

Metformin Increases the Chemosensitivity of Pancreatic Cancer Cells to Gemcitabine by Reversing EMT Through Regulation DNA Methylation of miR-663

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Background: Pancreatic cancer is a devastating malignancy with poor prognosis. Metformin, a classic anti-diabetes drug, seems to improve survival of pancreatic cancer patients in some studies.

Methods: Cell counting kit-8 assay was used to detect the BxPC-3 and MIAPaCa-2 cell viability after treatment with gemcitabine only or with different concentrations of metformin. The methylation state and expression level of miR-663 were detected by methylation analysis and RT-PCR. Dual-luciferase reporter gene analysis, Western blot and RT-PCR were used to confirm the target of miR-663. Moreover, xenograft experiment was also performed to validate the role of metformin in chemosensitivity in vivo.

Results: We found that metformin increased the chemosensitivity of pancreatic cancer cells to gemcitabine, and epithelial–mesenchymal transition (EMT) progress caused by gemcitabine was suppressed by metformin. We further explored the possible molecular mechanisms and it was demonstrated that CpG islands of miR-663 were hypomethylated and relative expression level of miR-663 was up-regulated after treatment of metformin. miR-663, an important cancer suppressor miRNA, was confirmed to increase the chemosensitivity of pancreatic cancer cells by reversing EMT directly targeted TGF- β 1. Moreover, we identified that metformin increased the chemosensitivity through up-regulating expression of miR-663.

Conclusion: We demonstrated that metformin increased the chemosensitivity of pancreatic cancer cells to gemcitabine by reversing EMT through regulation DNA methylation of miR-663.

Keywords: metformin, pancreatic cancer, epithelial-mesenchymal transition; EMT, DNA methylation, miR-663

Introduction

Pancreatic cancer is one of the devastating malignancies of the digestive system. The incidence and mortality of pancreatic cancer have increased rapidly in recent years.¹ Patients, whether they received radical resection or were diagnosed at an advanced stage, are all recommended to receive comprehensive treatment including chemotherapy to improve prognosis.² However, the effect of chemotherapy in pancreatic cancer is still not satisfactory.³ The overall 5-year survival rate of pancreatic cancer is about 8%.^{1,4} Increasing the chemosensitivity of pancreatic cancer is one of the hotspots in research field of pancreatic cancer. In recent

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years, some evidence shows that metformin, a classic drug for diabetes mellitus, seems to improve survival of pancreatic cancer patients.^{5,6} The combination of metformin and gemcitabine has synergistic effect, and metformin can be used as an adjuvant drug to improve the clinical efficacy of gemcitabine of pancreatic cancer patients.⁷⁻⁹

It was found that epithelial-mesenchymal transition (EMT) can lead to chemotherapeutic resistance to gemcitabine in pancreatic cancer.¹⁰ When EMT process occurs in cancer cells, cell division is partly blocked, which reduces the efficiency of gemcitabine, a cell division dependent drug. Therefore, inhibiting EMT process can increase the sensitivity to gemcitabine, which provides new clues and ideas for reversing drug resistance in pancreatic cancer.^{11,12} We observed that metformin could reverse the gemcitabine-induced EMT process in pancreatic cancer cells. Thus, it is speculated that the reversal of the EMT process may be the mechanism of metformin to increase the sensitivity to gemcitabine. We further studied the molecular mechanism caused by metformin in pancreatic cancer cells.

Metformin can alter the DNA methylation status of cancer cells in a variety of ways.^{13,14} Our previous study found that the promoter region of miR-663 was abnormally hypermethylated in pancreatic cancer cells, which inhibit the expression of miR-663.¹⁵ After treatment with metformin in pancreatic cancer cells, the methylation status of miR-663 and its expression level was detected. We took miR-663 into further research, and then focused on its role in the chemosensitivity of pancreatic cancer cells. Our results indicate that metformin can increase the chemosensitivity to gemcitabine in pancreatic cancer cells by reversing EMT through regulation DNA methylation of miR-663.

Materials and Methods

In vitro Cell Culture

Pancreatic cancer cell lines BxPC-3 and MIAPaCa-2 were purchased from Shanghai Cell Bank and cultured in DMEM (Invitrogen, Carlsbad, CA, USA) with 10% fetal bovine serum (Sigma, St Louis, MO, USA) in a 5% CO₂ incubator at 37°C. The gemcitabine/metformin was dissolved in PBS with different concentrations.

Quantitative Real-Time Polymerase Chain Reaction

Pancreatic cancer cells were treated with TRIzol reagent (Invitrogen, Thermo Fisher Scientific, Inc) to extract RNA

according to the manufacturer's protocol. The expression level of miRNA was detected using All-in-One Customized Human qPCR Array (GeneCopoeia Inc, Rockville, Md, USA). The TGF- β 1 mRNA level was quantified using SYBR-Green PCR Master Mix kit (Takara, Dalian, China) normalized to β -actin. The primer sequences used were as follows. TGF- β 1, forward, 5'-GGACACCAACTATTGCTTCAG-3', TGF- β 1 reverse, 5'-TCCAGGCTCCAAATGTAG-3', β -actin, forward, 5'-AGAAAATCTGGCACCACACC-3', β -actin, reverse, 5'-TAGCACAGCCTGGATAGCAA-3'.

Western Blot

Total protein was extracted using RIPA lysis buffer. Protein samples were subjected to separation on SDS-PAGE gels and then transferred onto polyvinylidene difluoride membranes, which were blocked using 5% non-fat milk. The membranes were incubated with antibodies against TGF- β 1 (1:1000 dilution, Abcam, Cambridge, MA), E-cadherin (1:500 dilution, Santa Cruz, Dallas, TX, USA), Vimentin (1:1000 dilution, Abcam, Cambridge, MA), and β -actin (1:1000 dilution, Beyotime, Nantong, China) at 4°C overnight. The membranes were then incubated with secondary antibody at room temperature for 2 h. The protein bands were visualized using an ECL chemiluminescent kit (Pierce, Rockford, IL, USA). Quantification of Western blots was conducted using Index of Density.

Cell Transfection

miR-663 mimic or inhibitor and negative control was purchased from RiboBio (Guangzhou, Guangdong, China). Pancreatic cancer cell lines BxPC-3 and MIAPaCa-2 cells were cultured in 6-well plates for 24 h to 50–60% confluence and transfected with 30nmol/L miRNA or negative control using Lipofectamine 2000 (Invitrogen, Carlsbad, CA, USA).

Cell Counting Kit-8 (CCK-8) Assay

The BxPC-3 and MIAPaCa-2 were seeded into 96-well plates. After being treated with gemcitabine, or gemcitabine+metformin for 48 h, CCK-8 reagent was added to each well and further cultured for 2 h. Absorbance values were measured at 550 nm wavelength with a microplate reader (Tecan, Shanghai, China).

