

# Leptin receptor mediates the proliferation and glucose metabolism of pancreatic cancer cells via AKT pathway activation

YINGJIE XU<sup>1\*</sup>, MEIYU TAN<sup>2\*</sup>, XIAOYU TIAN<sup>2</sup>, JUN ZHANG<sup>3</sup>, JIE ZHANG<sup>2</sup>,  
JIAJIE CHEN<sup>2</sup>, WEIHONG XU<sup>2</sup> and HUIMING SHENG<sup>2</sup>

Departments of <sup>1</sup>Surgery and <sup>2</sup>Clinical Laboratory, Shanghai Tongren Hospital, Shanghai 200336;

<sup>3</sup>Digestive Disease Research Institute, Shanghai Huashan Hospital, Shanghai 200041, P.R. China

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**Abstract.** Pancreatic cancer (PC) is the fourth leading cause of cancer-related mortality worldwide. Leptin is an adipokine that is significantly increased in obese patients and that functions in various biological processes of cancer, such as tumor growth and metastasis. However, its role in PC cell proliferation and glucose metabolism and the underlying mechanisms remain unclear. In the present study, *in vitro* leptin treatment significantly promoted cell proliferation and increased glucose uptake and lactate production of human PC and healthy pancreas cells in a dose-dependent manner, accompanied by increased expression of the glycolytic enzymes hexokinase II and glucose transporter 1. Furthermore, leptin receptor-specific short hairpin RNAs were used to silence leptin receptor expression in PC cells, which had the opposite effect to leptin stimulation and decreased AKT phosphorylation. In addition, the effects of leptin stimulation were significantly counteracted by the AKT inhibitor LY294002, whereas the effects of leptin silencing were counteracted by AKT activator insulin-like growth factor 1. The results of the present study suggested that leptin may contribute to cell proliferation and glucose metabolism of human PC cells, which may be through activation of the AKT pathway.

## Introduction

As the fourth leading cause of cancer mortality, pancreatic cancer (PC) is an aggressive disease, and its global prevalence has been increasing (1,2). The 5-year overall survival rate of

patients diagnosed with PC is <5% (3). It has been reported that obesity is one of the risk factors for PC development, and over the past decade, the rate of obesity has risen at an unprecedented rate (4,5). Chronic obesity changes the production and secretion of adipokines, a type of cytokine secreted by the adipose tissue (6-9).

The adipokine leptin has been reported to be highly expressed in obese patients (6,7). Leptin regulates a variety of biological processes, such as satiety, food intake and energy expenditure, through leptin receptor (LepR) (10-12). Glucose metabolism, controlled by cellular ATP and metabolite levels, offers energy and required substance for tissue anabolism and catabolism, and is essential for the growth and development of cells and tissues (13,14). In addition, increasing evidence showed that glucose is an important regulator for leptin expression and secretion (15-17). LepR is composed of several isoforms including the short isoform (LepR-short) and long isoform (LepR-long). It has been reported that LepR can trigger the activation of several pathways, including PI3K/AKT and Janus-activated kinase/STAT (18-20). LepR common variant (LepR-common) is reported to influence gestation glycemic traits (21). Leptin is highly expressed in subsets of cancer patients including breast, colon and prostate cancer, and stimulates various biological activities in cancer cells, including cell proliferation, migration and invasion (22-25). Leptin is a growth factor for colon epithelial cells, and it has been demonstrated to promote motility and invasiveness in human colon cancer cells (22,23). Leptin also promotes invasion and migration of breast cancer cells through trans-activation of epidermal growth factor receptor (23). Pancreatic  $\beta$ -cells have been demonstrated to express functional LepRs, and LepR-short and LepR-long in PC cells (4,26). A previous study reported that leptin signaling enhances human PC cell invasion and metastasis by increasing matrix metalloproteinase-13 (MMP-13) (27). However, the effects of LepR on the proliferation and glucose metabolism of human PC, as well as its underlying mechanisms remain unclear.

In the present study, *in vitro* leptin stimulation significantly promoted cell proliferation and enhanced glucose metabolism of human PC and normal pancreas cells in a dose-dependent manner, accompanied by an increase in the expression levels of the glycolytic enzymes hexokinase II (HKII) and glucose transporter 1 (GLUT1). Silencing of LepR decreased AKT

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**Correspondence to:** Dr Weihong Xu or Dr Huiming Sheng, Department of Clinical Laboratory, Shanghai Tongren Hospital, 1111 Xianxia Road, Changning, Shanghai 200336, P.R. China  
E-mail: xwh3066@126.com  
E-mail: shm2783@shtrhospital.com

\*Contributed equally

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phosphorylation. Additionally, the induction of leptin stimulation was significantly counteracted by treatment with an AKT inhibitor (LY294002), whereas the effect of LepR silencing was counteracted by the AKT activator insulin-like growth factor 1 (IGF-1). The results of the present study suggested that leptin may contribute to human PC cell proliferation and glucose metabolism, which may be through activation of the AKT pathway.

