

Metformin suppresses the invasive ability of pancreatic cancer cells by blocking autocrine TGF- β 1 signaling

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Received January 20, 2018; Accepted June 18, 2018

DOI: 10.3892/or.2018.6518

Abstract. Pancreatic ductal adenocarcinoma (PDAC) is a highly invasive neoplasm with a 5-year survival rate of <8%. Metformin, the most widely used antidiabetic drug in the world, has been shown to exert anticancer activities in epidemiological and animal studies. Our previous studies revealed that metformin suppressed desmoplasia in PDAC by reducing TGF- β 1 production in cancer cells. The aim of the present study was to investigate the effects of metformin on invasion and epithelial-mesenchymal transition (EMT) in pancreatic cancer and to reveal the underlying mechanisms. In the present study, we revealed that metformin suppressed migration, invasion and EMT changes in pancreatic cancer cells. Furthermore, metformin reduced TGF- β 1 production and Smad2/3 phosphorylation in pancreatic cancer cells. In addition, treatment with recombinant TGF- β 1 recovered the metformin-mediated invasion inhibition and EMT changes. Treatment with metformin also suppressed tumor growth, invasion and EMT in *LSL-Kras^{G12D/+}*, *Trp53^{fl/+}* and *Pdx1-Cre* (KPC) transgenic mice that harbor spontaneous pancreatic cancer. Collectively, our study revealed a new possible mechanism for the antitumor effects of metformin via autocrine TGF- β 1/Smad2/3 signaling in PDAC.

Introduction

Pancreatic ductal adenocarcinoma (PDAC) is one of the most aggressive forms of human pancreatic cancer. Although in-depth knowledge of the disease has been elucidated and an extensive amount of research has been conducted in the past few decades, the effectiveness of treatment for this malignancy

has not substantially improved. PDAC remains the fourth leading cause of cancer-related mortality in the United States, with a 5-year survival rate of <8% (1). PDAC may go undetected until patients have progressed to an advanced stage. More than 80% of patients have already missed the opportunity for radical surgery at the time of initial diagnosis (2). Due to the aggressive nature of this malignancy and its non-sensitivity to radiotherapy and chemotherapy, local recurrence and distant metastasis after initial 'curative' resection is an unresolved clinical problem for PDAC patients (3).

Active invasion and metastasis have been recognized as hallmarks of cancer (4). Accumulating evidence has indicated that epithelial-mesenchymal transition (EMT), which initiates an invasion-metastasis cascade, is a feature of aggressive tumors (5,6). During EMT, cancer cells disassemble their epithelial junctions and suppress the expression of junctional proteins. Consequently, the cancer cells acquire the abilities to invade, resist apoptosis and disseminate (7). E-cadherin, a component of adherent junctions, is lost during EMT and cancer progression. Conversely, another adherent junction protein, N-cadherin is upregulated in cancer cells. A previous study revealed that cadherin switching was a main event in EMT and was necessary for the increased motility of cancer cells (8). Accumulating evidence has revealed that EMT plays a crucial role in the invasion and metastasis of pancreatic cancer (9,10). Recently, several studies have revealed that many growth factors and cytokines, as well as cellular signaling pathways, could induce EMT. One such cytokine is transforming growth factor- β 1 (TGF- β 1) (11).

Compared with other treatments, metformin, the first-line treatment for managing type 2 diabetes mellitus, has been associated with reduced cancer burden in epidemiological studies in diabetic patients (12,13). There is also evidence that metformin use is associated with a reduced risk of prostate (14), colon (15) and pancreatic cancer (16). Given the anticancer properties and the safety profile of metformin, this treatment has attracted great interest from cancer researchers worldwide (17). In the past, exciting preclinical studies have shown that metformin can inhibit cancer cell growth both *in vitro* and *in vivo* (18-20). Using the *LSL-Kras^{G12D/+}*, *Trp53^{fl/+}* and *Pdx1-Cre* (KPC) mouse models, our previous study indicated that metformin suppressed the initiation and progression of pancreatic cancer (21). In another study (22), our data revealed

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Key words: metformin, TGF- β 1, epithelial-mesenchymal transition, metastasis, pancreatic ductal adenocarcinoma, *LSL-Kras^{G12D/+}*, *Trp53^{fl/+}*, *Pdx1-Cre* (KPC) transgenic mice

that metformin-mediated AMPK activation in cancer cells inhibited pancreatic cancer progression by suppressing the desmoplastic reaction in tumor tissues. Further mechanistic experiments revealed that metformin reduced TGF- β 1 production in cancer cells, thus suppressing the TGF- β 1-induced activation of pancreatic stellate cells (PSCs) (22). Based on these observations, we hypothesized that metformin could inhibit the invasion and metastasis of pancreatic cancer cells by blocking the autocrine TGF- β 1 signaling. In the present study, we used the KPC transgenic mouse model of pancreatic cancer to demonstrate that metformin inhibited the invasion and metastasis of pancreatic cancer by suppressing TGF- β 1/Smad2/3 signaling.