Wound Healing Assay

Cells were cultured in 6-well plates for 24 h to 70–80% confluence. The artificial wound was created using a 10 μ L

sterile pipette tip. After washed using PBS, cells were incubated in FBS-free DMEM for 24 h. The migrated cells were imaged at 0, 12 and 24 h. The cell migration ability was calculated by the proportion of the wound.

Cell Invasion Assay

Cell invasion assay was performed with a Transwell assay kit and invasion chambers (Corning, Corning, NY, USA). The upper chamber of 8 μ m pore transwells were overlaid with 1mg/mL Matrigel-coated membrane matrix. 0.25×10^5 cells per well (BxPC-3) or 0.5×10^5 cells per well (MIAPaCa-2) were plated in the upper chamber with serum-free medium. The lower chamber contained DMEM supplemented with 10% fetal bovine serum. After 24 h, cells invaded through the membrane were stained with 0.1% crystal violet for 30 min at 37 °C. Stained cells were soaked in 33% ice-cold acetic acid and absorbance values were measured at 550 nm wavelength with a microplate reader.

Dual-Luciferase Reporter Gene Analysis

miRanda, TargetScan, and TarBase were used to predict the miRNA targets. The mutant or wild-type 3'-UTR of TGF- β 1 sequence was cloned into the pGL3-control vector (Promega, Madison, WI, USA). After co-transfected with vector and miRNA, the luciferase activity was detected using the Dual-Luciferase Reporter Assay System (Promega).

DNA Isolation and Methylation Analysis

Genomic DNA was extracted from pancreatic cancer cell lines using the DNeasy kit (Qiagen, Hilden, Germany). Bisulfite modification procedure was according to the manufacturer's instructions of Epiect Bisulfite kit (Qiagen).

The PCR primers were designed as follows: miR-615 forward, 5'-AGTTAAGAGTAGTGGGGAGATTAA-3', reverse, 5'-AAATTTTTTTCTTTATTTACCCC-3', miR-663 forward, 5'-GTTGAGGAGGGTTTTTTTGTA-3', reverse, 5'-CCTCGATAAAAAACCTTCTCT-3', miR-663b forward, 5'-TATTTTATTAAGGGGAAGTGG-3', reverse, 5'-CCTCGATAAAAAACCTTCTCT-3', miR-675 forward, 5'-GGGTTTTGGAGGTTAGTGT-3', reverse, 5'-CTTAACAAACAATACAACATCCA-3'. The PCR products of sodium bisulfite-treated DNA were cloned into pTA2 vector using the TA cloning kit (Toyobo, Osaka, Japan), and single clones of six samples were sent for sequencing.

Xenograft Experiment

Six-week-old BALB/c nude mice were obtained from the Shanghai Experimental Animal Center. There were two groups, gemcitabine group and gemcitabine + metformin group. Five mice were assigned per group. A total of 1×10^7 MIAPaCa-2 cells were injected subcutaneously into the neck area of each mouse. After cell injection, the mice were treated with drugs, and the scheme was as following, gemcitabine only group (100 mg/kg of gemcitabine injected intraperitoneally twice weekly), combination gemcitabine with metformin group (gemcitabine 100 mg/kg injected intraperitoneally twice weekly and 800 mg/L metformin in drinking water each day). The tumor width, length, and height were measured by external caliper every 5 days and calculated as follows: Volume (mm^3) = width \times length \times height/2. The mice were sacrificed by cervical dislocation following 25 days of treatment and the tumors were collected, fixed, and embedded. The animal experiments were approved by the Animal Care and Use Committee of The Second Affiliated Hospital of Nanjing Medical University, approval number 2018-KY-027, and in accordance with the Guide for the Care and Use of Laboratory Animals (eighth edition, 2011, National Institutes of Health, USA).

Immunohistochemistry

Immunohistochemistry was used to examine TGF- β 1 expression in tumor xenografts. After fixed in formalin, embedded in paraffin blocks, tumor tissues were incubated with primary antibody against TGF- β 1 (1:500 dilution, Abcam, Cambridge, MA) overnight at 4 °C. Goat anti-rabbit IgG antibody conjugated to HRP (1:200 dilution, Santa Cruz, Dallas, TX, USA) were incubated for 2 h at room temperature. Subsequently, the sections were stained with DAB and counterstained with hematoxylin.

Statistical Analysis

All experimental results were shown as the means \pm standard deviation of three separate experiments. The statistical significance of differences was performed using the Student *t* tests when only two groups were compared. The difference among multiple groups was compared using one-way ANOVA using Dunnett's multiple comparison test. $P < 0.05$ was considered to indicate a significant.

Results

Metformin Increased the Chemosensitivity of Pancreatic Cancer Cells to Gemcitabine by Reversing EMT *in vitro*

To study the role of metformin in the chemosensitivity of pancreatic cancer cells to gemcitabine, the CCK-8 assay was used to detect the BxPC-3 and MIAPaCa-2 cell viability after treatment with gemcitabine only or with different concentrations of metformin. It showed that the survival of pancreatic cancer cells was inhibited by gemcitabine, and when combined with different concentrations of metformin, cell viability was further inhibited which exhibited synergistic effect (Figure 1A). The difference was statistically significant. We used CalcuSyn software

to evaluate the synergistic effect of gemcitabine and metformin, and found that the best CI value was 0.549 when the concentration of gemcitabine and metformin was 143.2nmol/L and 6.8mmol/L for MIAPaCa-2, and the best CI value was 0.645 when the concentration of gemcitabine and metformin was 633.9nmol/L and 4.8mmol/L for BxPC-3.

Before and after treatment of gemcitabine only or with metformin, morphological changes were taken by microscope of BxPC-3 and MIAPaCa-2. BxPC-3 and MIAPaCa-2 cells clustered together, and after treatment of gemcitabine, cells gradually transitioned from epithelial-like cells to mesenchymal-like cells, which were elongated, and grew individually. While it was treated with gemcitabine and metformin, EMT progress was suppressed and epithelial-like cells reappeared. (Figure 1B).