## Materials and methods

**Cell culture.** Two human PC cell lines, BxPC3 and Panc-1, and normal pancreas cells HPC-Y5 were purchased from The Cell Bank of Type Culture Collection of the Chinese Academy of Sciences. The cells were cultured in a 37°C, 5% CO<sub>2</sub> incubator (Thermo Forma 3111; Thermo Fisher Scientific, Inc.) in RPMI-1640 medium (cat. no. SH30809.01B; HyClone; GE Healthcare Life Sciences), which contained 10% FBS (cat. no. 16000-044; Gibco; Thermo Fisher Scientific, Inc.) and 1% penicillin/streptomycin (cat. no. P1400-100; Beijing Solarbio Science & Technology Co., Ltd.). The medium was refreshed every two days during incubation.

**Lentiviral transduction.** Short hairpin RNA (shRNA)-1 and shRNA-2 targeting two different sites of LepR (GenBank no. BC131779.1; Table I) were constructed and inserted into the *AgeI*/*EcoI* restriction sites of a pLKO.1-puro vector (Addgene, Inc.). Following confirmation by DNA sequencing (Shanghai Majorbio Pharmaceutical Technology Co., Ltd.), LepR-shRNA-1 or LepR-shRNA-2 was co-transfected with the viral packaging plasmids psPAX2 and pMD2G (Addgene, Inc.) into 293T (The Cell Bank of Type Culture Collection of the Chinese Academy of Sciences) cells using Lipofectamine® 2000 (Invitrogen; Thermo Fisher Scientific, Inc.) in a 37°C incubator. The virus particles were collected by ultracentrifugation 48 h post-transfection.

BxPC3 cells were transduced with lentiviruses expressing shRNA negative control (shNC), LepR-shRNA-1 and LepR-shRNA-2; medium-treated cells were used as the untransfected control. Following 48 h of transduction by lentivirus infection in a 37°C, 5% CO<sub>2</sub> incubator, the silencing efficiency of LepR-shRNA-1 and LepR-shRNA-2 were evaluated by reverse transcription-quantitative PCR (RT-qPCR) and western blot analysis. Subsequently, cell proliferation, glucose uptake and lactate production, as well as the expression of several related genes and proteins including LepR-short, LepR-long, HKII, GLUT1, AKT and p-AKT, were measured at 48 h of transduction.

**Experimental groups.** BxPC3, Panc1 or HPC-Y5 cells were treated with 0, 20, 50 or 100 ng/ml Leptin. Subsequently, cell proliferation, glucose uptake, lactate production as well as the expression of HKII and GLUT1 were detected. The treatment groups for BxPC3 cells were: i) RPMI-1640 medium (control); ii) shRNA negative control (shNC); iii) LepR-shRNA-1 (shRNA-1); iv) LepR-shRNA-2 (shRNA-2), or i) Medium + DMSO (control); ii) 50 ng/ml leptin + DMSO; iii) medium + 25 µmol/l LY294002 (AKT inhibitor); and iv) 50 ng/ml leptin + 25 µmol/l LY294002. A subset of shNC and shRNA-2-transfected BxPC3 cells were further treated with DMSO or

50 ng/ml IGF-1 (AKT activator). Following treatment, cell proliferation, glucose uptake, lactate production, and the expression of several related genes (HKII, GLUT1, AKT and p-AKT) were examined.

**Proliferation assay.** The proliferation of treated human PC cells (BxPC3 and Panc1) and normal pancreas cells HPC-Y5 were evaluated by the Cell Counting Kit-8 assay (CCK-8; cat. no. CP002; SAB Biotherapeutics, Inc.). Human PC cells were seeded in 96-well plates (3×10<sup>3</sup> cells/well) and cultured in a 37°C, 5% CO<sub>2</sub> incubator overnight. Following treatment for 24 h, 100 µl CCK-8 solution (diluted 1:10 in serum-free medium) was added and incubated for 1 h in a 37°C, 5% CO<sub>2</sub> incubator. Subsequently, the absorbance value [optical density (OD)] at 450 nm was determined using a microplate reader (DNM-9602; Perlong Medical Equipment Co., Ltd.).