Materials and methods

Cell culture and reagents. Human PDAC cell lines Panc-1 and BxPC-3 were purchased from The Chinese Academy of Sciences Cell Bank of Type Culture Collection (CBTCCAS; Shanghai, China). The cells were maintained in the appropriate medium (DMEM; HyClone Laboratories, Logan, UT, USA) for Panc-1 and RPMI-1640 medium for BxPC-3 (HyClone Laboratories) supplemented with 10% fetal bovine serum (FBS; HyClone Laboratories; GE Healthcare Life Sciences, Logan, UT, USA) and 1% penicillin/streptomycin in a humidified atmosphere containing 5% CO₂ at 37°C. Primary antibodies for E-cadherin (1:1,000; cat. no. 3195), N-cadherin (1:800; cat. no. 13116), MMP-2 (1:1,000; cat. no. 40994) and vimentin (1:1,000; cat. no. 5741) were purchased from Cell Signaling Technology, Inc. (Danvers, MA, USA). Antibodies against β -actin (1:1,000; cat. no. A5441) were obtained from Sigma-Aldrich; Merck (St. Louis, MO, USA). Primary antibodies for TGF- β 1 (1:800; cat. no. ab92486), Smad2/3 (1:1,000; cat. no. ab202445) and p-Smad2/3 (1:1,000; cat. no. ab63399) were purchased from Abcam (Cambridge, MA, USA) and anti- α -SMA (1:800; cat. no. A03744) was obtained from Boster Biological Technology (Wuhan, China). All secondary antibodies (goat anti-rabbit IgG-HRP (1:10,000; cat. no. ssc-2004) and goat anti-mouse IgG-HRP (1:10,000; cat. no. sc-2005) were purchased from Santa Cruz Biotechnology (Santa Cruz, CA, USA). Recombinant TGF- β 1 (T7039) was obtained from Sigma-Aldrich. Metformin (Sigma-Aldrich; Merck) was dissolved in phosphate-buffered saline (PBS) as a stock solution of 100 mM. Working dilutions of metformin were made in culture medium immediately before use.

Wound healing assays. Wound healing assays were conducted to assess the migration ability of cancer cells. Briefly, cancer cells were seeded in 6-well plates and when they almost covered the well, the cells were serum-starved and then treated with metformin (2 mM) for 24 h. Subsequently, the cells were scratched using a 200-ml sterile pipette tip. Images of the matched-pair wound regions at 0 and 24 h were obtained at a magnification of x100 using a light microscope (Nikon Instruments, Inc., Tokyo, Japan).

Matrigel invasion assay. Matrigel invasion assays were performed to investigate the invasive ability of cancer cells. In brief, cancer cells were serum-starved overnight and

then pretreated with or without metformin plus TGF- β 1 (2 ng/ml) for 24 h. Subsequently, the cells were trypsinized, and 5x10⁴ cells and 200 μ l of serum-free medium were added to Matrigel-coated inserts (BD Biosciences, Franklin Lakes, NJ, USA). Complete medium (500 μ l) was added to the lower chamber. Following incubation for 48 h, the invaded cells were stained with 0.1% crystal violet solution for 15 min at room temperature, and 10 randomly selected fields were photographed with a light microscope (Nikon Instruments, Inc.) at a magnification of x200.

Immunofluorescence analysis. Following the designated treatment, immunohistochemical analyses were performed according to our previously described protocol (23). Images were pseudocolored using a fluorescence microscope (Nikon Eclipse Ti-s; Nikon Instruments, Inc.) with the appropriate excitation and emission (blue, 440–450 nm; green, 530–550 nm; red, 630–650 nm) spectra at a magnification of x400. Image-Pro Plus software (version 6.0; Media Cybernetics Inc., Rockville, MD, USA) was used for further analysis.

Western blotting. Total cellular protein was extracted using RIPA lysis buffer (Beyotime Institute of Biotechnology, Guangzhou, China). Protein concentrations were analyzed with a BCA protein assay kit (Beyotime Institute of Biotechnology) according to the manufacturer's instructions. The same amounts of protein (150 μ g) from the samples were separated by 10% SDS-PAGE and then transferred onto polyvinylidene difluoride (PVDF) membranes as previously described (24). The membranes were blocked with 5% non-fat dry milk in TBST [10 mM Tris-HCl (pH 8.0), 150 mM NaCl, 0.05% Tween-20]. Then membranes were incubated with primary antibodies overnight at 4°C. After three washes of 10 min each in TBST, the membranes were incubated with HRP-conjugated secondary antibodies for 1 h and subsequently washed again. Protein expression was visualized using enhanced chemiluminescence (EMD Millipore, Billerica, MA, USA) and the ChemiDoc XRS imaging system (Bio-Rad Laboratories, Inc., Hercules, CA, USA). β -actin was used to ensure equivalent protein loading.