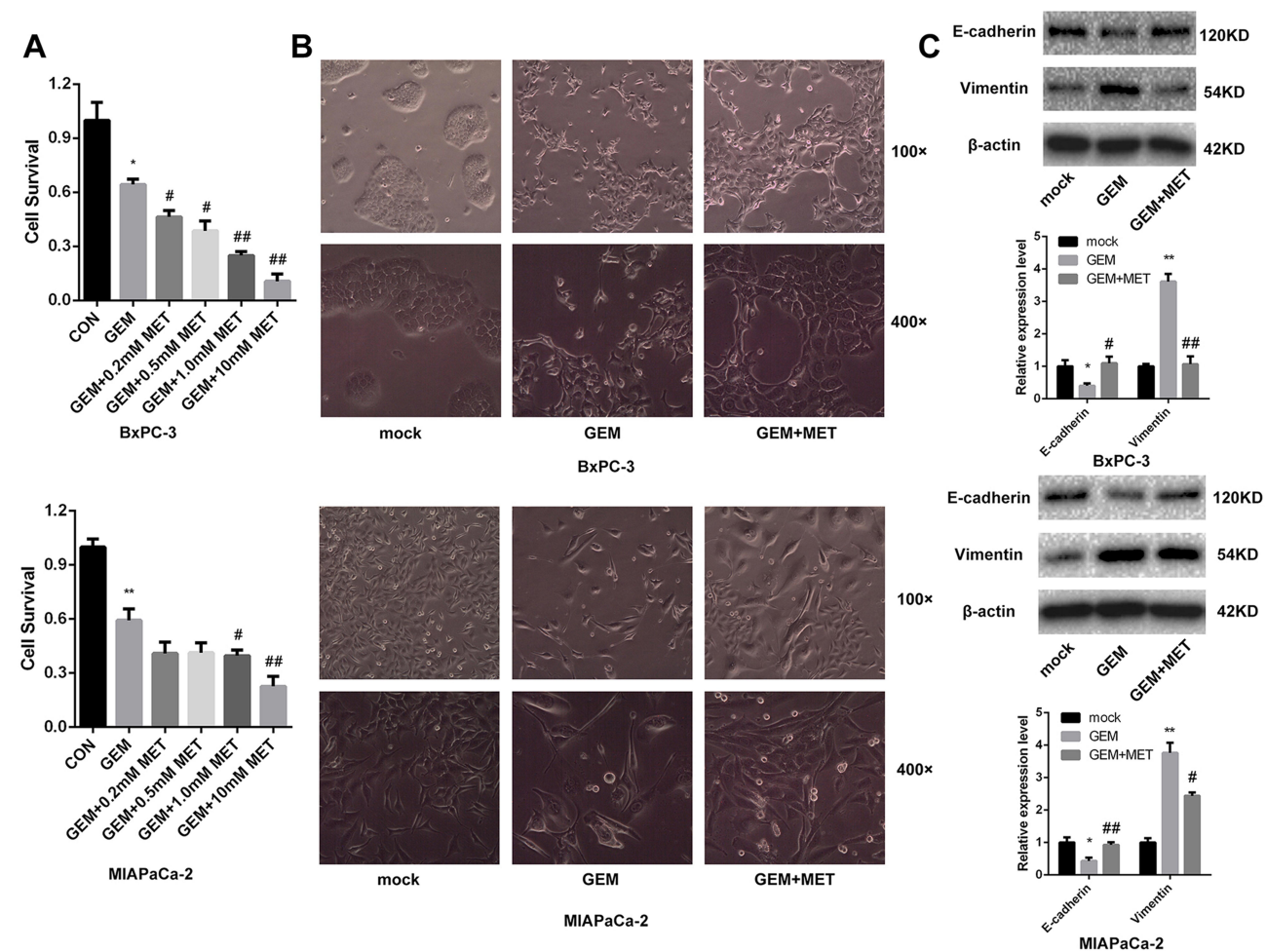


Figure 1 Metformin increased the chemosensitivity of pancreatic cancer cells by reversing EMT. (A) Cell viability was determined by CCK-8 assay after treated with gemcitabine only or with different concentrations of metformin. * P_{CON} vs. GEM < 0.05, ** P_{CON} vs. GEM < 0.01, # P_{GEM} vs. GEM+MET < 0.05, ## P_{GEM} vs. GEM+MET < 0.01. (B) Morphology changes of BxPC-3 and MIAPaCa-2 cells were taken before and after treated with gemcitabine only or with metformin. (C) The protein levels of E-cadherin and Vimentin in BxPC-3 and MIAPaCa-2 cells were determined by Western blot analyses. * P_{CON} vs. GEM < 0.05, ** P_{CON} vs. GEM < 0.01, # P_{GEM} vs. GEM+MET < 0.05, ## P_{GEM} vs. GEM+MET < 0.01.

EMT-related protein E-cadherin and Vimentin were detected in pancreatic cancer cells using Western blot.

The lower expression level of E-cadherin and higher expression level of Vimentin was found after gemcitabine treatment in both BxPC-3 and MIAPaCa-2 cells, while it was treated with gemcitabine and metformin, E-cadherin was increased and Vimentin was decreased comparing with metformin only group (Figure 1C). The difference was statistically significant.

CpG Islands of miR-663 Were Hypomethylated and Relative Expression of miR-663 Was Up-Regulated After Treatment of Metformin in Pancreatic Cancer Cells

As we described before,¹⁵ CpG islands of miR-615, miR-663, miR-663b and miR-675 were confirmed to be hypermethylated. It had been reported that metformin could alter the DNA methylation status of cancer cells, so after treated of metformin for BxPC-3, the percentage of methylated CpG sites of the four miRNAs was re-examined and miR-663 was shown to be hypomethylated (80.3±5.5% vs. 30.3±13.8%, the difference was statistically significant), while there was no significant change of miR-615, miR-663b and miR-675 (Figure 2A). Furthermore, the expression of miR-663 in BxPC-3 and MIAPaCa-2 cells was up-regulated by 5.7±0.3 and 5.1±0.6 fold after treatment of metformin (Figure 2B), and the difference was statistically significant.

miR-663 Increased the Chemosensitivity of Pancreatic Cancer Cells to Gemcitabine by Reversing EMT in vitro

Because miR-663 was up-regulated after treatment of metformin, we further evaluated the role of miR-663 in chemosensitivity of pancreatic cancer cells to gemcitabine. CCK-8 assay revealed that up-regulation of miR-663 significantly increased the chemosensitivity to gemcitabine of BxPC-3 and MIAPaCa-2 cells (Figure 3C). The difference was statistically significant.

Morphological changes was also taken after treatment of gemcitabine on the miR-663-overexpressed BxPC-3 and MIAPaCa-2 cells, and the EMT progress caused by gemcitabine was suppressed by miR-663 (Figure 3A). The expression levels of EMT-related protein were detected, and it was found that E-cadherin was increased and

Vimentin was decreased after overexpression of miR-663 (Figure 3B). The difference was statistically significant.

It was reported that EMT was also related to invasion and metastasis of pancreatic cancer. The migratory ability of BxPC-3 and MIAPaCa-2 cells measured by the wound healing assay demonstrated that miR-663 significantly inhibited the cell migration (Figure 4A), and miR-663 reduced the cell invasive ability using a Matrigel invasion assay (Figure 4B). The difference was statistically significant.

miR-663 Directly Targeted TGF-β1

In order to investigate the role of miR-663 in pancreatic cancer, we used Targetscan to explore the potential targets of miR-663, and EMT-related gene TGF-β1 was selected for further analysis (Figure 5A). Two groups of luciferase reporter constructs were used to confirm the prediction. It was shown that the luciferase activity was significantly reduced to 65.3±4.9% of BxPC-3 and 26.0±5.0% of MIAPaCa-2 in miR-663 mimic + TGF-β1_WT co-transfected group comparing with WT+NC group (Figure 5B).