**Detection of glucose uptake and lactate production.** Human PC cells (BxPC3 and Panc1) and normal pancreas cell HPC-Y5 in the logarithmic growth phase were inoculated in 6-well plates (5×10<sup>5</sup> cells/well) and cultured in a 37°C incubator overnight. After treatment with a range of leptin concentrations (0, 20, 50 and 100 ng/ml) or according to the grouping aforementioned, the cells were cultured for 3 h in low-glucose DMEM, followed by washing with glucose-free Krebs-Ringer bicarbonate buffer (containing 2% BSA) at 37°C. The cells were subsequently incubated in glucose-free DMEM containing 100 µM 2-NBDG (cat. no. 0467597-16; Cayman Chemical Company) for 45 min, and glucose uptake was evaluated using a 2-NBDG Glucose Uptake Assay kit (Nanjing Jiancheng Bioengineering Institute). Lactate production was evaluated using a Lactate Assay Kit (Nanjing Jiancheng Bioengineering Institute). The supernatant of the treated cells was prepared according to the manufacturer's protocol and the OD was measured at 530 nm using a spectrophotometer.

**RT-qPCR.** Following treatment, total RNA from human PC cells (BxPC3 and Panc1) and normal pancreas cells HPC-Y5 was extracted using TRIzol reagent (cat. no. 1596-026; Invitrogen; Thermo Fisher Scientific, Inc.). Following RNA quantification and confirmation of RNA integrity by electrophoresis with 1% gel, 1 µg of RNA was reverse transcribed into cDNA using the RevertAid First strand cDNA synthesis kit (cat. no. K1622; Fermentas; Thermo Fisher Scientific, Inc.). RT-qPCR was performed in triplicate using an ABI-7300 Real-Time PCR System (Applied Biosystems, Thermo Fisher Scientific, Inc., USA) and the Maxima SYBR Green/ROX qPCR Master Mix kit (cat. no. K0223; Thermo Fisher Scientific, Inc.). GAPDH was used as an internal reference. mRNA expression levels of HKII, GLUT1, LepR-common, LepR-short and LepR-long were analyzed using the 2<sup>-ΔΔCT</sup> method (28). The primers used were as follows: HKII, forward 5'-ACGACATCATTTGTTAAGG-3', reverse 5'-TTTGGCAAAGTGAGGATGTAG-3'; GLUT1, forward 5'-TGCAGGAGATGAGGAAG-3', reverse 5'-CAATGGTGGCATAACACAG-3'; LepR-common, forward 5'-TTGTGCCAGTAATTATTTCTCTT-3', reverse 5'-CACACCAAAGAATGAAAAAGCTAT-3'; LepR-short, forward 5'-TTCCTGGGCACAAGG

Table I. Leptin receptor interference target design results.

Name	Sequence
Leptin receptor target site 1 (842-860)	Sense 5'-GGGUACUGAGGUAACCUAUUU-3' Antisense 5'-AUAGGUUACCUCAGUACCCUU-3'
Leptin receptor target site 2 (863-881)	Sense 5'-GGACGAAAGCCAGAGACAAUU-3' Antisense 5'-UUGUCUCUGGCUUUCGUCCUU-3'

ACTTA-3'- reverse 5'-GCTCCAAAAGAAGAGGACCA-3'; LepR-long, forward 5'-TTCCTGGGCACAAGGACTTA-3', reverse 5'-TTTGTGTCCCTGGGTACTTGA-3'; GAPDH, forward 5'-AATCCCATCACCATCTTC-3', reverse 5'-AGGCTGTTGTCATACTTC-3'. The thermocycling conditions were as follows: 95°C for 10 min; followed by 40 cycles of 95°C for 15 sec and 60°C for 45 sec (29).

