Abstract. The relationship between diabetes mellitus and pancreatic cancer is complex. Diabetes has been postulated to be both an independent risk factor and a consequence of pancreatic cancer. Our previous study confirmed that curcumin plays a vital role in inhibiting the epithelial-mesenchymal transition of pancreatic cancer cells. However, whether curcumin attenuates hyperglycemia-induced cancer invasive and migratory abilities and the underlying mechanisms are not yet well understood. As high glucose is able to induce the expression of epidermal growth factor (EGF), which is intimately related with tumor progression, the aim of this study was to evaluate whether curcumin could influence the high glucose-induced EGF/EGFR pathway and the biological activity of pancreatic cancer cells. Human pancreatic cancer BxPC-3 cells were exposed to high glucose or EGF, with or without curcumin, LY 294002 (an Akt inhibitor) or PD 98059 (an ERK inhibitor). MTT, Transwell invasion and wound healing assays were used to detect the proliferation, invasion and migration potential of cancer cells. The activation of p-EGFR, p-ERK and p-Akt was determined by western blot analysis. The expression levels of uPA and E-cadherin were examined using qRT-PCR and western blot analysis. The results showed that high glucose could not only promote the proliferation, invasion and migration of pancreatic cancer cells, but also induce the expression of EGF and activation of EGFR, ERK and Akt. These effects of high glucose were counter-balanced by curcumin. EGF-induced proliferative, invasive and migratory abilities of BxPC-3 cells were abrogated by curcumin, LY 294002 and PD 98059. In addition, EGF-modulated activation of EGFR, ERK and Akt, as well as the expression of uPA and E-cadherin were inhibited by curcumin. Taken together, the present study indicates that curcumin suppresses hyperglycemia-driven EGF-induced invasion and migration of pancreatic cancer cells by inhibiting the EGF/EGFR signaling pathway and its downstream signaling molecules including ERK and Akt. Curcumin is a potential anticancer agent for pancreatic cancer.

Introduction

Pancreatic cancer is a highly aggressive and fatal malignant digestive tumor for which the 5-year relative survival rate is 8%. The poor prognosis of this severe disease is mainly due to the local invasion and distant metastasis at an early stage. In 2018, 55,440 individuals were estimated to be newly diagnosed with pancreatic cancer, which may result in more than 44,000 cancer-related deaths in the United States (1). In China, pancreatic cancer accounts for the second leading upward trend of age-standardized mortality rates from 2000 to 2011 (2). Routine treatments for this severe disease include surgery, radiation and chemotherapy. However, at present, approximately 80% of pancreatic cancer patients who are diagnosed do not qualify for surgical resection due to early relapse or metastatic spread of the disease. Therefore, the exploration of risk factors and novel effective therapeutic options are urgently needed to improve the treatment outcome in patients with pancreatic cancer (3).

Diabetes mellitus could be both a risk factor and a consequence of pancreatic cancer. A collaborative analysis from the International Pancreatic Cancer Case-Control Consortium analyzed more than 8,000 pancreatic cancer cases and confirmed that individuals with diabetes mellitus have an almost 2-fold increased risk of developing pancreatic cancer (4). In a Chinese retrospective cohort study, both male and female type 2 diabetes mellitus patients had an increased risk to develop pancreatic cancer compared with the general population with the standardized incidence ratios of 2.973 and 2.687, respectively (5). Diabetes mellitus improves following
pancreatectomy, suggesting that diabetes may be induced by pancreatic cancer (6). Our previous studies confirmed that a hyperglycemic condition could promote the proliferation, invasion, epithelial-mesenchymal transition (EMT) as well as the metastasis of pancreatic cancer (7,8).