In vivo study. All experimental protocols were approved by the Ethical Committee of The First Affiliated Hospital of Medical College, Xi'an Jiaotong University (Xi'an, China). *LSL-Kras^{G12D/+}; Trp53^{fl/+}; Pdx1-Cre* (KPC) mice were generated and maintained as previously described (21). A total of 20, 6-week-old mice (female; average weight, 20 \pm 1.79 g) were divided into two groups of 10 mice each. Mice in the metformin group were gavaged with vehicle (sterile water) or metformin (200 mg/kg) daily for 4 weeks. All mice were housed under pathogen-free conditions with a 12:12-h dark/light cycle under constant temperature (24°C). At the termination of the experiment, the mice were euthanized by 30% CO₂ asphyxiation, followed by cervical dislocation. Tissues were harvested and fixed in formalin. Tumor samples were prepared for further histological analysis.

Statistical analysis. Each experiment was performed at least three times. All quantitative data were analyzed using SPSS version 15.0 (SPSS, Inc., Chicago, IL, USA) and are expressed

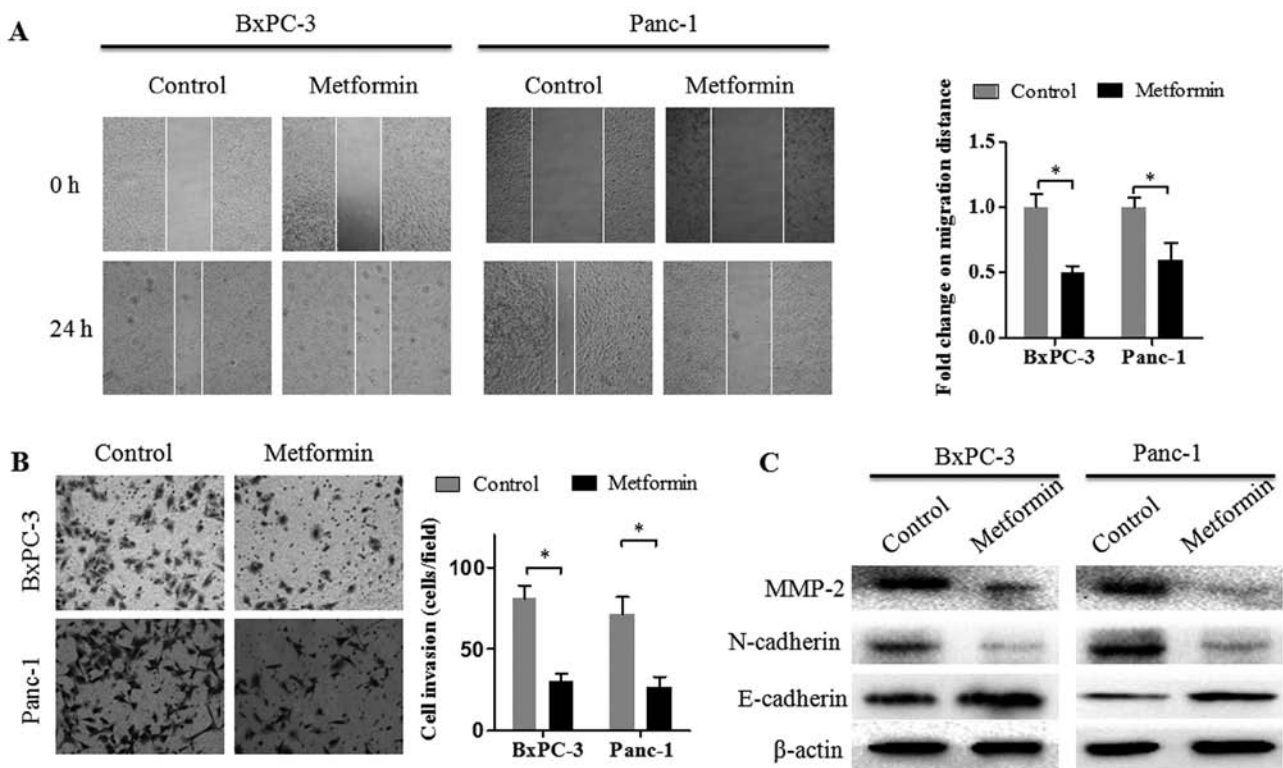


Figure 1. Metformin inhibits the migration and invasion of pancreatic cancer cells. (A) Wound healing assays were performed to assess the migration ability of Panc-1 and BxPC-3 cells following 2 mM metformin treatment. Images were acquired at 0 and 24 h at a magnification of x100; * $P < 0.05$. (B) Panc-1 and BxPC-3 cells were treated with 2 mM metformin for 24 h. Cell invasion ability was evaluated by Matrigel invasion assays. Error bars display the mean \pm SD of three independent experiments. * $P < 0.05$, magnification, x200. (C) Western blot assays were performed to evaluate the effects of metformin on the EMT markers E-cadherin and N-cadherin and the invasion-related molecule MMP-2.

as the mean \pm standard deviation (SD). Two-tailed unpaired Student's t-tests were used to analyze the data between groups. $P < 0.05$ was considered to indicate a statistically significant difference.