**Western blot analysis.** Following the various treatments, total protein from human PC cells (BxPC3 and Panc1) and normal pancreas cells HPC-Y5 were extracted using RIPA buffer supplemented with protease and phosphatase inhibitors (cat. no. R0010; Beijing Solarbio Science & Technology Co., Ltd.), followed by quantification by the BCA Kit (cat. no. PICPI23223; Thermo Fisher Scientific, Inc.). Proteins (25 µg) were separated using 10 (spacer) and 8% (separation) SDS-PAGE, and transferred to PVDF membranes (cat. no. HATF00010; EMD Millipore). Subsequently, the membranes were blocked in 5% skimmed milk (cat. no. BYL40422; BD Biosciences) for 1 h at room temperature and incubated with primary antibodies against GLUT1 (1:1,000; cat. no. ab115730; Abcam), HKII [1:1,000; cat. no. 2867; Cell Signaling Technology (CST)], AKT (1:1,000; cat. no. 2920; CST), phosphorylated (p)-AKT (1:2,000; cat. no. 4060; CST), LepR-long (1:250; cat. no. sc-1835; Santa Cruz Biotechnology, Inc., California, USA), LepR-short (1:250; catalog no. sc-8325; Santa Cruz Biotechnology, Inc.) and GAPDH (1:1,000; cat. no. 5174; CST) overnight at 4°C with gentle agitation. Following 5-6 washes in TBS + 0.1% Tween-20, the membranes were incubated for 2 h at room temperature with horseradish peroxidase-conjugated goat anti-rabbit secondary antibody (1:1,000; cat. no. ZB2301, OriGene Technologies, Inc.). The blots were visualized by ECL reagent (cat. no. WBKLS0100; EMD Millipore) and images were captured with a Tanon-5200 ECL imaging system (Tanon Science and Technology Co., Ltd.). Protein expressions levels were normalized to GAPDH and analyzed using ImageJ 1.47v (National Institutes of Health).

**Statistical analysis.** All statistical analyses and calculations in this study were carried out using GraphPad Prism 7.0 software (GraphPad Software, Inc.). The statistical significance of differences between the two groups was determined using two-tailed Student's t-test, and multiple comparisons were made by one-way ANOVA followed by Tukey's test for multiple comparisons. All experiments were performed in triplicate, and data are expressed as the mean ± standard

deviation.  $P < 0.05$  was considered to indicate a statistically significant difference.

## Results

**Leptin stimulates the proliferation and glucose metabolism of human PC cells.** Human PC cell lines, BxPC3 and Panc1, and normal pancreas cells were treated with increasing concentrations of leptin (0, 20, 50 and 100 ng/ml) for 24 h. *In vitro* treatment with leptin significantly stimulated the proliferation of BxPC3 and Panc1 cells as well as pancreas cells HPC-Y5 in a dose-dependent manner (Fig. 1A). These results indicated that leptin stimulation may promote the proliferation of human PC and pancreas cells. In addition, glucose uptake (Fig. 1B) and lactate production (Fig. 1C) of BxPC3, Panc1 and pancreas cells were markedly increased by leptin stimulation, accompanied by increased expression of HKII and GLUT1 (Fig. 1D and E). These data suggested that leptin stimulation may contribute to glucose metabolism and proliferation of human PC cells and healthy pancreatic cells.

**Silencing of LepR expression levels in BxPC3 cells.** Due to the significant increase of proliferation, glucose uptake and lactate production in both BxPC3 and Panc1 cells, in the present study, one cell line, BxPC3, was selected for further investigation. BxPC3 cells were infected with lentiviruses expressing shNC, LepR-shRNA-1 and LepR-shRNA-2; untreated cells were used as a control. mRNA expression of LepR-common, LepR-short and LepR-long (Fig. 2A), as well as the protein expression levels of LepR-short and LepR-long, were significantly downregulated by LepR-shRNA-1 and LepR-shRNA-2 (Fig. 2B). Therefore, the LepR-shRNA-1 and LepR-shRNA-2 vectors were used for subsequent experiments.

**Silencing of LepR inhibits proliferation and glucose metabolism of human PC cells.** To investigate the role of leptin in human PC cells, proliferation, glucose uptake and lactate production were assessed in BxPC3 cells following LepR silencing. The results indicated that the proliferation of BxPC3 cells was notably suppressed by silencing of LepR, compared with the control groups (Fig. 3A). Glucose uptake (Fig. 3B) and lactate production (Fig. 3C) of BxPC3 cells were also inhibited, accompanied by decreased protein expression levels of HKII, GLUT1 and p-AKT, whereas total AKT protein expression was unaltered (Fig. 3D), suggesting an inhibitory effect of leptin silencing on AKT activation. These results demonstrated the beneficial effects of leptin in glucose metabolism

