenosine triphosphate (ATP) synthesis. Mitochondrial damage is characterized by a series of changes, including cytoplasmic Ca^{2+} accumulation, ROS production and MMP decrease (26). We first examined mitochondria in PC cells treated with Pro A by transmission electron microscopy. In Pro A-treated PC cells, mitochondria became long rod-shaped and swollen (*Figure 5A*). DCFH-DA staining showed that ROS levels increased significantly in PC cells treated with Pro A for 48 h and in a dose-dependent manner (*Figure 5B*, *Table 1*). Intracellular Ca^{2+} levels, assayed using fluo-4-Ca^{2+} imaging, also increased with Pro A treatment (*Figure 6A*, *Table 1*). MMP in PC cells was analyzed using JC-1 staining, which showed significant decrease in MMP after Pro A treatment (*Figure 6B*, *Table 1*). The data showed that Pro A induced mitochondrial damage.
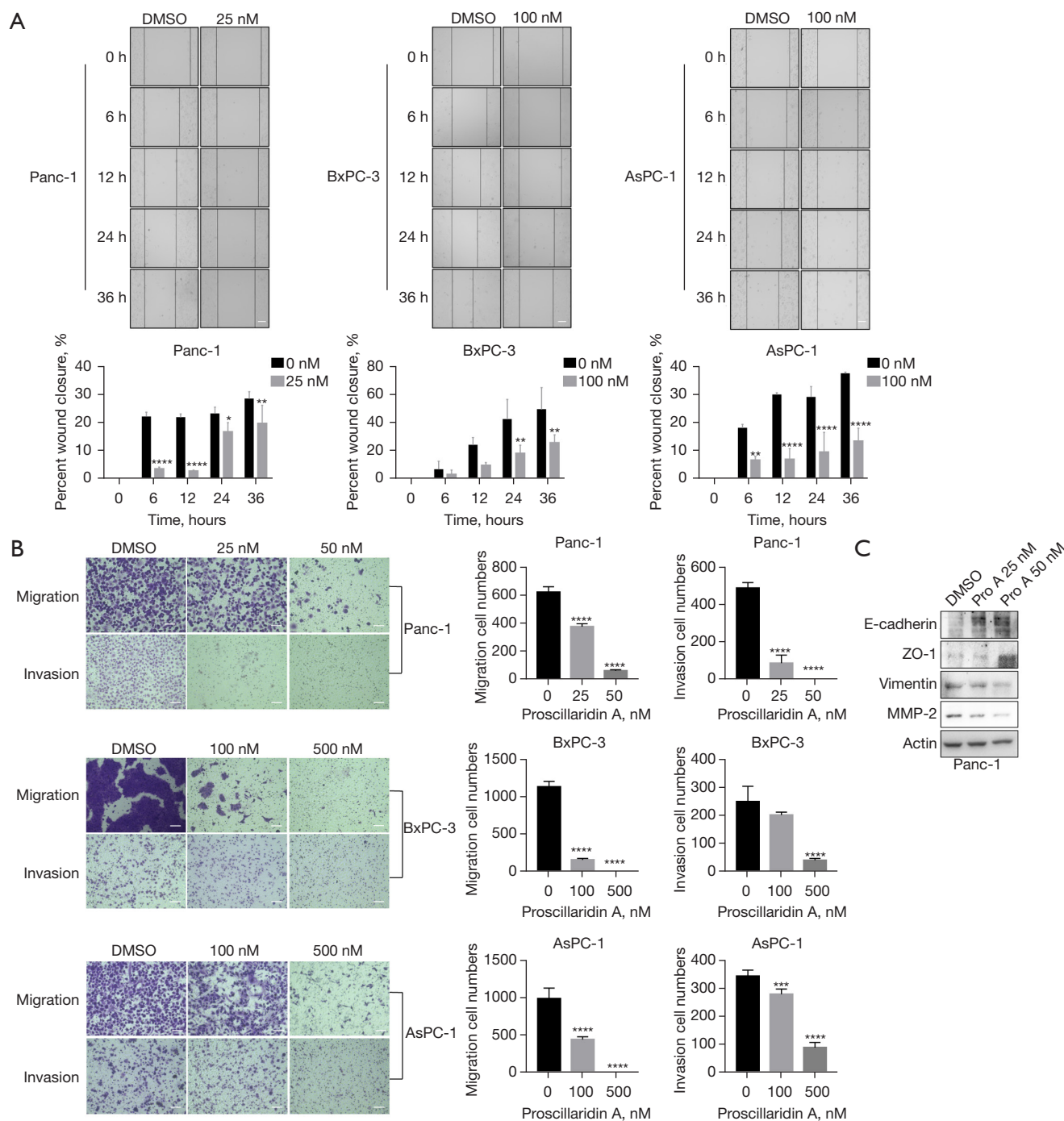
Figure 2 Pro A inhibits migration and invasion of PC cells *in vitro*. (A) PC cells were cultured in serum-free medium and treated with indicated concentrations of Pro A. Wound assay was performed and healing was measured at 0, 6, 12, 24 and 36 h. Bars: 100 μm. (B) Transwell migration and invasion assays were