Results

Metformin attenuates the migration and invasion abilities of pancreatic cancer cells. First, the effect of metformin on the migration ability of pancreatic cancer cells was assessed by wound healing assays. Similarly, with our previous study, 2 mM metformin was also used in the present study (22). As displayed in Fig. 1A, the migration ability of pancreatic cancer cells was significantly suppressed by 2 mM metformin treatment. To investigate the effect of metformin on the invasion ability of pancreatic cancer cells, Matrigel invasion assays were performed. The results revealed that the invasion ability of pancreatic cancer cells was decreased by 2 mM metformin treatment (Fig. 1B). These results indicated that metformin inhibited the invasion and migration capacities of pancreatic cancer cells *in vitro*. Available evidence has indicated that active EMT in cancer cells enhances their invasion and metastasis abilities. To explore whether metformin has an effect on EMT-related molecules, the protein expression levels of E-cadherin, N-cadherin and MMP-2 were detected in pancreatic cancer cells following exposure to 2 mM metformin for 48 h. The western blotting results revealed that metformin treatment decreased the protein expression levels of N-cadherin and MMP-2 and increased the protein

expression levels of E-cadherin in both BxPC-3 and Panc-1 cells (Fig. 1C).

Metformin suppresses TGF- β 1/Smad2/3 signaling in pancreatic cancer cells. Canonical TGF- β 1/Smad2/3 signaling has been revealed to play an important role in EMT in various epithelial cells (11). Our previous study (22) revealed that metformin inhibited TGF- β 1 production in pancreatic cancer cells by inducing AMPK activation. To further explore the effects of metformin on TGF- β 1/SMAD2/3 signaling in pancreatic cancer cells, BxPC-3 and Panc-1 cells were treated with 2 mM metformin. Following 48 h, total cell protein was extracted and subjected to western blot analysis. As anticipated, the protein expression levels of TGF- β 1 were decreased following metformin treatment. Additionally, the protein expression levels of p-Smad2/3 were also decreased by metformin treatment in both BxPC-3 and Panc-1 cells (Fig. 2A). In addition, this phenomenon was confirmed by immunofluorescence assays (Fig. 2B). The results revealed that the expression levels of TGF- β 1 and p-Smad2/3 were decreased by metformin treatment in pancreatic cancer cells. Notably, p-Smad2/3 was observed primarily in the cytoplasm following metformin treatment. In a subcutaneous tumor model, the immunofluorescence data revealed that TGF- β 1 staining in tumor tissue, as well as α -SMA expression, was reduced by the oral administration of 200 mg/kg metformin (Fig. 2C). These results were consistent with those of our previous study (22). These findings revealed that metformin suppressed TGF- β 1/Smad2/3 signaling in pancreatic cancer cells.