Epidermal growth factor (EGF) is a polypeptide that regulates various cellular functions involving proliferation, survival, differentiation, angiogenesis and metastasis in many types of cancer. EGF binds with a high affinity receptor located in the cellular membrane and stimulates rapid activation of protein kinase activity (9). Overexpression of epidermal growth factor receptor (EGFR) and ligands has been shown in pancreatic ductal adenocarcinoma and pancreatic cancer cells, which is essential for the initiation and progression of this disease (10).

In our previous study, it was shown that high glucose could promote pancreatic cancer cell proliferation via the induction of EGF expression and transactivation of EGFR (11). Curcumin, a natural polyphenol compound derived from turmeric, has multiple biologic properties, including health maintenance and cancer prevention (12). Curcumin exhibits its antitumor effects via targeting multiple signaling pathways, including mitogen-activated protein kinase (MAPK), protein kinase B (Akt) and others (13). In a recent study, it was indicated that curcumin plays an important role in suppressing superoxide dismutase-induced EMT of pancreatic cancer by inhibiting the PI3K/Akt/NF-κB signaling pathway (14). It was also confirmed that curcumin could restrain hypoxia-induced EMT via suppression of the hedgehog signaling pathway in pancreatic cancer cells (15). However, whether curcumin is able to inhibit high glucose-induced progression of pancreatic cancer and the related mechanisms have not yet been elucidated.

In the present study, the hypothesis that curcumin is able to inhibit high glucose-driven EGF-induced invasive and migratory abilities in pancreatic cancer cells was tested. The effect of curcumin on high glucose-induced activation of EGFR, extracellular signal-regulated kinase (ERK) and Akt signaling pathways was also investigated. In addition, the effect of curcumin on EGF-modulated activation of EGFR, ERK and Akt, as well as the expression of uPA and E-cadherin was demonstrated. Results from this study suggest that curcumin is a potential anticancer agent for the therapy of pancreatic cancer via the suppression of the EGF/EGFR signaling pathway and its downstream signaling molecules including ERK and Akt.

**Materials and methods**

**Cell culture and reagents.** Human BxPC-3 pancreatic cancer cell line was obtained from the American Type Culture Collection (ATCC; Manassas, VA, USA). The cells were cultured in Gibco™ DMEM (Thermo Fisher Scientific, Inc., Waltham, MA, USA), which contained 10% dialized heat-inactivated FBS, 100 U/ml penicillin as well as 100 µg/ml streptomycin in a humidified atmosphere of 5% CO₂ at 37°C. Exponentially growing cells in complete medium were treated with 50 ng/ml EGF, 20 µM curcumin, 10 µM LY 294002 (a PI3K inhibitor), and/or 50 µM PD 98059 (an ERK inhibitor) in normal culturing conditions (5.5 mM glucose) or high glucose (25 mM) conditions for indicated time intervals according to the aim of the experiment. Curcumin, EGF, PD 98059 and LY294002 were purchased from Sigma-Aldrich; Merck KGaA (Darmstadt, Germany). Millicell® culture plate inserts for the Transwell assay were obtained from EMD Millipore (Billerica, MA, USA). Matrigel was purchased from BD Biosciences (Bedford, MA, USA). Primary antibodies [dilution at 1:100 in phosphate-buffered saline (PBS)-Tween-20] against EGFR (cat. no. sc-374255), E-cadherin (cat. no. sc-52328) and uPA (cat. no. sc-59727) were obtained from Santa Cruz Biotechnology (Santa Cruz, CA, USA). The anti-EGFR (cat. no. 4267), anti-phospho-EGFR (Tyr 1068, cat. no. 2234), anti-Akt (cat. no. 9272), anti-phospho-Akt (Ser473, cat. no. 4060), anti-ERK (cat. no. 9102) and anti-phospho-ERK (Thr202/Tyr204, cat. no. 9106) were purchased from Cell Signaling Technology (Beverly, MA, USA). Other reagents were purchased from common commercial sources. All drug solutions were freshly prepared on the day of testing.