We further examined the regulation of miR-663 on TGF-β1 expression in pancreatic cancer cells. The results indicated that miR-663 significantly inhibited the expression level of TGF-β1 mRNA (Figure 5C) and protein (Figure 5D) in BxPC-3 and MIAPaCa-2 cells. These data indicated that TGF-β1 was the direct target of miR-663. The difference was statistically significant.

Metformin Increased the Chemosensitivity of Pancreatic Cancer Cells Through Up-Regulating Expression of miR-663

In order to test our hypothesis that the effect of metformin resulted from up-regulating the expression of miR-663, BxPC-3, and MIAPaCa-2 cells were co-treated with miR-663 inhibitor and metformin. CCK-8 assay revealed that down-regulation of miR-663 significantly decreased the chemosensitivity to gemcitabine of BxPC-3 and MIAPaCa-2 cells. When co-treated with metformin and miR-663 inhibitor, there were no statistical differences between control group and miR-663 inhibitor + metformin group (Figure 6A).

The expression level of TGF-β1 protein was detected in pancreatic cancer cells using Western blot. Compared with control group, TGF-β1 protein was increased in gemcitabine

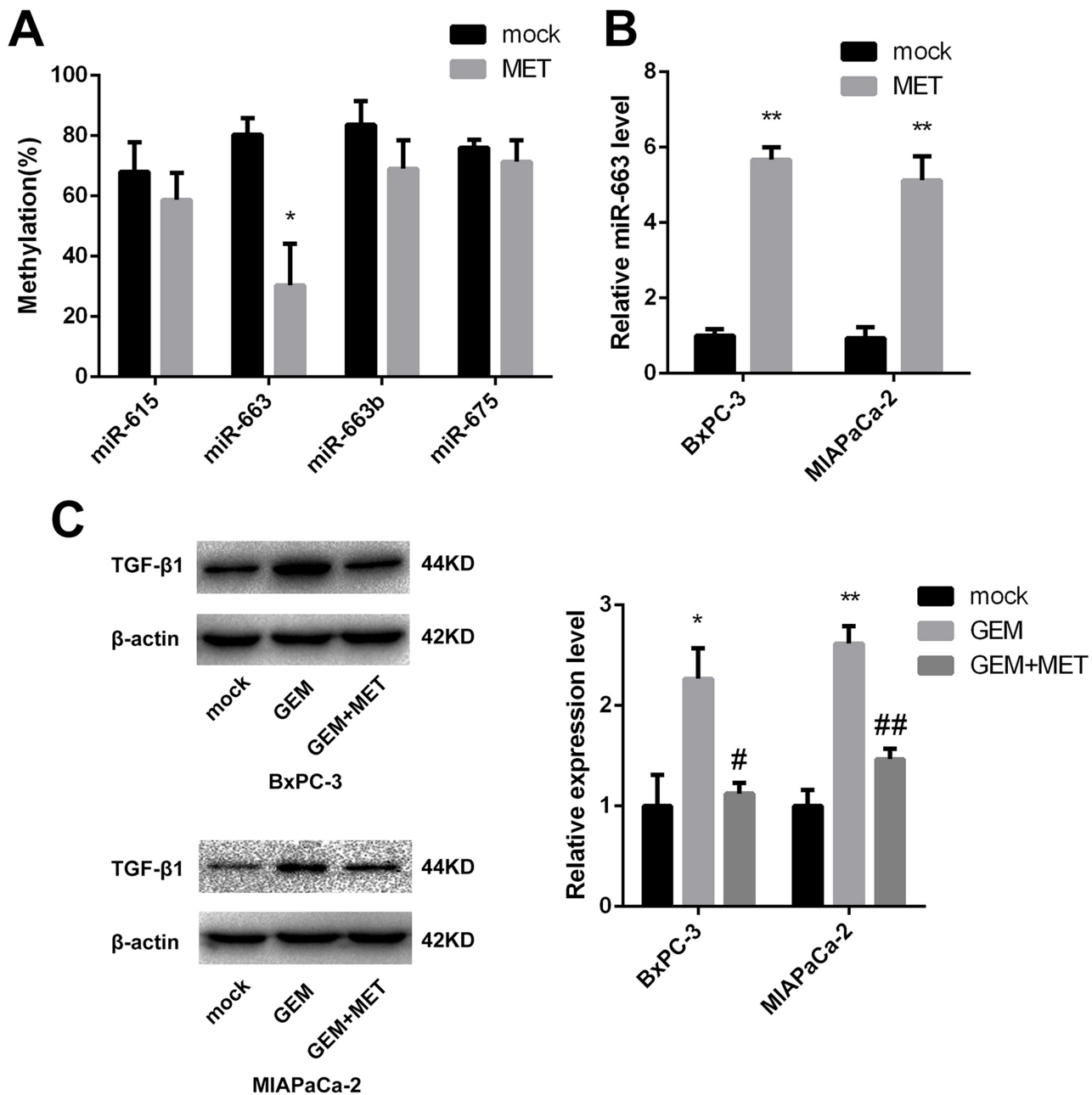


Figure 2 DNA methylation status and relative expression of miR-663 was altered after treatment of metformin. **(A)** The percentage of methylated CpG sites of miR-615, miR-663, miR-663b and miR-675 was shown for BxPC-3 before and after treatment of metformin. * $P < 0.05$. **(B)** Relative expression of miR-663 was shown for BxPC-3 and MIAPaCa-2 before and after treatment of metformin. ** $P < 0.01$. **(C)** The protein level of TGF- β 1 in BxPC-3 and MIAPaCa-2 cells were determined by Western blot analyses. * P_{CON} vs. GEM < 0.05 , ** P_{CON} vs. GEM < 0.01 , # P_{GEM} vs. GEM+MET < 0.05 , ## P_{GEM} vs. GEM+MET < 0.01 .

group. When treated with gemcitabine and metformin, the increased TGF- β 1 protein trend disappeared (Figure 2C). We also detected the TGF- β 1 protein in miR-663 inhibitor group and miR-663 inhibitor + metformin group of BxPC-3 and MIAPaCa-2 cells, the TGF- β 1 protein was down-regulated in the miR-663 inhibitor group compared with the control group (Figure 6B). The difference was statistically significant. When co-treated with metformin and miR-663 inhibitor, there were no statistical differences of TGF- β 1

protein between the control group and miR-663 inhibitor + metformin group (Figure 6C).