performed using Panc-1, BxPC-3 and AsPC-1 cells (1×10^5 cells/well in a 24-well plate) in the presence of indicated concentrations of Pro A. Cells on the membrane were fixed with 4% PFA and stained with 0.1% crystal violet. Bars: 100 μm. (C) Western blot analysis of epithelial and mesenchymal markers in Panc-1 cells treated with indicated concentrations of Pro A. Each experiment was repeated three times. Error bars represent SEM. *, $P<0.05$; **, $P<0.01$; ***, $P<0.001$; ****, $P<0.0001$. DMSO, dimethyl sulfoxide; PC, pancreatic cancer; Pro A, proscillaridin A; PFA, paraformaldehyde; SEM, standard error of mean.
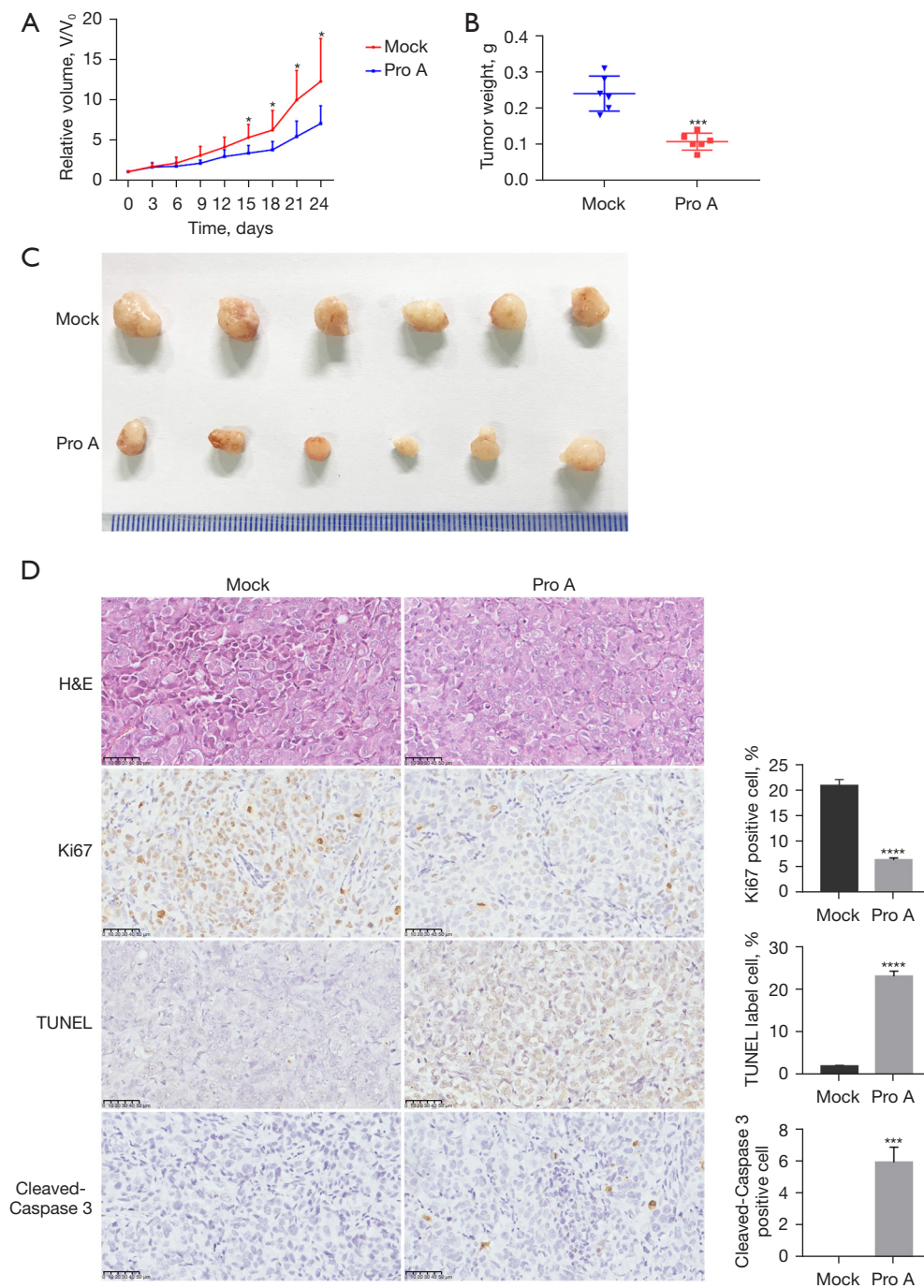
Figure 3 Pro A inhibits Panc-1 xenograft tumor growth in nude mice. Four-week-old female BALB/c nude mice weighing about 16 g were implanted with 5×10^6 Panc-1 cells, randomly divided into two groups (n=6); 7 days later and injected with vehicle (PBS) or Pro A (6.5 mg/kg) every two days for 3 weeks before sacrifice. (A) Tumor volume was measured every 3 days. (B,C) Tumors were surgically removed after sacrifice and weighed. Weights of each removed tumor as well as mean ± SEM for each group are shown. (D) Representative H&E, Ki67, cleaved-Caspase 