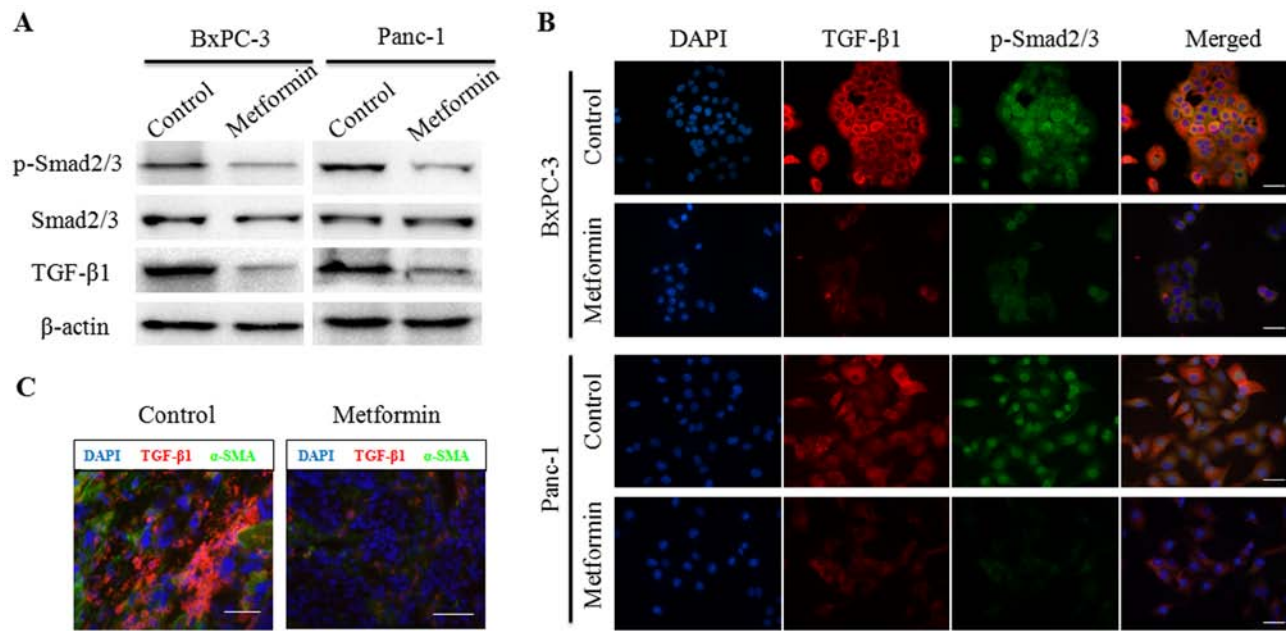


Figure 2. Metformin suppresses TGF- β 1 production and TGF- β 1/Smad2/3 signaling in pancreatic cancer cells. (A) Panc-1 and BxPC-3 cells were treated with 2 mM metformin for 48 h and then, total cell protein samples were prepared and analyzed by immunoblotting with antibodies against TGF- β 1, Smad2/3 and p-Smad2/3. (B) Immunofluorescence analyses were conducted to detect TGF- β 1 (red) and p-Smad2/3 (green) expression in Panc-1 and BxPC-3 cells following 2 mM metformin treatment for 48 h, magnification, x200. (C) Representative images displaying the immunofluorescence results of TGF- β 1 and α -SMA staining of tumor tissue from a subcutaneous tumor model after the mice were orally administered 200 mg/kg metformin for 4 weeks. Scale bar, 50 μ m.