**MTT proliferation assays.** BxPC-3 cells were seeded in 96-well plates at the density of 1x10⁴ cells per well. The cells were treated with curcumin, PD 98059 or LY 294002 in EGF or a high glucose condition. After incubation for 24 h at 37°C, 15 µl of MTT solution was added to each well and the cells were incubated for 4 h at 37°C. DMSO (100 µl) was then added to each well. The optical density (OD) value at 490 nm was determined using a spectrophotometer (Bio-Rad Laboratories, Inc., Hercules, CA, USA).

**Transwell Matrigel invasion assay.** The 8.0-µm pore inserts of the Transwell chambers were coated with 30 µl of Matrigel. After serum starvation for 24 h, BxPC-3 cells were suspended in the top chamber at a concentration of 5x10⁴ in DMEM containing 1% FBS in the absence or presence of high glucose, EGF, curcumin, PD 98059 and/or LY 294002. Simultaneously, in the lower chambers, 500 µl of DMEM containing 20% FBS was placed. The Matrigel invasion chamber was incubated for 48 h in a humidified tissue culture incubator. The non-invading cells were then removed from the upper surface. After staining with crystal violet, the stained cells on the bottom surface were counted under an Nikon inverted microscope DIAPHOT-TMD (Nikon, Tokyo, Japan) to test the invasion ability of the cancer cells. Three random fields were captured at x20 magnification (n=3).

**Wound healing assay.** BxPC-3 cells (1.0x10⁵ cells/500 µl) were seeded in 24-well plates. A sterile pipette tip was used to produce a wound line between the cells that grew to 90-100% confluence and cellular debris was removed. Cells were allowed to migrate for 24 h. Images were captured at time 0 and 24 h post-wounding under a Nikon DIAPHOT-TMD microscope (x10 magnification). The relative distance traveled by the leading edge from 0 to 24 h was assessed using Photoshop software (Adobe Systems, Inc., San Jose, CA, USA) (n=5).

**Real-time quantitative PCR (qRT-PCR).** After being extracted from the BxPC-3 cells using the Fastgen 200 RNA isolation system (Fastgen, Shanghai, China) according to the manufacturer's protocol, the total RNA was reverse-transcribed into cDNA using the Fermentas RevertAid™ kit (MBI Fermentas, Burlington, ON, Canada). The primer sequences...
were as follows: uPA-F, 5′-TAA GAG CTG GTT GAT TG-3′ and uPA-R, 5′-TTG GAT GAA CTA GGC TAA AA-3′; E-cadherin-F, 5′-ATTCCTGA TTTGCTGCTCTGGT-3′ and E-cadherin-R, 5′-AGTCCTGGTCTCCTCTCC-3′; β-actin-F, 5′-GATTAGTGTGCTTACCCCTTCT-3′ and β-actin-R, 5′-GAACGGT GAAGGTGACAGCAGT-3′. The PCR reactions consisted of 30 sec at 95˚C, followed by 40 cycles of 95˚C for 5 sec, 60˚C for 30 sec and 72˚C for 30 sec. After each qRT-PCR experiment, a dissociation curve analysis was conducted. The relative gene expression was calculated using the previously described 2-∆∆Cq method (16).

Western blot analysis. Total protein from BxPC-3 cells was extracted from cultured cells in Radio-immunoprecipitation assay (RIPA) lysis buffer on ice for 25 min. Insoluble materials were removed by centrifugation at 4˚C with 15,000 x g for 15 min. Subsequently, supernatants were collected and total protein concentrations were measured using the BCA assay kit (Pierce, Rockford, IL, USA). Clarified protein lysates (30-80 µl) from the BxPC-3 cancer cells were electrophoretically resolved on a denaturing SDS-polyacrylamide gel (10-12%) and electrotransferred onto polyvinylidene difluoride (PVDF) membranes. The membranes were initially blocked with 5% non-fat dry milk in Tris-buffered saline (TBS) for 2 h and then probed with antibodies against E-cadherin, uPA, EGF, EGFR, p-EGFR, Akt, p-Akt, ERK, p-ERK and β-actin (Control). After incubation with the primary antibodies at 4˚C overnight, the membranes were hybridized with secondary goat anti-mouse (cat. no. 7056) or goat anti-rabbit (cat. no. 7074) antibodies at 1:1,000 dilution (Cell Signaling Technology, Beverly, MA, USA) for 2 h at room temperature. Immunopositive bands were developed using an enhanced chemiluminescence (ECL) detection system (Amersham, Piscataway, NJ, USA). Immunodetection was visualized on a Gel Doc 2000 Imaging System (Bio-Rad Laboratories). All analyses were conducted in triplicate. The data were analyzed as (p-protein/control)/(total protein/control).