3, and TUNEL staining of tissue sections from removed tumors. Bars: 50 µm. Error bars represent SEM. *, P<0.05; ***, P<0.001; ****, P<0.0001. Pro A, proscillaridin A; HE, hematoxylin-eosin; PBS, phosphate buffer saline; SEM, standard error of mean.
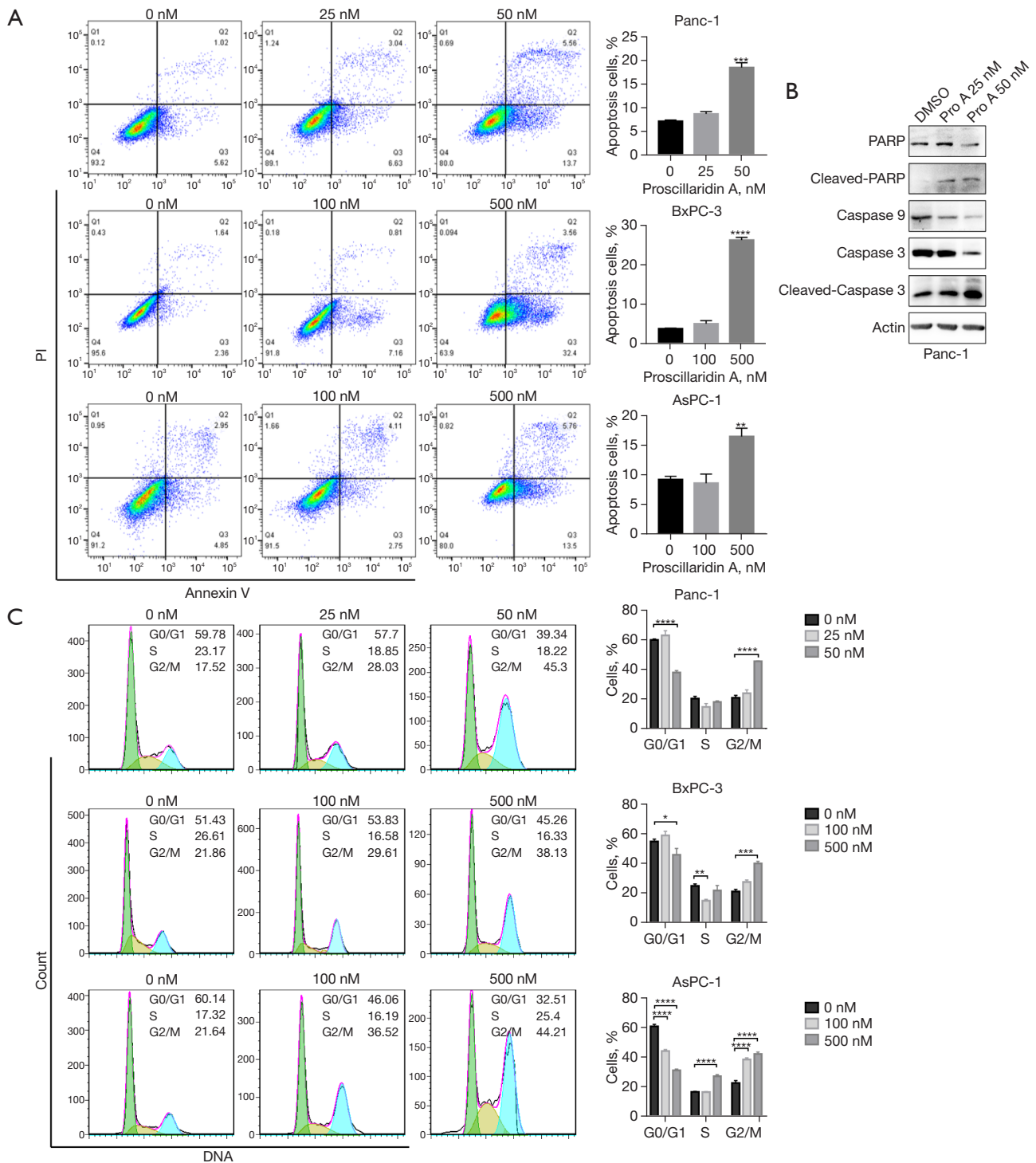
Figure 4 Pro A induces apoptosis and blocked cell cycle progression. Panc-1, BxPC-3, AsPC-1 cells were treated with DMSO or Pro A at the indicated concentration for 48 h. (A) PC cells were subjected to apoptosis analysis using Annexin V/PI double staining followed by flow cytometry. Percentage of early and late apoptotic cells were calculated and displayed on the right. (B) Western blot analysis of apoptotic markers in Pro A-treated Panc-1 cells. (C) Cell cycle analysis by PI staining of PC cells treated with indicated concentrations of Pro A. Each experiment was repeated three times. Error bars represent SEM. *, P<0.05; **, P<0.01; ***, P<0.001; ****, P<0.0001. PC, pancreatic cancer; PI, propidium iodide; DMSO, dimethyl sulfoxide; Pro A, proscillaridin A; SEM, standard error of mean.
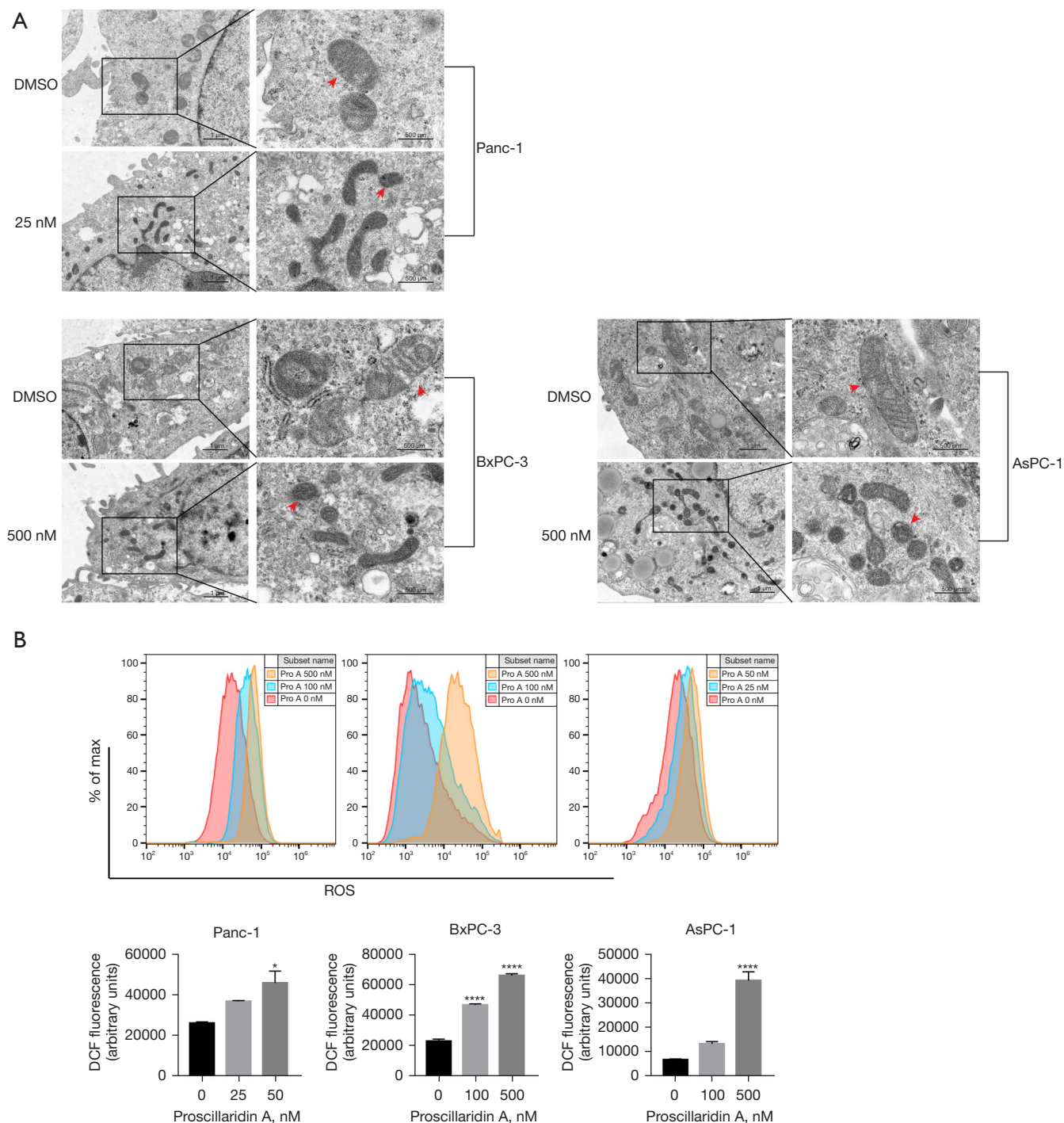