Metformin Increased the Chemoresensitivity of Pancreatic Cancer Cells to Gemcitabine in vivo

To investigate the role of metformin in chemoresensitivity of pancreatic cancer cells to gemcitabine in vivo, xenograft

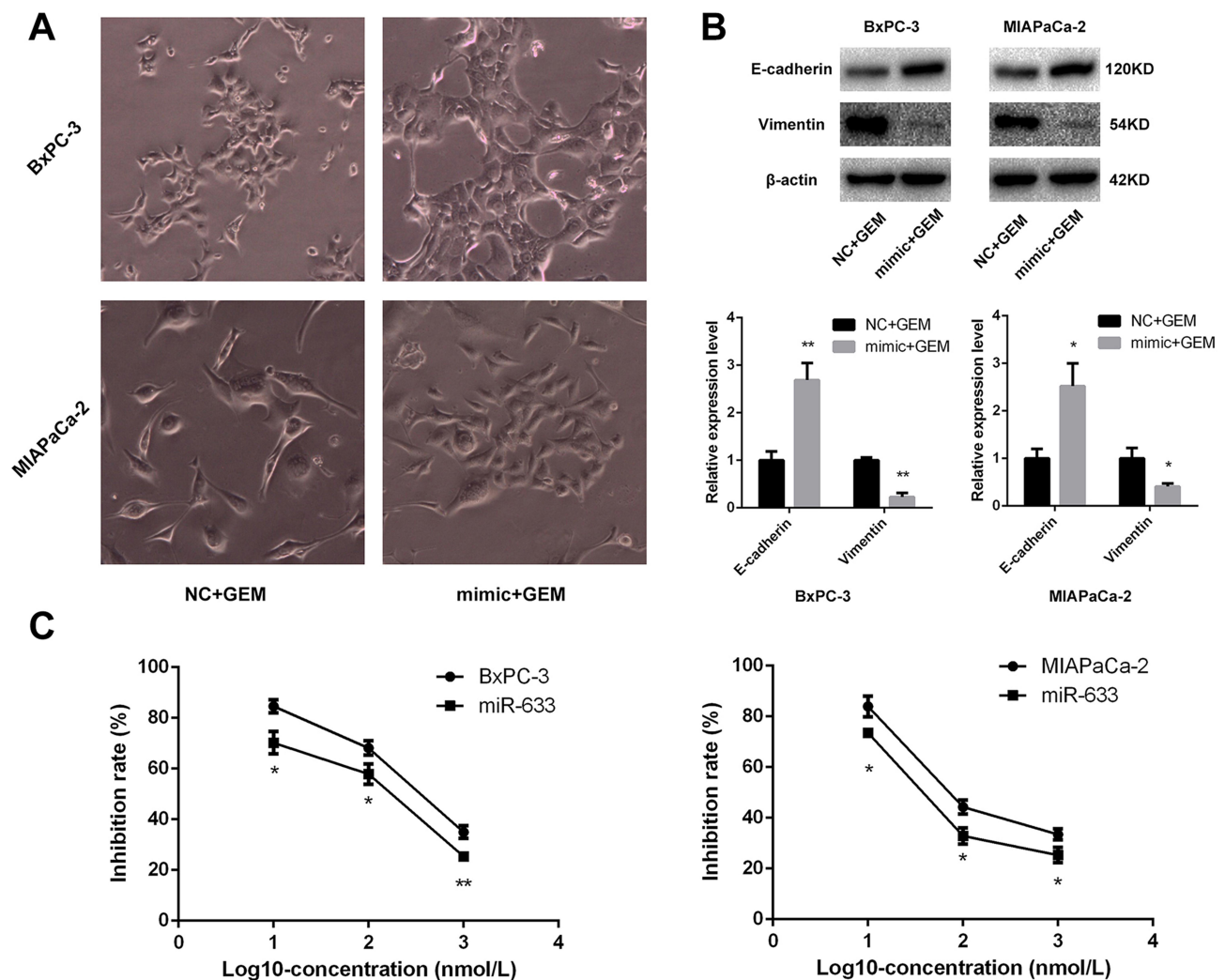


Figure 3 miR-663 increased the chemosensitivity of pancreatic cancer cells by reversing EMT. (A) Morphology changes of BxPC-3 and MIAPaCa-2 cells were taken treating with gemcitabine after transfected with miR-663 mimic or negative control (NC). Magnification: $\times 400$. (B) The protein levels of E-cadherin and Vimentin in BxPC-3 and MIAPaCa-2 cells were determined by Western blot analyses. * $P < 0.05$, ** $P < 0.01$. (C) Cell inhibition rate was determined by CCK-8 assay treated with different concentrations of gemcitabine after transfected with miR-663 mimic or NC. * $P < 0.05$, ** $P < 0.01$.

experiments were conducted. The tumor volume was measured every 5 days, and tumor growth was slower in the gemcitabine group than in the gemcitabine + metformin group (Figure 7A, 7B). The difference was statistically significant. Mice were sacrificed at the 25th day and the weights of the tumors were measured. The average weight of tumors from the gemcitabine + metformin group was significantly less than that from the gemcitabine group (360.0 ± 85.4 vs. 180.0 ± 60.0 mg, the difference was statistically significant) (Figure 7C). Furthermore, the protein expression levels of TGF- $\beta 1$ in tumor tissues were examined by immunohistochemistry and it was markedly down-regulated in the gemcitabine + metformin group than in the gemcitabine group ($83.3 \pm 8.6\%$ vs. $45.3 \pm 6.7\%$, the difference was statistically significant) (Figure 7D, 7E).

Discussion

Chemotherapy is widely used in all stages of pancreatic cancer patients, which is an important part of comprehensive treatment of pancreatic cancer. However, the effect of chemotherapy in pancreatic cancer is still not satisfactory, and drug resistance is the main reason for the failure of chemotherapy. EMT process appeared in the process of gemcitabine treatment of pancreatic cancer cells.¹⁶ The volume of epithelial cells reduced gradually, and the intercellular junction decreased, which turned to the characteristics of mesenchymal cells. The expression level of epithelial marker E-cadherin was down-regulated and mesenchymal marker Vimentin was up-regulated, which resulted in the loss of cell polarity in epithelial cells. EMT process can lead to resistance of gemcitabine in pancreatic cancer.¹⁷