Statistical analysis. Statistical analysis was performed using SPSS software (version 17.0; SPSS Inc., Chicago, IL, USA). Data are presented as the means ± SEM of three replicate assays. Differences between the groups were analyzed by analysis of variance [ANOVA followed by least significant difference (LSD)]. Statistical significance was set at P<0.05. All experiments were repeated independently at least three times.

Results

Curcumin inhibits high glucose and EGF-induced proliferation in pancreatic cancer cells. Our previous study demonstrated that a hyperglycemic condition promotes the proliferation of pancreatic cancer cells via the induction of EGF expression as well as the transactivation of EGFR (11). As our previous studies found that curcumin shows a 50% inhibitory concentration (IC50) of ~20 µM and this concentration exhibited no cytotoxic effects in BxPC-3 cells, the treatment concentration of 20 µM of curcumin was used in the present experiments (17). Here, it was shown that both a high glucose condition and EGF stimulation accelerated the growth of BxPC-3 cells, which was able to be counter-balanced by curcumin. In addition both PD 98059 and LY 294002 inhibited the high glucose- and EGF-induced cellular proliferation, which indicated that the growth of tumor cells was mediated via the ERK and Akt pathways (Fig. 1).

Curcumin inhibits high glucose-induced invasive ability and wound closure of pancreatic cancer cells. Invasion and migration of cancer cells are considered detrimental aspects in the development of tumor metastasis. In order to confirm whether curcumin influences high glucose-induced pancreatic cancer cell invasion and migration, Transwell invasion and wound-healing assays were conducted. As shown in Fig. 2, the average cell numbers that invaded into the lower chamber were significantly increased in the high glucose condition after incubation for 48 h and this increase was reversed by co-treatment with curcumin. The migratory ability of BxPC-3 cells after incubation for 24 h was also suppressed by curcumin.

Curcumin downregulates high glucose-induced activation of EGF/ERK and EGF/Akt pathways. In order to form metastases, cancer cells undergo various steps, including local invasion, intravasation, survival in the circulation, arrest at distant organ site, micrometastasis formation and metastatic colonization (18). Different molecular pathways are activated to pass these important steps, such as the ERK pathway and PI-3K/Akt pathway (18). The ERK pathway, which belongs to the mitogen-activated protein kinase (MAPK) signaling pathway, is involved in tumor proliferation, differentiation, migration and invasion (19). The PI3K/Akt pathway promotes cancer development, angiogenesis as well as metastasis (20).

Our previous study confirmed that high glucose could promote the expression of EGF and transactivation of EGFR (11). Hyperglycemic conditions could also activate the ERK and p38 MAPK signaling pathways via the production of H2O2 in
Figure 2. Effects of curcumin on the high glucose-induced invasive and migratory ability of cancer cells. Pancreatic cancer cells were exposed to normal glucose (5.5 mM), high glucose (25 mM) condition as well as 20 µM curcumin for 24 or 48 h. The number of migrated cells was quantified by counting the number of cells at x200 magnification after 48 h treatment. The confluent monolayer was wounded with a sterile pipette tip and the cells were allowed to migrate for 24 h. *P<0.05 as compared with the 5.5 mM glucose group; #P<0.05 as compared with the 25 mM glucose group.