Figure 5 Pro A induces mitochondrial damage and ROS generation in PC cells. (A) Transmission electron micrographs of PC cells with DMSO or Pro A treatment for 48 h. The arrowheads indicate the morphology of mitochondria in different cells. Magnification: ×7,000, ×20,000. (B) PC cells were treated with indicated concentrations of Pro A for 48 h and stained with DCFH-DA to determine intracellular ROS level. Each experiment was repeated three times. Error bars represent SEM. *, P<0.05; ****, P<0.0001. DMSO, dimethyl sulfoxide; PC, pancreatic cancer; Pro A, proscillaridin A; ROS, reactive oxygen species; DCFH-DA, 2',7'-dichlorofluorescein diacetate; SEM, standard error of mean.
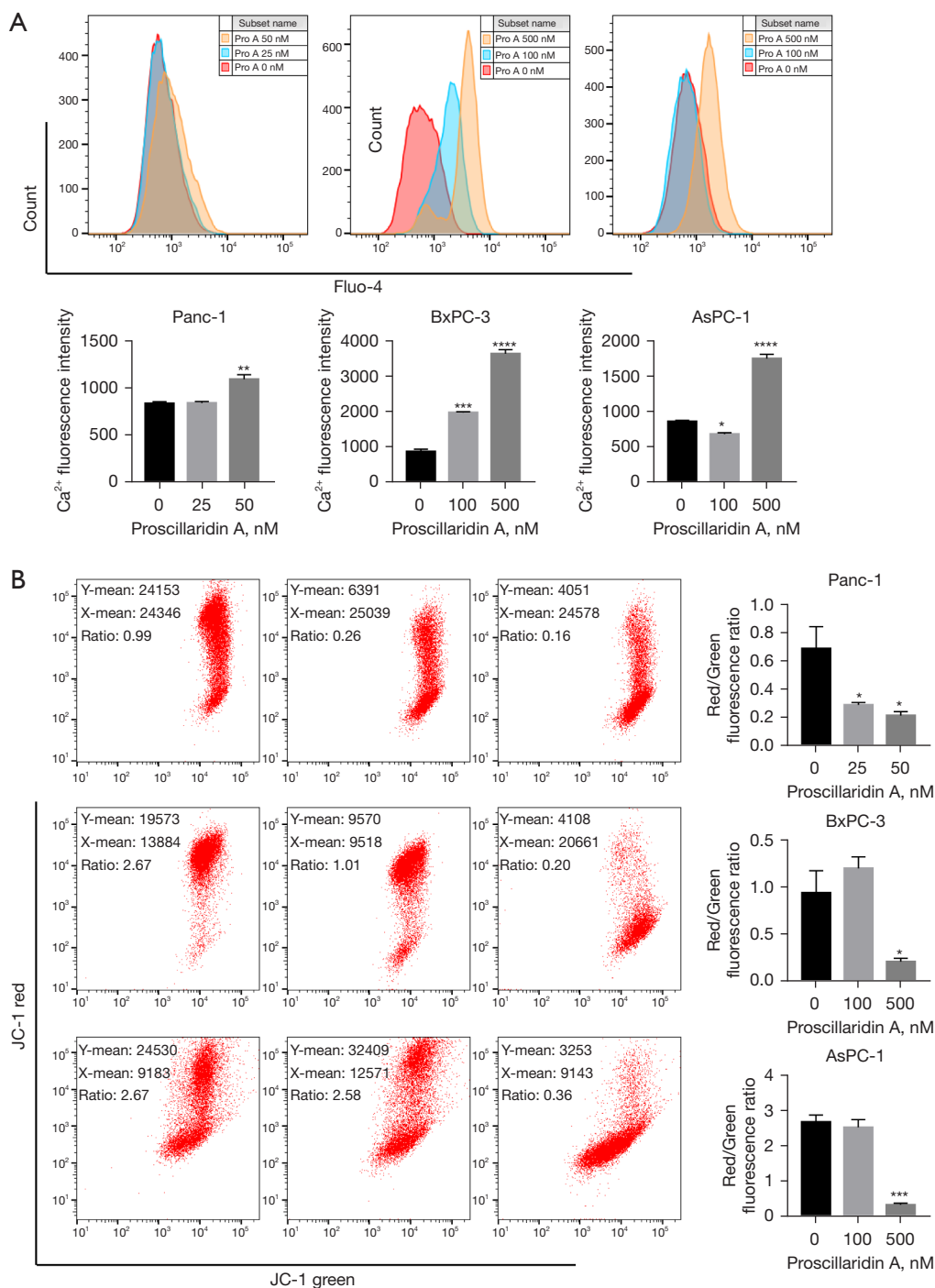
Figure 6 Pro A induces Ca^{2+} accumulation and MMP loss in PC cells. (A) PC cells were treated with indicated concentrations of Pro A for 48 h and stained with Fluo-4 AM to measure intracellular free Ca^{2+}. (B) Cells were treated with indicated concentrations of Pro A for 48 h and stained with JC-1 solution to measure MMP. Each experiment was repeated three times. Error bars represent SEM. *, P<0.05; **, P<0.01; ***, P<0.001; ****, P<0.0001. Pro A, proscillaridin A; PC, pancreatic cancer; MMP, mitochondrial membrane potential; SEM, standard error of mean.