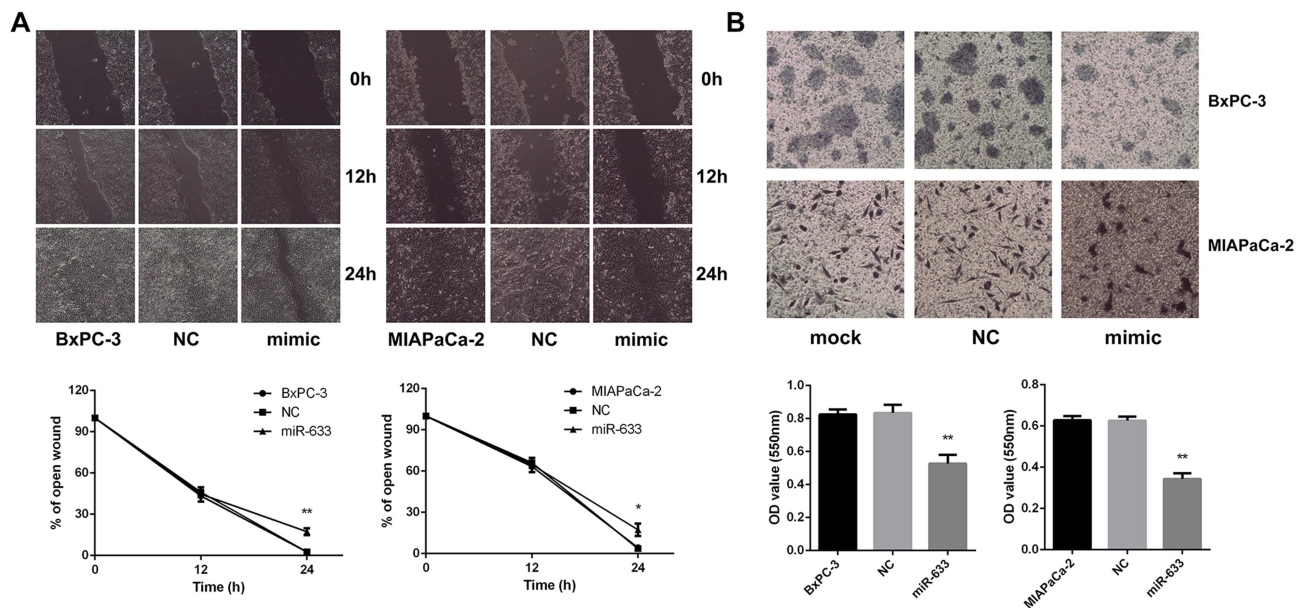


Figure 4 miR-663 inhibited cell migration and invasion of BxPC-3 and MIAPaCa-2 in vitro. **(A)** Wound healing assay was performed on BxPC-3 and MIAPaCa-2 cells after transfected with miR-663 mimic or NC. Magnification: $\times 200$. * $P < 0.05$, ** $P < 0.01$. **(B)** Transwell invasion assay was performed on BxPC-3 and MIAPaCa-2 cells after transfected with miR-663 mimic or NC. Magnification: $\times 200$. ** $P < 0.01$.

Tumor cells show mesenchymal transition in the process of acquired drug resistance, and EMT process has gradually been considered as an important mechanism of drug resistance of tumor cells.¹⁰ More and more studies have shown that there are common regulatory signal pathways between EMT process and drug resistance of tumor cells.^{18,19} The inactivation and imbalance of EMT process-related signaling pathway play an important role in drug resistance. Therefore, reversal of the process of EMT provides a new clue to solve the problem of drug resistance.

Metformin has been proven to increase chemosensitivity to gemcitabine in pancreatic cancer, while its mechanism is not yet clear. Zhong et al¹³ found that metformin affected the expression of DNA methylase 3B by regulating the activity of S-adenosylhomocysteine (SAHH), thus altering the percentage of genomic DNA methylation. Cuyas et al¹⁴ found that metformin could increase the S-adenosylmethionine (SAM)/SAHH ratio and regulate the percentage of DNA methylation. In our previous study, we used methylation chip to detect the DNA methylation status of pancreatic cancer cells. The results showed that the promoter regions of 11 microRNAs were abnormally methylated in pancreatic cancer cells compared with normal pancreatic tissues. Pyrosequencing and TA clone sequencing was then used to confirm the presence of aberrant methylation in the promoter regions of miR-615, 663, 663b and 675.¹⁵ We explored whether metformin

could alter the methylation status and the expression of the above microRNAs. After treatment with metformin in pancreatic cancer cells, the percentage of methylation of miR-663 was down-regulated and its expression level was up-regulated.

miR-663 is an important miRNA in tumorigenesis. The role of miR-663 in gastric cancer,²⁰ breast cancer,²¹ liver cancer,²² lung cancer,²³ bladder cancer,²⁴ nasopharyngeal cancer,²⁵ and other malignant tumors has been found, and its main functions focus on regulation of tumor proliferation, invasion, metastasis, and apoptosis. The expression level of miR-663 is varied in different types of tumors, and we noticed that it is regulated by DNA methylation²⁶ and drugs (retinoic acid,²⁷ fluoxetine)²⁸ in some reported research. At the same time, the abnormal expression of miR-663 also has been found in drug-resistant breast cancer cells²¹ and castrated refractory prostate cancer.²⁹ In pancreatic cancer, Zhang et al³⁰ reported that the expression of miR-663 was lower than that of normal tissues, and miR-663 inhibited the proliferation and invasion of cancer cells. However, there are few reports about the function of miR-663 in chemotherapy of cancers. Our previous study found that abnormal DNA methylation inhibits the expression of miR-663 in pancreatic cancer cells. We over-expressed miR-663 using miRNA mimic in pancreatic cancer cell lines, and found that miR-663 significantly increased the chemosensitivity to gemcitabine.