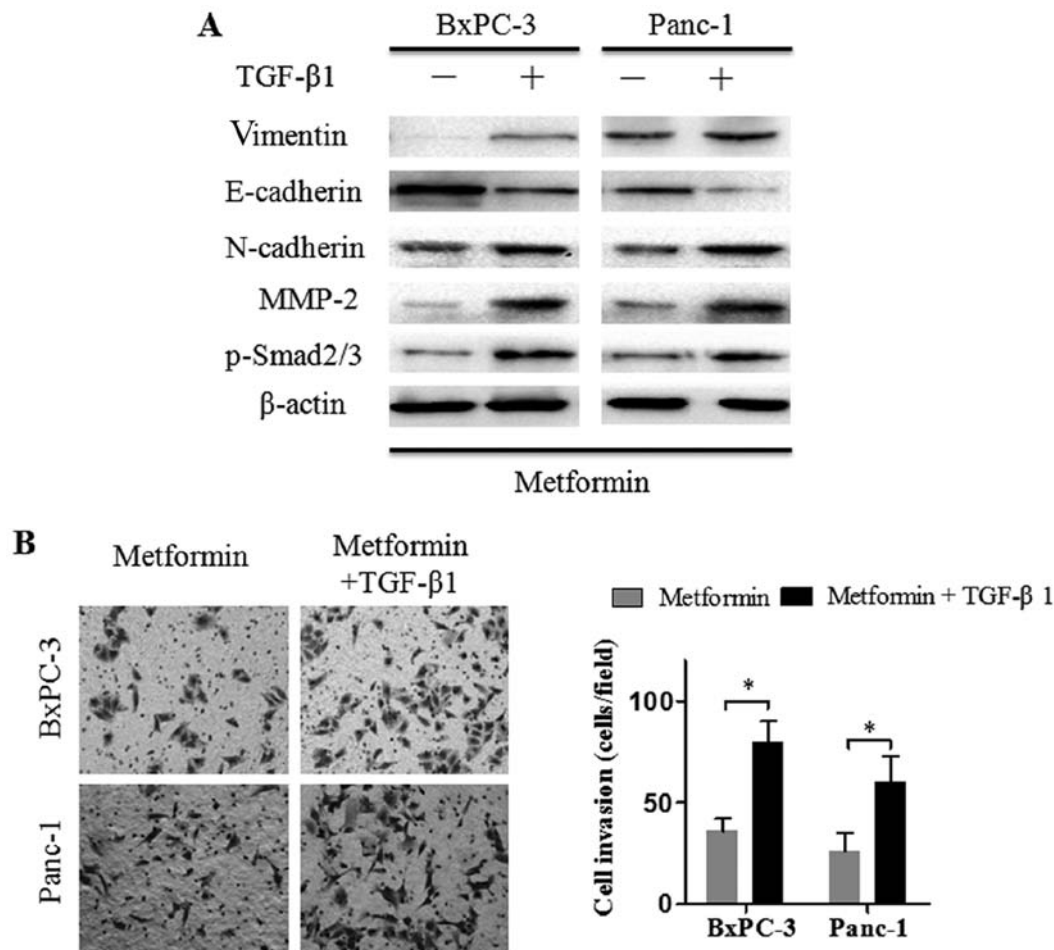


Figure 3. Recombinant TGF- β 1 reverses the metformin-induced EMT changes and invasion suppression in pancreatic cancer cells. (A) The expression levels of E-cadherin, N-cadherin, vimentin, MMP-2 and p-Smad2/3 were assessed by western blot assays after Panc-1 and BxPC-3 cells were treated with metformin (2 mM) or metformin plus TGF- β 1 (2 ng/ml) for 48 h. (B) Matrigel invasion assays were performed to investigate the invasion ability of pancreatic cancer cells following treatment with metformin or metformin plus TGF- β 1 for 24 h. Images are representative of three independent experiments. * P <0.05, magnification, x200.

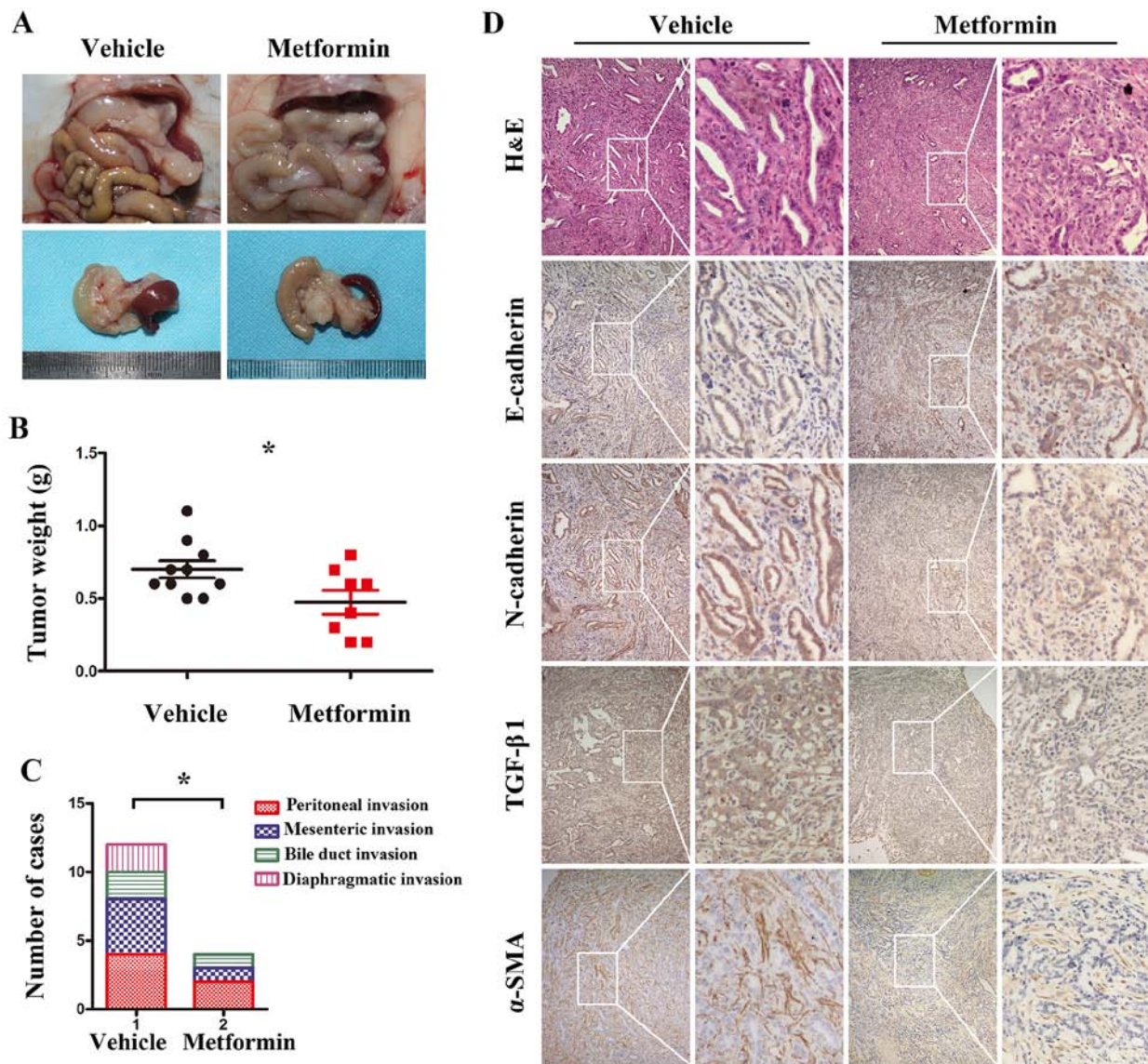


Figure 4. Metformin suppresses the growth, invasion and EMT of pancreatic cancer in a KPC transgenic mouse model. (A) Representative images displaying the macroscopic pancreatic tissue specimens from KPC mice treated with vehicle or metformin. (B) Tumor weights were quantified in mice treated with vehicle or metformin at the end of the experiment; $P < 0.05$. (C) The occurrence of abdominal invasion (peritoneal invasion, mesenteric invasion, bile duct invasion and diaphragmatic invasion) was analyzed in KPC mice treated with vehicle or metformin; $P < 0.05$. (D) Representative images of hematoxylin and eosin (H&E) staining and immunohistochemical staining for E-cadherin, N-cadherin, TGF- β 1 and α -SMA in tumor tissues from KPC mice treated with vehicle or metformin.