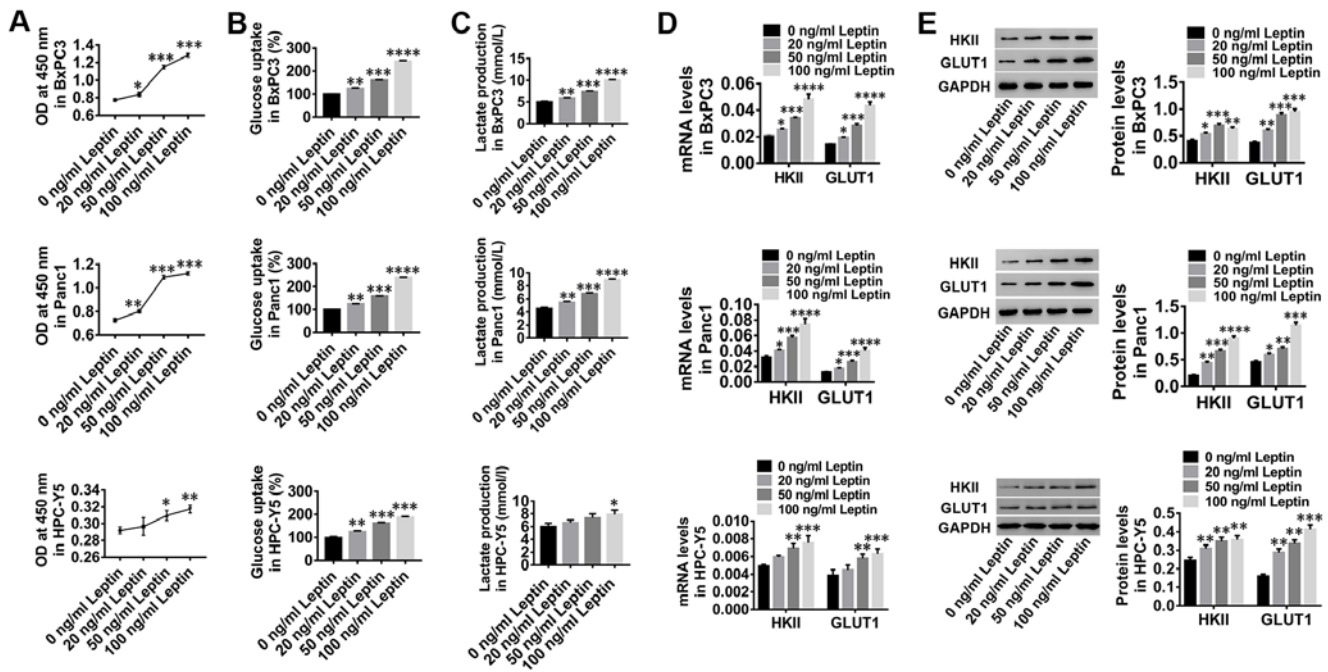


Figure 1. Leptin stimulates the proliferation and glucose metabolism of human PC and normal pancreas cells. (A) The proliferation of BxPC3 (top), Panc1 (middle) and normal pancreas cells HPC-Y5 (bottom) was assessed by Cell Counting Kit-8. (B) Glucose uptake and (C) lactate production were respectively detected by a 2-NBDG kit and a Lactate Assay Kit, respectively. (D) mRNA and (E) protein levels of GLUT1 and HKII were determined by reverse transcription-quantitative PCR and western blotting, respectively. \* $P<0.05$ ; \*\* $P<0.01$ ; \*\*\* $P<0.001$ ; \*\*\*\* $P<0.0001$  vs. 0 ng/ml leptin. GLUT1, glucose transporter 1; HKII, hexokinase II; OD, optical density.

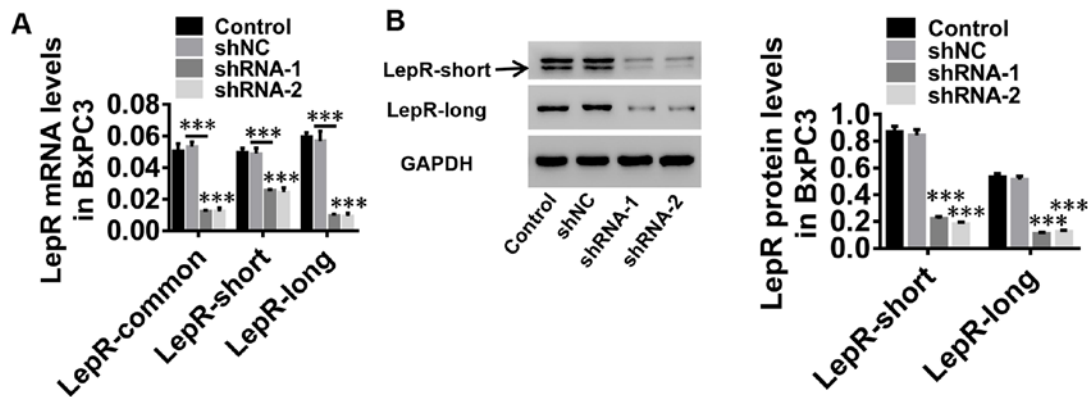


Figure 2. Silencing of LepR expression levels in BxPC3 cells. (A and B) BxPC cells were infected with lentiviruses expressing shNC, LepR-shRNA-1 and shRNA-2; untreated cells were used as control. (A) mRNA expression of LepR-common, LepR-short and LepR-long was detected by reverse transcription-quantitative PCR. (B) Protein levels of LepR-short and LepR-long were detected by western blot analysis. \*\*\* $P<0.001$  vs. shNC. LepR, leptin receptor; shNC, negative control shRNA; shRNA, short hairpin RNA.