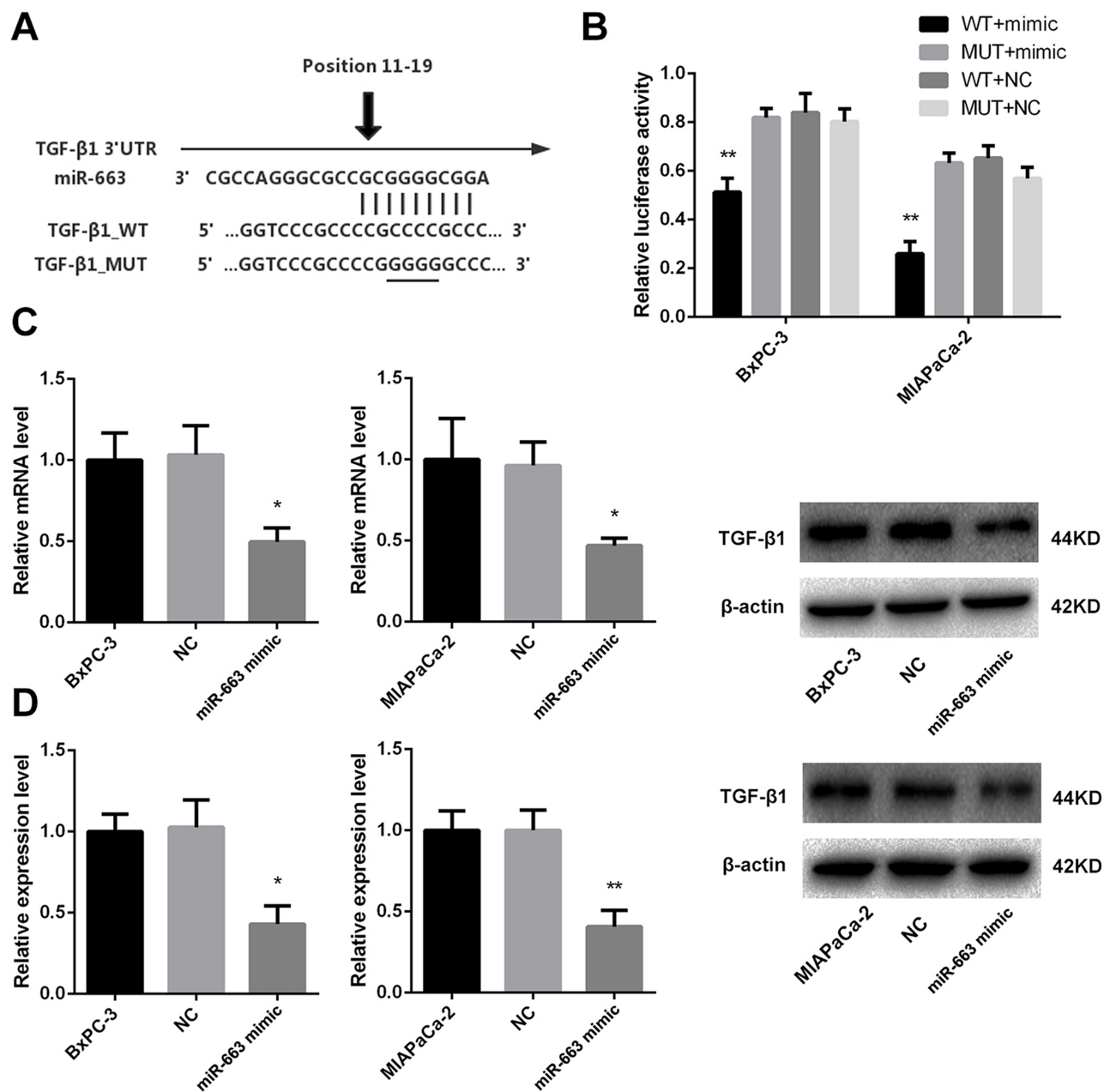


Figure 5 miR-663 inhibited expression of TGF-β1 by targeting 3'-UTR of TGF-β1 in pancreatic cancer cells. **(A)** Sequence of the miR-663 binding site with TGF-β1 3'-UTR (TGF-β1_WT) and the mutated TGF-β1 3'-UTR sequence (TGF-β1_MUT). **(B)** BxPC-3 and MIAPaCa-2 cells were co-transfected with miR-663 mimic or NC and luciferase reporter vectors containing the TGF-β1_WT or TGF-β1_MUT sequences of the miR-663 binding sites. Luciferase activities were measured 48 h after transfection. $**P < 0.01$. **(C)** The relative expression of TGF-β1 mRNA was measured by real-time PCR in BxPC-3 and MIAPaCa-2 cells. $*P < 0.05$, $**P < 0.01$. **(D)** TGF-β1 protein in BxPC-3 and MIAPaCa-2 cells was detected by Western blot. $*P < 0.05$, $**P < 0.01$.

The reported target genes of miR-663 in cancer cells include JUND, JUNB, PI3K, EEF1A2, which are closely related to multiple cellular functions.³⁰⁻³² TGF-β1, a recognized gene that induces EMT process,³³ is also the target gene of miR-663. TGF-β1 can activate Smad and bind with related nuclear transcription factors through the classical signaling pathway, then regulates the expression of downstream genes. Thereby, the signaling pathway

adjusts the junctional structure between epithelial cells, and makes epithelial cells lose polarity and turn into mesenchymal cells.³⁴ Bioinformatics prediction software showed that the 18-24 base of 3'-UTR region of TGF-β1 paired with miR-663, and when it was over-expressed of miR-663 in pancreatic cancer cells, the expression of TGF-β1 was down-regulated at both the level of mRNA and protein. Luciferase reporter gene experiment further