Exogenous TGF- β 1 reverses the effects of metformin on the invasion ability of pancreatic cancer cells. The aforementioned observations revealed that metformin suppressed the invasion ability and TGF- β 1/Smad2/3 signaling in pancreatic cancer cells. To ascertain that TGF- β 1 signaling inhibition attenuated cancer cell invasion, Panc-1 and BxPC-3 cells were treated with metformin (2 mM) alone or metformin plus recombinant TGF- β 1 (2 ng/ml) for 24 h and then, EMT- and invasion-related markers were detected by western blot assays. The results revealed that the addition of recombinant TGF- β 1 to the culture medium induced Smad2/3 phosphorylation (Fig. 3A). Furthermore, N-cadherin, vimentin, and MMP-2 protein expression levels were increased, and E-cadherin protein expression levels were decreased in Panc-1 and BxPC-3 cells treated with metformin plus TGF- β 1, compared to cells treated with metformin alone. Matrigel invasion assays revealed that the metformin-mediated

suppression of cancer cell invasion was almost recovered in the presence of TGF- β 1. Collectively, these observations indicated that TGF- β 1 downregulation may be responsible for the metformin-mediated invasion and EMT changes in pancreatic cancer cells.

Metformin suppresses pancreatic cancer tumor growth and metastasis in KPC transgenic mice. The KPC model is a well-validated, clinically relevant model of PDAC that recapitulates the spectrum of pancreatic cancer from pancreatic intraepithelial neoplasia (PanIN) to invasive PDAC (21). To determine the effects of metformin *in vivo*, KPC transgenic mice were generated and treated orally with vehicle (sterile water, daily for 4 weeks) or metformin (200 mg/kg, daily for 4 weeks). At the end of the experiment, KPC mice were sacrificed, and the tumor samples were prepared (Fig. 4A).

The results revealed that the average tumor weight in the metformin-treated group was lower than that in the vehicle-treated group (Fig. 4B). In addition, compared with vehicle treatment, metformin treatment significantly decreased the incidence of abdominal invasion, including peritoneal invasion, mesenteric invasion and bile duct invasion as well as diaphragmatic invasion (Fig. 4C). The immunohistochemistry results (Fig. 4D) revealed that the TGF- β 1 immunoreactivity in the metformin-treated group was lower than that in the vehicle-treated group. In addition, increased E-cadherin and decreased N-cadherin expression levels were observed in the metformin-treated group. Furthermore, compared with vehicle treatment, metformin treatment reduced the area of α -SMA-positive staining. Collectively, these results indicated that metformin inhibited pancreatic cancer tumor growth, EMT and metastasis *in vivo*.

Discussion

PDAC is typically associated with a poor prognosis even after curative resection with postoperative adjuvant chemotherapy. Despite advances in our understanding of the molecular and genetic mechanisms of pancreatic cancer, managing the disease remains a clinical challenge due to its poor response to most chemotherapeutic agents (25). Local recurrence and metastasis are the primary causes of treatment failure in cancer patients and of cancer-related deaths. Recently, mounting evidence from both observational and laboratory studies indicated that metformin treatment may be associated with a decreased risk of developing cancer and a better response to chemotherapy (12,26). The potential mechanisms underlying the antitumor properties of metformin remain unclear. Our previous studies have indicated that metformin can block the interaction between cancer cells and PSCs by reducing cancer cell-derived cytokines (22). In the present study, we found that metformin treatment significantly attenuated the migration and invasion properties of pancreatic cancer cells. In addition, metformin suppressed EMT in pancreatic cancer cells. This outcome was also produced in KPC transgenic mice. Furthermore, our data revealed that these effects of metformin occurred in part via the downregulation of TGF- β 1/Smad2/3 signaling in pancreatic cancer.