Figure 3. Role of curcumin in high glucose-induced EGF expression and phosphorylation of EGFR, ERKs as well as Akt. BxPC-3 cells were treated with 20 µM curcumin for 24 h under high glucose (25 mM) or normal glucose (5.5 mM) condition to evaluate the expression of EGF and the phosphorylation levels of EGFR, ERK and Akt. *P<0.05 as compared with the 5.5 mM glucose group; †P<0.05 as compared with the 25 mM glucose group. EGF, epidermal growth factor; EGFR, epidermal growth factor receptor; ERK, extracellular signal-regulated kinase; Akt, protein kinase B.
pancreatic cancer (21). In this study, it was shown that a high glucose condition activated ERK and Akt signaling pathways, as both ERK and Akt phosphorylation were strongly increased. Curcumin was able to suppress the expression of EGF and activation of EGFR, ERK and Akt in the high glucose condition (Fig. 3).

In order to assess whether the high glucose-induced activation of ERK and Akt signaling pathways was EGF/EGFR-dependent, BxPC-3 cells were treated with EGF. As shown in Fig. 4, EGF significantly promoted the phosphorylation of EGFR, ERK and Akt, while curcumin was able to counter-balance these effects of EGF.

**Curcumin inhibits EGF-induced invasive ability of pancreatic cancer cells.** Growth factors, such as EGF and vascular endothelial growth factor (VEGF), are intimately related with cancer cell migration, angiogenesis, regulation of cell adhesion and EMT (22). Activation of ErbB ligands and overexpression of EGFR significantly promote tumor metastasis via chemotaxis and invasation (23).

In order to confirm whether curcumin inhibits EGF-induced BxPC-3 cell invasion, a Transwell invasion assay was utilized. As shown in Fig. 5, the number of cells that invaded into the lower chamber was increased with the addition of EGF after incubation for 48 h. This increase was reversed by co-treatment with curcumin. In addition, both PD 98059 and LY 294002 also suppressed the effect of EGF which indicated that EGF-induced invasion was related to the ERK and Akt pathways.

**Curcumin suppresses EGF-induced wound closure of pancreatic cancer cells.** A classic wound healing assay was used to evaluate the effect of curcumin on EGF-induced BxPC-3 cell motility. The results showed that EGF significantly increased the migratory ability of the BxPC-3 cells after incubation for 24 h. Curcumin counter-balanced these effects of EGF. Both PD 98059 and LY 294002 inhibited EGF-induced wound closure of pancreatic cancer cells. Curcumin exerts its inhibitory effect on cellular motility, which might be attributed to the suppression of the EGF/ERK and EGF/Akt pathways (Fig. 6).
Curcumin inhibits EGF-modulated expression of metastatic-related factors. Our previous study demonstrated that a high glucose condition could induce the invasive and migratory abilities of pancreatic cancer cells by regulating metastatic-related factors, including uPA and E-cadherin (24). In the present study, it was shown that EGF direct stimulation also promoted the mRNA expression of uPA and down-regulated the mRNA level of E-cadherin. Curcumin significantly reversed these EGF-induced effects (Fig. 7). In addition, curcumin also reversed the EGF-modulated metastatic-related factors at the protein level, and the trend was consistent with the mRNA results. Taken together, these results indicate that curcumin inhibits invasion and migration of pancreatic cancer cells under high glucose conditions, which maybe attributed to the EGF/ERK and EGF/Akt signaling pathways.