and proliferation of human PC cells, which may be involved in AKT pathway activation.

*Leptin stimulates the proliferation and glucose metabolism of human PC cells via activation of the AKT pathway.* The molecular mechanisms of leptin in mediating glucose metabolism and proliferation of human PC cells were explored. Leptin-stimulated proliferation, glucose uptake and lactate production were effectively counteracted by co-treatment with the AKT inhibitor LY294002 (Fig. 4A-C, respectively). The increased protein expression levels of HKII, GLUT1 and p-AKT induced by leptin stimulation were also significantly decreased (Fig. 4D), whereas total AKT was unchanged

(Fig. 4D). Leptin stimulation potentially attenuated the effects of AKT inhibitor, LY294002, suggesting that there was a rescue effect of leptin stimulation on AKT inhibition. In addition, the inhibition of proliferation, glucose uptake and lactate production induced by Lep2-shRNA-2 transfection were significantly counteracted by co-treatment with the AKT activator IGF-1 (Fig. 4E-G, respectively), accompanied by an increased expression of GLUT1 and HKII protein (Fig. 4H). In the present study, Lep2-shRNA-2 was used due to its better downregulation efficiency compared with shRNA-1. These results further demonstrated that leptin may stimulate glucose metabolism and proliferation of human PC cells potentially through activating the AKT pathway.



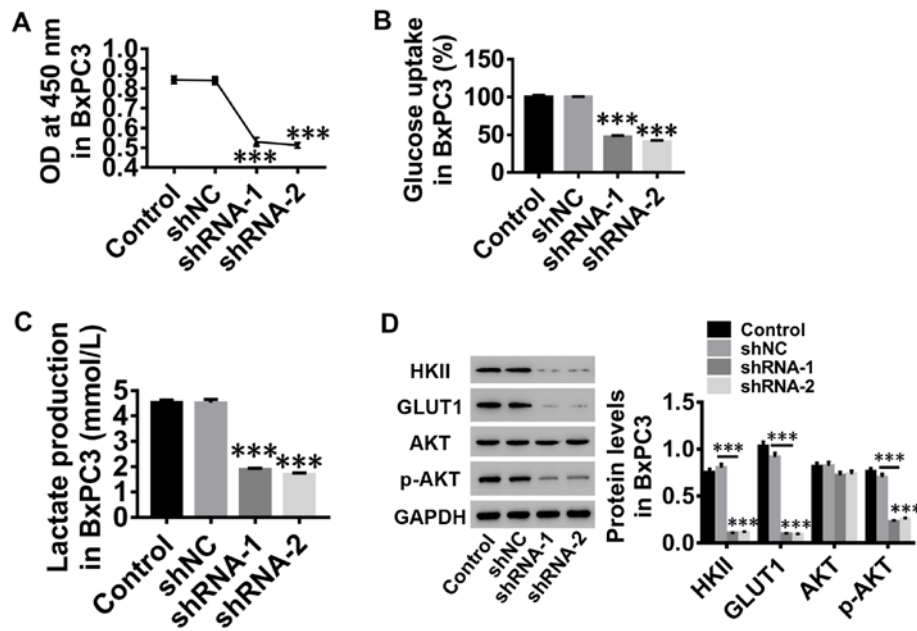


Figure 3. Silencing of LepR reduces the proliferation and glucose metabolism of human PC cells. (A-C) BxPC3 cells were transduced with LepR-shRNA-1 and LepR-shRNA-2. (A) Cell proliferation was assessed by the Cell Counting Kit-8 assay, which demonstrated that LepR silencing reduced cell proliferation. (B and C) Glucose uptake and lactate production were also decreased by the shRNAs. (D) Reduced expression levels of GLUT1, HKII, p-AKT and AKT proteins were detected following lentiviral infection with shRNA. \*\*\*P<0.001 vs. shNC. GLUT1, glucose transporter 1; HKII, hexokinase II; p-AKT, phosphorylated AKT; shNC, negative control shRNA; shRNA, short hairpin RNA.