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Pro A induces autophagy in PC cells

Autophagy is a cellular mechanism for coping with stress, defense and recycling. When autophagy occurs, the cleavage of LC3 by ATG4 produces the soluble form LC3-I, which is then bound to the membrane of emerging autophagosomes by ATG3 and ATG7, where LC3-I is transformed to the lipidated form LC3-II.

To determine the role of Pro A in autophagy activation, we used the GFP-LC3-fluorescent indicator system (27). PC cells were transfected with pGFP-LC3 and treated with Pro A for 24 h. GFP-LC3 aggregation and increased LC3-II levels were observed in Pro A-treated PC cells by a fluorescent microscopy and western blot, respectively (*Figure 7A,7B*). mTOR and Raptor, which inhibit autophagy, were down-regulated after 50 nM Pro A treatment in Panc-1 cells (*Figure 7C*). To assess whether autophagy affects Pro A-mediated cytotoxicity, shATG7 was stably transfected into Panc-1 cells to reduce the expression of ATG7. Knockdown of ATG7 expression inhibited the production of LC3-II and the cytotoxicity of Pro A was partially relieved (*Figure 7D,7E*), suggesting that Pro A partially reduces PC cell viability through induction of autophagy.

SMAD4 expression partially determines the sensitivity of PC cells to Pro A

To explore why Panc-1 is more sensitive to Pro A treatment than BxPC-3 and AsPC-1, we used qPCR to detect the expression of important cancer-related genes in the three PC cell lines (*Figure 8A*). Panc-1 cells had significantly higher *SMAD4* mRNA expression than BxPC-3 and AsPC-1 cells, which was confirmed at the protein level (*Figure 8B*). In order to further study the relationship between Pro A's anti-tumor activity and SMAD4 expression, we constructed three stably transfected cell lines, in which *SMAD4* protein was down-regulated in Panc-1 and overexpressed in BxPC-3 and AsPC-1 (Figure S2). The cytotoxicity of Pro A was slightly alleviated with *SMAD4* knockdown in Panc-1 cells, and became slightly aggravated with *SMAD4* overexpressed in BxPC-3 and AsPC-1 cells (*Figure 8C*). The results showed that *SMAD4* expression may be related to the cytotoxicity of Pro A for PC cells.

Interestingly, Pro A treatment also reduced endogenous SMAD4 protein levels in Panc-1 cells in a dose-dependent and time-dependent manner (*Figure 9A,9B*). To understand how Pro A reduces SMAD4 protein expression, we treated Panc-1 cells with cycloheximide (CHX) and measured protein level of SMAD4. The results showed that the SMAD4 protein levels decreased faster in the presence of Pro A, which indicates that Pro A might affect the degradation of SMAD4 protein (*Figure 9C*). Consistent with this observation, SMAD4 protein levels in Panc-1 cells treated with proteasome inhibitor MG132 were almost unaffected by Pro A (*Figure 9D*). These results suggest that Pro A might promote SMAD4 degradation through the ubiquitination-proteasome pathway.