Discussion

Due to local recurrence, lymphnode and liver metastases as well as peritoneal dissemination, pancreatic cancer is one of the most aggressive malignant diseases, the hallmarks of which includes poor outcome, short survival duration and resistance to therapy (25). In China, pancreatic cancer is the seventh deadliest disease with annual mortality rates almost equal to incidence rates, of which the 5-year relative survival rate is 4.1% and the median survival time is 3.9 months (6). The exploration of risk factors and newer effective therapeutic options are important to improve the treatment outcome for pancreatic cancer patients. Diabetes mellitus can be both a risk factor and a consequence of pancreatic cancer. Our previous study confirmed that high glucose could promote pancreatic cancer cell proliferation via the induction of epidermal growth factor (EGF) expression and transactivation of the epidermal growth factor receptor (EGFR) (11). Hyperglycemic conditions could worsen the prognosis of pancreatic cancer by enhancing invasive ability and promoting epithelial-mesenchymal transition (EMT) through the production of hydrogen peroxide (8). In addition, diabetes mellitus was also found to enhance perineural invasion in pancreatic cancer patients and to aggravate a poor prognosis (7). It has been verified that the addition of 5.5 and 25 mM glucose resulted in balanced osmotic pressure inside and outside the cell membrane (21), thus we used 5.5 and 25 mM of glucose as it has been applied worldwide in the related field as a classical high glucose model in the cell culture. In the present study, we focused on whether curcumin is able to suppress high glucose-induced cancer proliferation, migratory and invasive abilities and the underlying mechanism.

Our data showed that a high glucose condition could promote the proliferation, migration and invasion of pancreatic cancer cells. High glucose was not only able to increase the expression of EGF and transactivation of EGFR, but also activate the ERK and Akt signaling pathways. Curcumin was able to abrogate these effects of a high glucose condition. In addition, in order to ascertain whether the effect of a high glucose condition on pancreatic cancer cells was EGF-dependent, we also treated BxPC-3 cells with EGF directly. Data showed that EGF treatment significantly promote the proliferative, migratory and invasive abilities of the BxPC-3 cells. EGF treatment also modulated the expression levels of p-EGFR,
p-ERK, p-Akt, uPA and E-cadherinin pancreatic cancer cells. Curcumin was also able to suppress the effects of EGF treatment. The addition of PD 98059 and LY 294002 to the cell culture resulted in the inhibition of cellular growth and invasion. Our results indicate that curcumin inhibits high glucose-induced invasion and migration of pancreatic cancer cells, which might be attributed to the EGF/ERK and EGF/Akt signaling pathways.

EGF belongs to the EGF family, which includes EGF, transforming growth factor α (TGFA), amphiregulin, betacellulin, cripto, heparin-binding EGF (HB-EGF), epigen, epiregulin and neuroglial EGF. EGF exerts its biological effects by binding to the receptors in an autocrine or paracrine manner. The ErbB family consists of four receptors: EGFR (ErbB-1), ErbB-2, ErbB-3 and ErbB-4. After binding of EGF, the four members of the family are able to form various heterodimers, which lead to autophosphorylation and further activated downstream signaling pathways, including the PI3K/Akt pathway and MAPKs, such as the ERK pathway (9). The signaling cascades activated by EGF/EGFR could further regulate different cellular responses such as proliferation, survival, migration, invasion, intravasation and metastasis (26).

The ERK MAPK signaling pathway is involved in the regulation of several cancer cell processes, including proliferation, differentiation, survival, cellular motility and metastasis. Aberrant signaling drives tumor initiation and progression. The PI3K/Akt/mTOR signaling pathway also promotes cancer development, migration, invasion, angiogenesis and metastasis (18). Overexpression or mutational activation of EGFR has been found in cancer cells leading to activation of the ERK and PI3K/Akt signaling pathways (27). Various EGFR inhibitors have been developed for cancer treatment, including EGFR monoclonal antibodies (cetuximab, panitumumab) and tyrosine kinase inhibitors, including erlotinib, gefitinib and lapatinib. These drugs have been successfully approved for the treatment of metastatic colorectal cancer (cetuximab and panitumumab), locally advanced, unresectable, or metastatic pancreatic cancer (erlotinib), breast cancer (lapatinib) and non-small cell lung cancer (gefitinib and erlotinib) (28). In the present study, it was found that high glucose and EGF stimulation could activate EGFR, ERK and Akt, which further induce cancer proliferation, migration and invasion. As a type of natural polyphenol compound derived from turmeric, curcumin was able to abrogate these effects leading to cancer prevention.