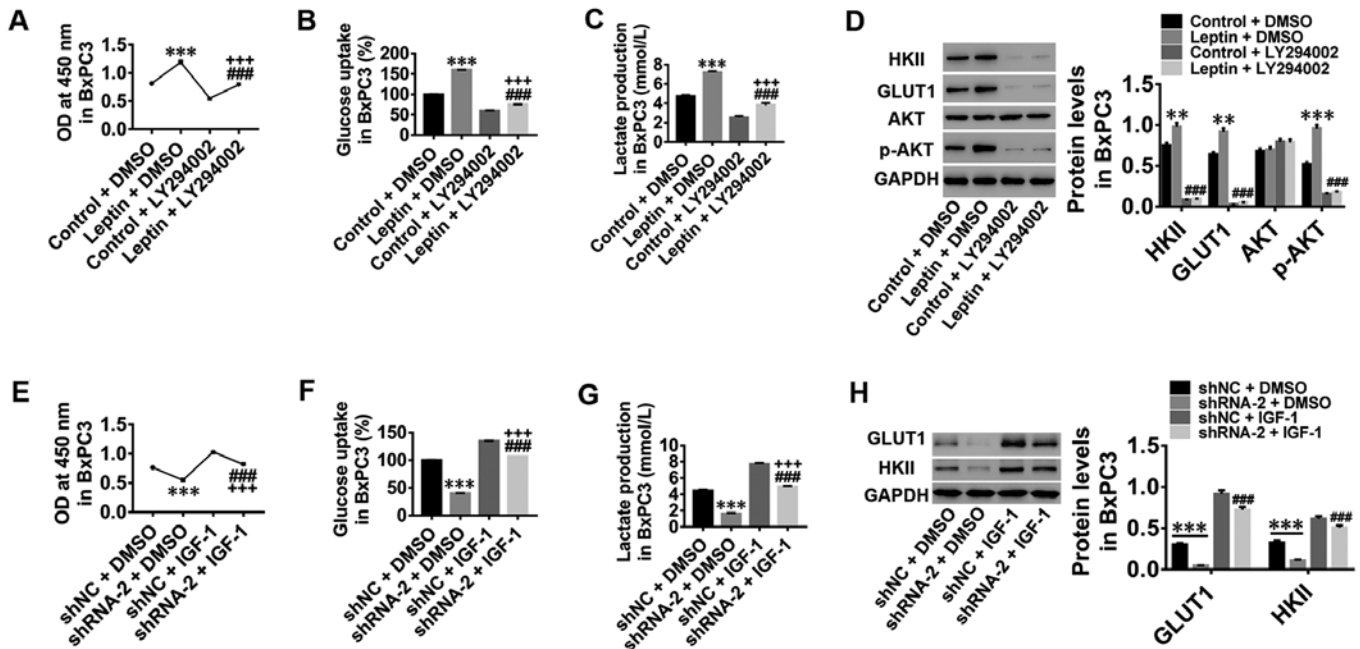


Figure 4. Leptin stimulates the proliferation and glucose metabolism of human PC cells via activation of the AKT pathway. BxPC3 cells were treated with medium (control), 50 ng/ml leptin, medium + 25  $\mu$ mol/l LY294002 (an AKT inhibitor), or 50 ng/ml leptin + 25  $\mu$ mol/l LY294002. (A) Cell proliferation was assessed. (B) Glucose uptake was calculated. (C) Lactate production was calculated. (D) The protein levels of GLUT1, HKII and the p-AKT/total AKT ratio were detected. BxPC3 cells were treated with shNC, shRNA-2, shNC + 50 ng/ml IGF-1 (an AKT activator), and shRNA-2 + 50 ng/ml IGF-1. (E) Cell proliferation was assessed. (F) Glucose uptake was calculated. (G) Lactate production was calculated. (H) The protein levels of GLUT1, HKII were detected. \*\*P<0.01 and \*\*\*P<0.001 vs. control + DMSO or shNC + DMSO; \*\*\*P<0.001 vs. leptin + DMSO or shRNA-2 + DMSO; \*\*\*P<0.001 vs. control + LY294002 or shNC + IGF-1. GLUT1, glucose transporter 1; HKII, hexokinase II; p-AKT, phosphorylated AKT; shNC, negative control shRNA; shRNA, short hairpin RNA.