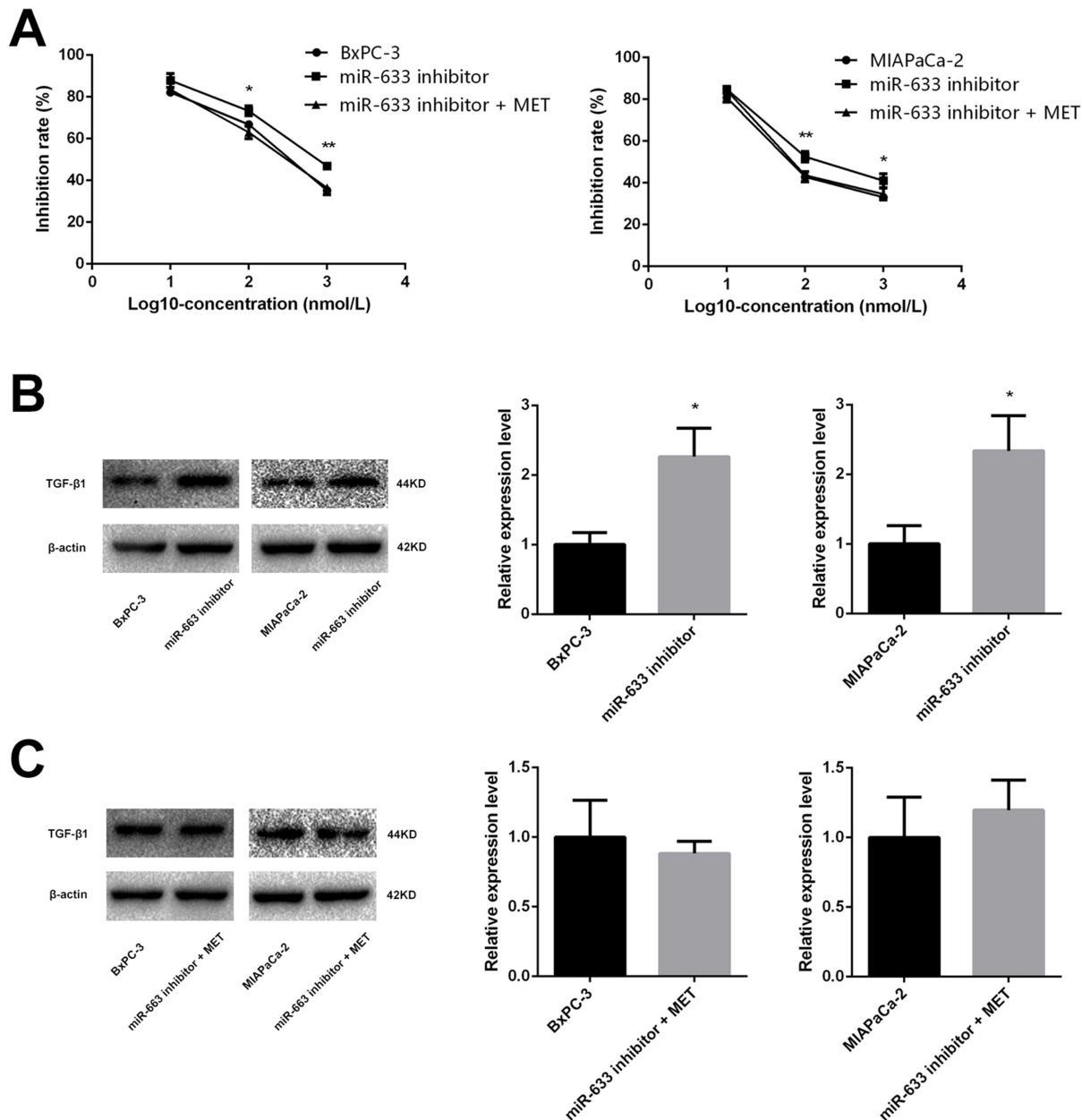


Figure 6 Metformin increased the chemosensitivity through up-regulating expression of miR-663. **(A)** Cell inhibition rate was determined by CCK-8 assay treated with different concentrations of gemcitabine after transfected with miR-663 inhibitor only or with metformin. * P_{CON} vs. inhibitor < 0.05, ** P_{CON} vs. inhibitor < 0.01. **(B and C)** The protein level of TGF-β1 in BxPC-3 and MIAPaCa-2 cells were determined by Western blot analyses. * P < 0.05.

confirmed the targeting pairing relationship between miR-663 and the 3'-UTR region of TGF-β1. Therefore, we speculated that the expression of TGF-β1 was regulated by miR-663, and then inhibited the process of EMT in pancreatic cancer cells.

It has been shown that another metformin-like drug AICAR is effective in combination with mTOR inhibitor-Rapamycin as anti-cancer agent.³⁵ Interestingly, metformin activates AMPK while AMPK regulates the

mTOR pathway in a feedback loop manner.³⁶ In this study, we found that whether gemcitabine or gemcitabine + metformin treatment would affect the protein level of TGF-β, which is known to activate Akt through PI3K, that in turn activates the mTOR complex 1 (mTORC1).³⁷ It will be interesting to see how this gemcitabine and metformin combination treatment affects mTOR signaling pathways. This may be meaningful in possible future studies.

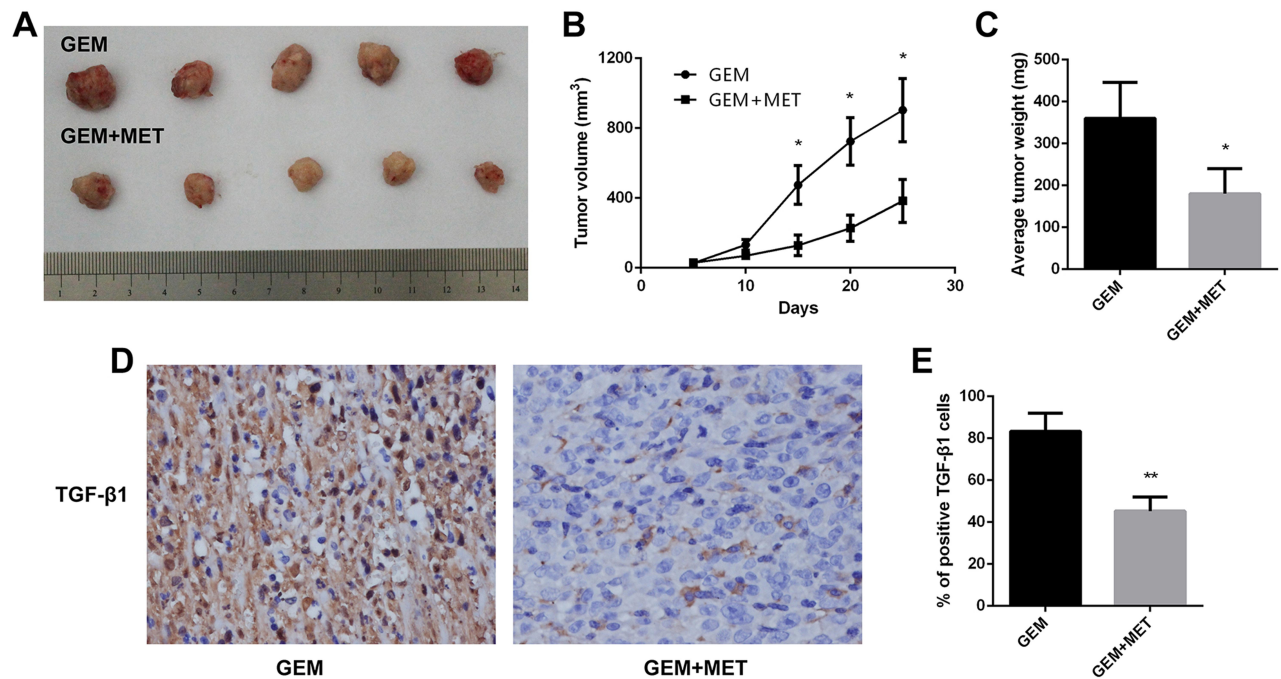


Figure 7 Metformin increased the chemosensitivity of pancreatic cancer in vivo. **(A)** Tumors were harvested from gemcitabine group and gemcitabine+metformin group of nude mice. **(B)** The tumor volume growth curve in vivo was shown. * $P < 0.05$. **(C)** The tumor weight of each group were calculated. * $P < 0.05$. **(D)** TGF- β 1 protein levels were measured by immunohistochemistry in the tumor tissues isolated from the mice. Magnification: $\times 40$. **(E)** The percentage of TGF- β 1 positive cells was calculated. ** $P < 0.01$.

The antitumor effect of metformin in pancreatic cancer patients is still controversial. Although the meta-analysis suggests that the usage of metformin is associated with a significantly lower mortality rate in pancreatic cancer patients,⁶ different results also existed that metformin use was not statistically significantly associated with improved survival.³⁸ Therefore, we suppose that some selected pancreatic cancer patients can benefit from the usage of metformin, and further studies are needed.

In conclusion, the present study demonstrated that inhibition of EMT process accompanied with up-regulation of miR-663 was found in pancreatic cancer cells when treated with metformin. Over-expression of miR-663 increased the sensitivity of pancreatic cancer cells to gemcitabine and inhibited the EMT process. The analysis of the mechanism showed that TGF- β 1 was the target gene of miR-663. Our results provide a new theoretical basis of metformin for improving chemosensitivity to gemcitabine in pancreatic cancer.

Author Contributions

All authors made a significant contribution to the work reported, whether that is in the conception, study design, execution, acquisition of data, analysis and interpretation, or in all these areas; took part in drafting, revising or

critically reviewing the article; gave final approval of the version to be published; have agreed on the journal to which the article has been submitted; and agree to be accountable for all aspects of the work.

Disclosure

The authors declare that they have no conflicts of interest for this work.

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