Accumulating evidence suggests that curcumin inhibits cancer initiation promotion and progression through regulation of multiple signaling pathways including EGF/EGFR, Akt/mTOR, NF-κB, Notch as well as MAPK pathways (29). Soung and Chung (30) showed that curcumin was able to inhibit cancer cell functions by disrupting the interaction between integrin α6β4 and EGFR. Cheng et al (12) demonstrated that curcumin could exhibit inhibitory effects on the proliferation, invasion and metastasis of prostate cancer via inhibition of MMP-9 activity, downregulation of cellular matrix metalloproteinases, suppression of the effects of androgens and EGFR ligands on the protease and promotion of protease shedding. Le et al (30) found that targeting EGFR with EGF-conjugated curcumin liposomes increased the antitumor activity of curcumin against pancreatic cancer cells. Treatment with curcumin effectively attenuated tobacco smoke-induced activation of the ERK and JNK MAPK pathways as well as EMT alterations in the mouse liver (31). Curcumin was also found to weaken pancreatic cancer cell growth, clonogenic potential, migration and invasion abilities by downregulation of the expression of two homologous transcriptional co-activators, YAP (Yes-associated protein) and TAZ (transcriptional coactivator with PDZ-binding motif) via the Notch signaling pathway (32). As an activator of the ligase anaphase-promoting complex/C (APC/C), cell division cycle 20 (Cdc20) plays an oncogenic role in tumorigenesis.

A recent study found that curcumin could suppress cell growth, induce apoptosis and trigger cell cycle arrest via inhibition of Cdc20 expression in pancreatic cancer cells (33). S-phase kinase associated protein 2 (Skp2) plays an oncogenic role in pancreatic cancer development and progression. Su et al (34) demonstrated that curcumin-induced cellular proliferation inhibition, cell cycle arrest, apoptosis and invasion suppression in pancreatic cancer cells was partly attributed to
the downregulation of Skp2. Curcumin exerts its anticancer effects both alone and in combination with other anticancer drugs, including gemcitabine and 5-fluorouracil, by modulating multiple therapeutic molecular targets and signaling pathways. For instance, curcumin sensitized gemcitabine-resistant cancer cells by inhibiting the expression of the PRC2 subunit EZH2 and its related IncRNA PVT1 (35). We demonstrated that curcumin suppressed hydrogen peroxide-induced cell migration and invasion through the inhibition of the ROS/ERK/NF-κB signaling pathway (17). Curcumin was also found to inhibit superoxide dismutase-induced cancer EMT via the suppression of the PI3K/Akt/NF-κB signaling pathway in pancreatic cancer cells (14). In the present study, our data showed that curcumin inhibited the invasive and migratory abilities of pancreatic cancer cells under high glucose conditions, which maybe attributed to the EGF/ERK and EGF/Akt signaling pathways. Our future experiments may include more in vivo experiments to further validate the findings.

In conclusion, the present study demonstrated that curcumin plays an important role in suppressing high glucose-induced proliferative, invasive and migratory abilities of pancreatic cancer cells via inhibiting the EGF/ERK and EGF/Akt signaling pathways. Curcumin might be a potential anticancer agent for the treatment of pancreatic cancer patients.

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Availability of data and materials

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

Authors' contributions

QM, ZWa and LC conceived and designed the study; WL, LH and XC conducted the experiments; and ZWu and LC performed the data analysis. WL wrote the paper. LC and QM reviewed and edited the manuscript. 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

Not applicable.

Patient consent for publication

Not applicable.

Competing interests

The authors state that they have no competing interests.

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