## Discussion

Leptin is a hormone that is highly expressed in overweight and obese individuals and may be involved in the progression

of many cancers. Previous studies have shown that leptin can stimulate the proliferation of various cancer cell lines (30-33). An increase in leptin was reported in plasma and pancreatic tissue, suggesting that leptin may be involved in tumor growth

in the pancreas (4). The distribution and high abundance of positive leptin or LepR in tumor tissue samples of PC patients have been described in our previous study (27). Leptin acts as a growth factor to promote proliferation in multiple cancers, such as breast and lung cancer (34,35). Leptin stimulates cell proliferation and survival in breast cancer cells through LepR (36,37). In the current study, leptin stimulation promoted the glucose metabolism and proliferation of human PC cells and non-cancerous cells. Silencing of LepR by LepR-shRNAs had the opposite effect as leptin stimulation on human PC cells, which agreed with a previous report (38) and suggested that LepR knockdown may attenuate human PC progression by mediating glucose metabolism and cell proliferation.

The underlying mechanisms of LepR on human PC cell growth were also explored. GLUT1 and HKII are two key enzymes that regulate cellular glycolysis and energy metabolism. HKII has been demonstrated to be the key enzyme to catalyze the first step in the glycolytic pathway (39,40), and GLUT1 typically regulates the first rate-limiting step of glucose metabolism when glucose flows into cells (41,42). The results of the present study demonstrated that GLUT1 and HKII expression in human PC cells and healthy pancreatic cells was significantly increased by leptin stimulation, but LepR silencing in human PC cells significantly decreased GLUT1 and HKII expression. Studies have showed that leptin is a cytokine with multiple biological roles including regulation of energy metabolism and plays an important role in cell proliferation, invasion, metastasis and survival in cancers including breast, esophageal and endometrial cancers (43-46). Thus, the increase of GLUT1 and HKII expression in healthy pancreas cells may be caused by the stimulation of leptin. In our previous study, compared with normal tissues, high abundance of positive leptin or LepR was found in tumor tissue samples of PC patients (27). These results demonstrated that leptin may be a molecular switch that regulates glycolytic-related proteins such as GLUT1 and HKII, thus regulating glucose metabolism of human PC cells. The PI3K/AKT pathway has been suggested to be the main signaling cascade in glucose metabolism and cell growth regulation. It is known that activation of AKT regulates cell growth and controls the rates of glucose uptake through the GLUT1 transporter. It has also been found that AKT can further affect glycolysis through HKII (47-50). Therefore, the AKT pathway was analyzed in the present study. GLUT1, HKII and p-AKT were significantly decreased by the silencing of leptin receptor, whereas AKT levels were unchanged. Additionally, high levels of GLUT1, HKII and p-AKT induced by leptin stimulation were effectively counteracted by co-treatment with the AKT inhibitor LY294002, whereas inhibition of GLUT1, HKII and p-AKT by LepR silencing was counteracted by IGF-1 co-treatment. Increased levels of GLUT1 have been reported to enhance glucose uptake, which in turn elevates the rate of glycolysis to enhance ATP production, ultimately enhancing the growth of tumors (51). These findings demonstrated that LepR may contribute to the glucose metabolism and proliferation of human PC cells *in vitro*, possibly through the activation of the AKT pathway. Compared with our previous studies and other related studies (27,52-54) which reported that leptin enhances PC invasion through the increase in MMP-13 production but does not affect PC cell proliferation, the present study

targeted LepR instead of leptin and revealed that LepR had an effect on the proliferation and metabolism of PC cells and further demonstrated that this regulation may be through the activation of the AKT signaling pathway. Therefore, targeting the LepR/AKT signaling pathway is a potential therapeutic strategy for PC. However, the current results are based solely on *in vitro* experiments and are therefore not comprehensive enough. Further *in vivo* studies should be conducted in the future to confirm these results.

In conclusion, the present study demonstrated that knock-down of LepR may attenuate the development of human PC by inhibiting glucose metabolism and cell proliferation. LepR has the potential to be a molecular switch that regulates glycolysis-related proteins such as GLUT1 and HKII through activation of the AKT pathway, further regulating glucose metabolism and proliferation in human PC cells. These findings offer the foundation for further study of the role of LepR in the relief or treatment of pancreatic cancer.

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### Availability of data and materials

All data generated or analyzed during this study are included in this published article.

### Authors' contributions

WX and HS conceived and designed the study. YX, MT, XT, JuZ, JiZ and JC performed the experiments. WX and HS wrote the manuscript. All authors read and approved the final manuscript.

### Ethics approval and consent to participate

Not applicable.

### Patient consent for publication

Not applicable.

### Competing interests

The authors declare that they have no competing interests.

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