Discussion

CGs are a class of natural products used traditionally to increase cardiac contractility in patients with congestive heart failure and arrhythmia by blocking the Na^+/K^+ ATPase of the heart. Previous studies have reported that CGs can induce cell death of senescent cells and increase the levels of intracellular Ca^{2+} and H^+ to cause anti-tumor effects (28-30). Pro A belongs to the CG family and comes from *Drimia maritima*. Its therapeutic efficacy was first described more than two centuries ago (31). Although it is no longer used clinically, Pro A was reported to have anti-tumor effects on several cancers. Pro A can inhibit STAT3 signaling in breast cancer cells (32) and trigger the activation of endoplasmic reticulum stress to slow the prostate cancer progression (33). Previous study has shown that Pro A at micro-molar concentrations can induce PC cell apoptosis (25). In this study, we found that Pro A at the nano-molar concentrations has cytotoxic effects on PC cells, including inhibiting the proliferation, migration and invasion of PC cells *in vitro* (*Figures 1,2*) and tumor growth in mouse models (*Figure 3*). In addition, Pro A induces cell death by promoting both apoptosis (*Figures 4-6*) and autophagy (*Figure 7*). It is worth noting that Panc-1, BxPC-3 and AsPC-1 cells react differently to the cytotoxicity of Pro A, with Panc-1 being much more sensitive to Pro A (*Table 1*), which may be partly due to SMAD4 expression in Panc-1 (*Figures 8,9*).