TGF- β 1 is a potent cytokine with marked functionalities, including the regulation of cell apoptosis, proliferation, differentiation and extracellular matrix production (27). TGF- β 1 binding to its receptor results in the phosphorylation and activation of the transcription factors Smad2/3. Once activated, Smad2/3 translocates into the nucleus to govern gene transcription. The role of TGF- β 1 during cancer initiation and progression is complex and paradoxical. TGF- β 1 functions as a tumor suppressor in normal and early-stage cancers by inducing cell-cycle arrest and apoptotic reactions and as a tumor promoter in late-stage cancers by promoting cancer growth, invasion and metastasis (28,29). Parallel studies have suggested that TGF- β 1 is a potent inducer of EMT (11,30). Adding TGF- β 1 to epithelial cells in culture is a convenient way to induce EMT in various epithelial cells (31). During EMT induction, TGF- β 1 rapidly activates PI3K, Akt, mTOR complex 1 (mTORC1) and S6 kinase, leading to increases in protein synthesis, cell motility and invasion (32). A previous

study has indicated that TGF- β 1 acted in an autocrine manner to enhance tumor cell invasion in pancreatic cancer by upregulating MMP-2 (33). In another study, a significant association was found between the expression of TGF- β 1 and lymph node involvement and the depth of invasion in pancreatic cancer (34). In the present study, we found that TGF- β 1 expression in pancreatic cancer cells was decreased by metformin. Notably, metformin treatment inhibited TGF- β 1-induced Smad2/3 phosphorylation in pancreatic cancer cells.

The desmoplastic reaction, which is a result of the proliferation of activated PSCs and the increased deposition of extracellular matrix (ECM) components, is a prominent pathological characteristic of pancreatic cancer (35). Accumulating evidence has indicated that the desmoplastic reaction in pancreatic cancer contributes to the aggressive nature of this malignancy by fostering tumor growth and metastatic spread, as well as enhancing chemoradiotherapy resistance (36–39). Thus, strategies targeting cancer-stroma interactions may serve as a potential approach for pancreatic cancer treatment (8). In pancreatic cancer, PSCs are activated by tumor-stromal interactions, including direct contact with pancreatic cancer cells and paracrine growth factors, such as TGF- β 1, secreted from pancreatic cancer cells (40,41). It has recently been reported that metformin reduces desmoplasia in pancreatic cancer by reducing the expression of inflammatory cytokines, resulting in reduced disease progression (42). Our previous study also indicated that metformin suppressed desmoplasia in pancreatic cancer by reducing TGF- β 1 production in pancreatic cancer cells and inhibiting paracrine-mediated PSC activation both *in vitro* and *in vivo* (22). The anti-stromal behavior of metformin has been previously verified in a genetically engineered mouse model of pancreatic cancer (21). In the present study, we also observed that α -SMA-positive cells were markedly reduced in cancer tissues from KPC mice following metformin treatment. Collectively, this evidence indicated the anti-stromal properties of metformin, and that reduced TGF- β 1 production by cancer cells may be the underlying mechanism.

In the past, we conducted a series of studies on the anti-cancer effect of metformin on pancreatic cancer. Our previous study revealed that metformin suppressed desmoplastic reaction in pancreatic cancer by reducing TGF- β 1 production in cancer cells and suppressing paracrine TGF- β 1-induced PSC activation (22). In addition, both our *in vitro* and *in vivo* experiments revealed that metformin inhibited the proliferation, invasion and migration of pancreatic cancer cells (22,43). Furthermore, by using a genetic mouse model of pancreatic cancer, our results confirmed the inhibitory effect of metformin on pancreatic cancer initiation and progression (21). However, whether the inhibitory effect of metformin on tumor invasion and metastasis is mediated by targeting the autocrine TGF- β 1/Smad pathway remains unknown. Based on the aforementioned research, in the present study, we investigated the effect of metformin on TGF- β 1/Smad in pancreatic cancer cells. Our results indicated that blocking autocrine TGF- β 1 signaling was partially responsible for metformin-reduced cell invasion properties. Therefore, the present study is a confirmation and supplement to our previous studies.

In conclusion, the results from the present study indicated that metformin suppressed the invasion, EMT and metastasis

of pancreatic cancer by preventing autocrine TGF- β 1/Smad2/3 signaling both *in vitro* and *in vivo*. Thus, our study revealed a new possible mechanism for the antitumor effects of metformin.

Acknowledgements

Not applicable.

Funding

The present study was supported by grants from the National Natural Science Foundation of China (nos. 81402971, 81672434 and 81702916).

Availability of data and materials

The datasets used during the present study are available from the corresponding author upon reasonable request.

Authors' contributions

QM and WD designed the experiments. WQ, CZ and JC performed the majority of the experiments. TQ, YX, LC and JL analyzed the data. KC and XL organized the figures. JM wrote the manuscript. KC, XL and JM were also involved in the conception of the study. QM and WD reviewed it. All authors read and approved the manuscript and agree to be accountable for all aspects of the research in ensuring that the accuracy or integrity of any part of the work are appropriately investigated and resolved.

Ethics approval and consent to participate

All experimental protocols were authorized by the Ethics Committee of the First Affiliated Hospital of Medical College, Xi'an Jiaotong University (Xi'an, China). The protocols also complied with the Declaration of Helsinki.

Patient consent for publication

Not applicable.

Competing interests

The authors declare that they have no competing interests.

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