SMAD4, which serves as the central mediator of TGF-β signaling, translocates to the nucleus as SMAD2/SMAD3-SMAD4 complex after TGF-β family receptor activation. *SMAD4* and related pathway components are inactivated in ~50% of PC and cooperate with activated $Kras^{G12D}$ to promote PDAC in mouse model (34), but the role of SMAD4-mediated pathway in PC progression has not been elucidated. Cell proliferation was reported to be increased in *SMAD4*-expressing Panc-1 cells and decreased in
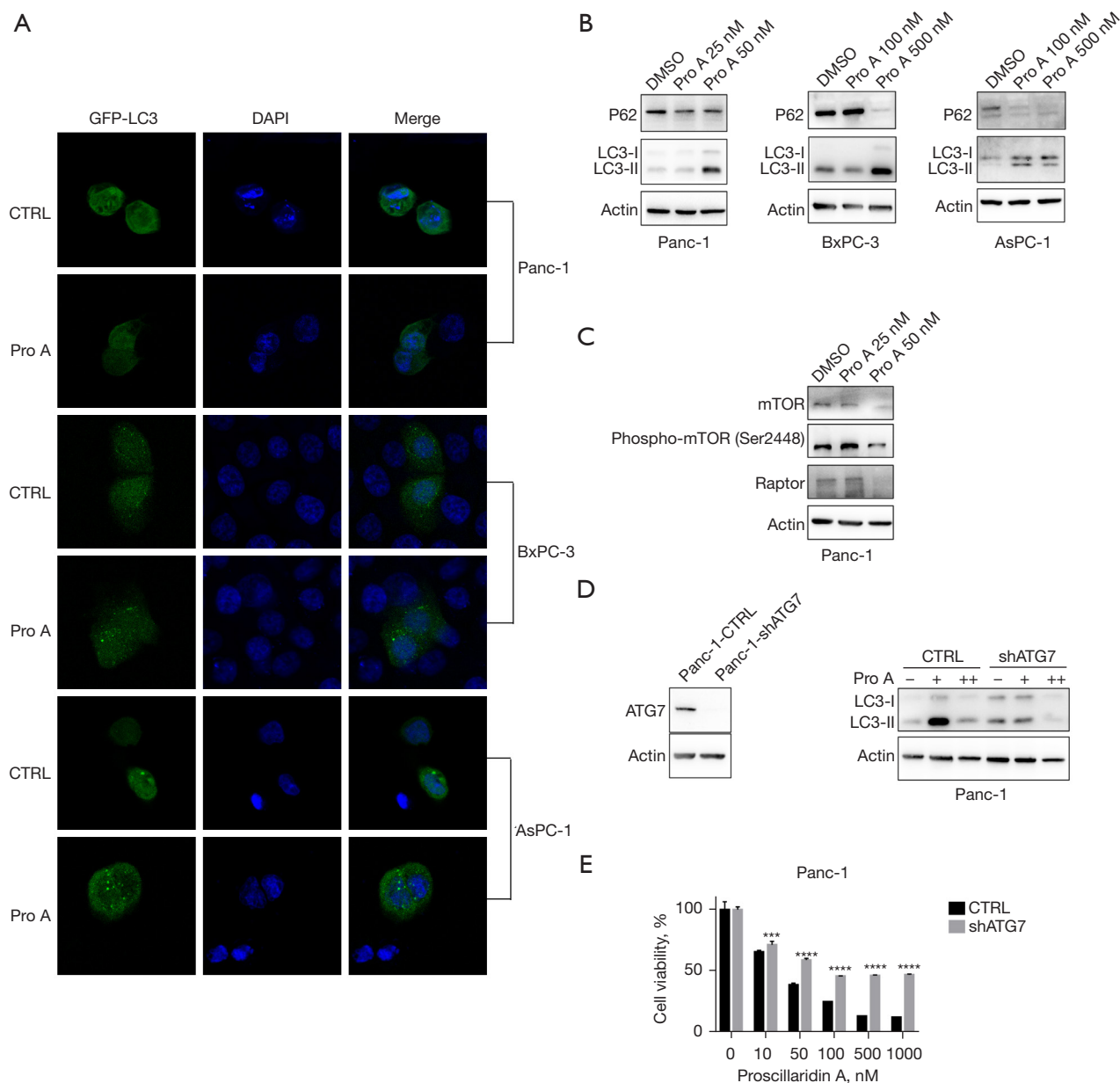
Figure 7 Pro A induces autophagy in PC cells. (A) GFP-LC3 was transfected to three PC cell lines and immunofluorescence of LC3 was observed after treated PC cells with Pro A or DMSO treatment for 24 h. Cells were stained with DAPI. Magnification: ×630. (B) Western blot analysis of p62 and LC3 was performed in PC cells treated with indicated concentration of Pro A for 48 h. (C) Western blot analysis of mTOR pathway associated proteins was performed in Panc-1 cells treated with indicated concentration of Pro A for 48 h. (D) Panc-1 cells were transfected with recombinant lentiviruses expressing shRNA targeting ATG7 or scrambled shRNA control. ATG7 knockdown was confirmed using western blot (left). Western blot analysis of LC3 was performed in shATG7 and shCTRL Panc-1 cells treated with Pro A for 48 h (right). (E) CCK-8 assay was used to determine the effects of ATG7 knockdown on Pro A-induced cytotoxicity in Panc-1 cells treated with indicated concentrations of Pro A for 72 h. Experiment was repeated three times. Error bars represent SEM. ***, $P<0.001$; ****, $P<0.0001$; –, absence; +, presence; ++, higher concentration than "+". DAPI, 4',6-diamidino-2'-phenylindole; Pro A, proscillaridin A; PC, pancreatic cancer; DMSO, dimethyl sulfoxide; CCK-8, cell counting kit-8; SEM, standard error of mean.
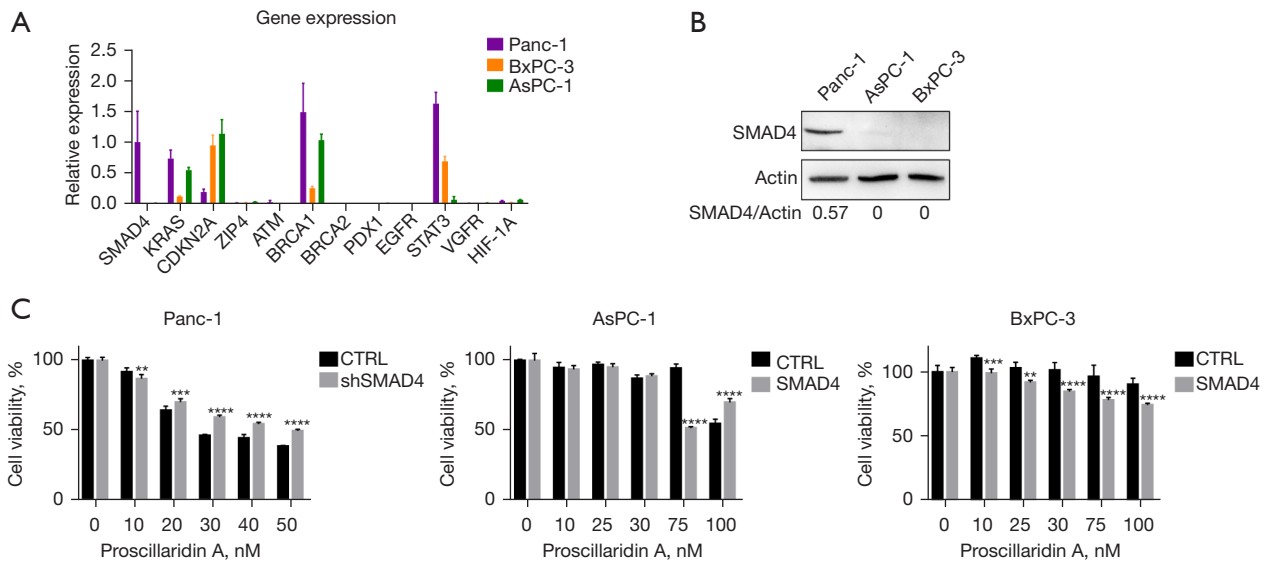
Figure 8 Relation between SMAD4 expression and sensitivity to Pro A of PC cells. (A) qPCR analysis of expression of indicated genes in PC cell lines. (B) Western blot analysis of SMAD4 protein in PC cell lines. (C) CCK-8 assay was used to determine the effects of SMAD4 on Pro A-induced cytotoxicity in PC cell lines. Each experiment was repeated three times. Error bars represent SEM. **, $P<0.01$; ***, $P<0.001$; ****, $P<0.0001$. PC, pancreatic cancer; Pro A, proscillaridin A; qPCR, quantitative polymerase chain reaction; CCK-8, cell counting kit-8; SEM, standard error of mean.
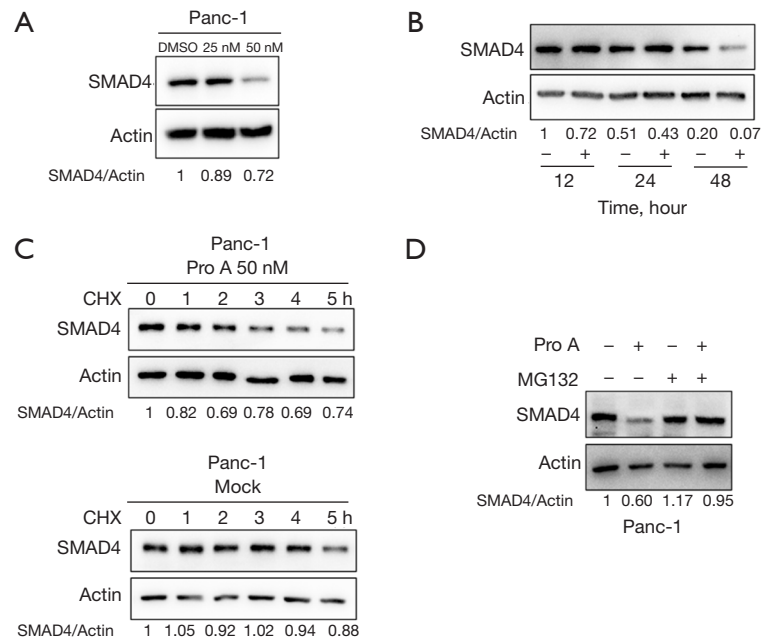
Figure 9 Pro A reduces SMAD4 protein stability in Panc-1 cells. (A) Western blot analysis of SMAD protein level in Panc-1 treated with indicated concentrations of Pro A for 48 h. (B) Western blot analysis of SMAD protein levels in Panc-1 treated with DMSO or Pro A for indicated times. (C) Panc-1 cells were treated with DMSO or Pro A and CHX for the indicated time and SMAD4 protein levels were analyzed by western blot. (D) SMAD4 protein levels in Panc-1 cells treated with DMSO or Pro A and with or without MG132, was determined by western blot analysis. Normalized densitometry scanning results are shown under the blots. –, absence; +, presence. DMSO, dimethyl sulfoxide; Pro A, proscillaridin A; CHX, cycloheximide.

SMAD4-null BxPC-3 cells upon TGFβ-1 stimulation (35). However, another report suggested that inactivation of TGF-β signaling in a mouse model enhanced PC initiation and progression in a SMAD4-dependant manner (36). In this work, Panc-1 displayed the highest sensitivity to Pro A among the three tested PC cell lines (*Figure 1A*). Previous genotyping reports showed that *SMAD4*, also named DPC4 for deleted in pancreatic carcinoma locus 4, is un-mutated in Panc-1, but lost due to homozygous deletion in BxPC-3 (37-39). There have been conflicting reports regarding the status of SMAD4 in AsPC-1 (40,41). Our data from mRNA and protein analyses show that Panc-1 was indeed the only PC cell line tested that displayed endogenous SMAD4 expression, which was related to sensitivity to Pro A, with downregulation of *SMAD4* expression in Panc-1 dampening Pro A's anti-tumor effect (*Figure 8C*). Furthermore, we showed that Pro A reduced the stability of SMAD4 (*Figure 9C,9D*). The wild-type SMAD4 has been shown to mainly undergo mono- and oligo-ubiquitination, which enhances its interaction with other SMADs and results in higher transcriptional activation, instead of poly-ubiquitination, which leads to degradation through proteasome (42). It is plausible that Pro A treatment, through as yet unknown mechanisms, promotes the poly-ubiquitination of SMAD4. On the other hand, there was no significant difference in the cell proliferation between the wild-type and *SMAD4* knockdown Panc-1 cells (Figure S3). These data suggest that SMAD4 expression partially determines the sensitivity of PC cells to Pro A. How SMAD4 impacts on Pro A's anti-tumor activity is currently unclear and warrants further study.

It is widely accepted that mitochondria play a crucial role in cell proliferation and apoptosis. Caspase family proteins and PARP proteins are considered to be the main participants in mitochondrial apoptosis. We found that the expression of cleaved caspase-3 and PARP increased in Pro A-treated PC cells (*Figure 4B*). The production of ROS, the disruption of MMP and the increase in intracellular Ca^{2+} level are the main characteristics of endogenous or mitochondrial apoptosis, which all manifested in PC cells treated with Pro A (*Figures 5,6*). The results confirm the involvement of mitochondrial apoptosis in Pro A's anti-tumor activity against PC cells. It's worth noting that intracellular Ca^{2+} levels in Panc-1 cells treated with cytotoxic concentrations of Pro A increased only moderately compared to the other two cell lines (*Figure 6A*) and this apparent discordance is more likely a result of differences between these cells in other mechanisms affecting Ca^{2+}

levels, for instance endoplasmic reticulum (ER) sequestration and plasma membrane Ca^{2+} pump(s). Moreover, it is indeed possible that different sensitivities to Pro A between the three PC cell lines used here might also be related to differences in Na^+/K^+ ATPase expression, its pump activity and downstream signaling.

Furthermore, our data demonstrate that Pro A not only induces apoptosis, but also induces a certain degree of autophagy. Autophagy is a major intracellular degradation system, which has opposing, context-dependent roles in cancers. Both stimulation and inhibition of autophagy have been proposed as cancer treatment (43). CGs have been shown to induce autophagy in some cancer types (44-46). Here we show for the first time that Pro A induces autophagy in PC cells, characterized by the changes in the expression of autophagy markers, LC3-I/II, p62 and mTOR pathway (*Figure 7*). In addition, ATG7-deficient Panc-1 cells showed reduced sensitivity to Pro A treatment. These data suggest that autophagy may also contribute to the anti-tumor activity of Pro A on PC cells.

In conclusion, our findings provide insights into the anti-tumor effect of Pro A on PC cells and a starting point on elucidation of the molecular mechanism of Pro A's differential cytotoxic effects on different PC cells, which will lay the basis for Pro A and similar molecules to be tested as novel drugs for PC treatment. More in-depth work is required to identify additional traits associated with sensitivity to Pro A, preferably using primary PC tumor cells, which would hopefully lead to the establishment of a practical subtyping criteria for screening candidates for initial testing of Pro A in human PC.

Acknowledgments

Footnote

Reporting Checklist: The authors have completed the ARRIVE reporting checklist. Available at https://atm.amegroups.com/article/view/10.21037/atm-22-1085/rc

Data Sharing Statement: Available at https://atm.amegroups.com/article/view/10.21037/atm-22-1085/dss

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Conflicts of Interest: All authors have completed the ICMJE uniform disclosure form (available at https://atm.amegroups.com/article/view/10.21037/atm-22-1085/coif). 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. Experiments were performed under a project license (No. 202103005S) granted by the Animal Ethics Committee of School of Basic Medical Sciences, Fudan University and were performed in accordance with the guidelines of the National Institutes of Health on the care and use of laboratory animals.

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