Hedgehog Signaling Regulates Epithelial-Mesenchymal Transition in Pancreatic Cancer Stem-Like Cells

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Abstract

Hedgehog (Hh) signaling is crucially involved in tumorigenesis. This study aimed to assess the role of Hh signaling in the regulation of epithelial-mesenchymal transition (EMT), stemness properties and chemoresistance of human pancreatic Panc-1 cancer stem cells (CSCs). Panc-1 cells were transfected with recombinant lentiviral vectors to silence SMO and serum-free floating-culture system was used to isolate Panc-1 tumorspheres. The expression of CSC and EMT markers was detected by flow cytometry, real-time RT-PCR and Western blot analysis. Malignant behaviors of Panc-1 CSCs were evaluated by tumorigenicity assays and nude mouse lung metastasis model. We found that tumorspheres derived from pancreatic cancer cell line Panc-1 possessed self-renewal, differentiation and stemness properties. Hh pathway and EMT were active in Panc-1 tumorspheres. Inhibition of Hh signaling by SMO knock-down inhibited self-renewal, EMT, invasion, chemoresistance, pulmonary metastasis, tumorigenesis of pancreatic CSCs. In conclusion, Hh signaling contributes to the maintenance of stem-like properties and chemoresistance of pancreatic CSC and promotes the tumorigenesis and metastasis of pancreatic cancer. Hh pathway is a potential molecular target for the development of therapeutic strategies for pancreatic CSCs.

Key words: Hedgehog, pancreatic cancer, cancer stem cell, epithelial-mesenchymal transition

Introduction

Pancreatic cancer (PC) is an aggressive malignancy with a 5-year survival rate < 6% [1]. Despite recent advances in the diagnosis, surgery and chemotherapy of PC, the prognosis of PC patients remains poor. At an advanced stage, 80-85% of PC patients are diagnosed when surgical resection is not feasible [2]. Therefore, understanding the molecular events underlying the development and progression of pancreatic cancer is urgently imperative to the development of novel therapeutic strategies.

Increasing evidence indicates that tumors contain a small subpopulation of cancer stem cells (CSCs) [3]. Cancer stem cells (CSCs) indicate multipotent cells exhibiting self-renewal capacity, multilineage differentiation ability and high carcinogenesis, and have been identified in several types of human cancers including pancreatic cancer [4-6]. The highly tumorigenic CSCs exhibit self-renewal by activating developmental pathways such as Wnt, Hedgehog (Hh), and Notch that participate in the maintenance of stemness [7]. As tumorigenic drivers, CSCs should be effectively targeted to achieve long-lasting therapeutic responses.

Recent studies indicate that epithelial-mesenchymal transition (EMT), a developmental process in which cells lose epithelial characteristics and acquire mesenchymal properties such as increased motility and invasion, can endow cells with stem-cell like characteristics. Hh signaling functions in the development and homeostasis of many organs...
and tissues. Aberrant activation of Hh signaling is associated with multifarious human tumors where the pathway is implicated in tumorigenesis, malignancy, metastasis and cancer stem cells [8]. Previous studies have shown that the activation of Hh pathway was a critical event in the tumorigenesis of pancreatic cancer [9,10].

In this study we aimed to assess the role of Hh signaling in the regulation of EMT, stemness properties and chemoresistance of pancreatic CSCs. We isolated human pancreatic Panc-1 tumorspheres and employed recombinant lentiviral vectors for the delivery of siRNA to knockdown SMO. We examined the expression of CSC and EMT markers and evaluated the malignant behaviors of Panc-1 CSCs in vitro and in vivo using nude mouse lung metastasis model.

Materials and methods

Reagents

Gemcitabine was obtained from Lilly. Fetal bovine serum (FBS) was from Gibeco. Antibodies for SMO, Snail, ALDH1A1 and N-cadherin were purchased from Abcam. Antibodies for E-cadherin were purchased from CST. Antibodies for Flow cytometry were purchased from Miltenyi. Cell Counting kit-8 were purchased from Dojindo, Kumamoto.

Cell culture

Pancreatic cancer cell line Panc-1 was received from the American Type Culture Collection and cultured in DMEM supplemented with 10% FBS, penicillin (100 U/ml) and streptomycin (100 µg/ml). A retroviral packaging cell line (ATCC 293T) was maintained in DMEM supplemented with 10% FBS. All cell lines were maintained at 37°C in a humidified 5% CO2 atmosphere.

Lentiviral-mediated delivery of siRNA

The RNAi targeting the human SMO gene (GenBank accession no.NM_005631.4) and the negative control sequence were designed and constructed by Invitrogen (Shanghai, China). The sequences of two different oligos targeting SMO were as follows: si-1

TGCTGTACACGACGATGAGTGGTGGCCACTGACATCTGGTCTGTA and CCGTATACGAGATAGGATGTCGAGTCCTAGTCGTA

GGCCAAAACCATCATCCTTTGTACGTA; si-2

TGCTGTACGATGACGAGATGAGATGGTGGCCACTGACATCTGGTCTGTA and CCGTATACGAGATAGGATGTCGAGTCCTAGTCGTA

GGCCAAAACCATCATCCTTTGTACGTA.

Lentiviral harboring luciferase reporter gene was generated using pBJ-luciferase, psPAX2 and pMD2.G (Addgene) in 293T packaging cells. Panc-1-NC and Panc-1-si cells were seeded in 6-well plates and transfected using the Lipofectamine system according to the manufacturer’s protocol when cells achieved about 50% confluence. Cells were selected with 1 ug/ml of puromycin for 2 weeks.

Real-time RT-PCR

Total RNA was extracted from cells using TRIZOL Reagent (Invitrogen). First strand cDNA was synthesized by using the ReverTra Ace® qPCR RT kit according to the manufacturer’s protocol (TOYOBO). Real-time RT-PCR was performed by using SYBR Green PCR Master Mix (TOYOBO). GAPDH was used as an internal control. The primer used were as follows: GAPDH 5’-ACGGATTTGGTGTAGTG-3’ and 5’-TGAATTTGGAGGAGCTGC-3’; SMO 5’-TGCTCCTGTGGGAGGCTACTT-3’ and 5’-ATCTTGCTGGGACGCTACTAC-3’; Gli1 5’-TGCTGGGACGCTACTAC-3’; Shh 5’-CCAATTACAACCCCGACATC-3’ and 5’-CATGGGAGGCCACTG-3’. Relative quantification of gene expression was calculated on triplicate reactions using the \( \triangle \Delta Ct \) method. Data were analyzed by ABI 7500 software v2.0.6 (Applied Biosystems).

Western blot analysis

Cells were lysed in lysis buffer containing 1 mmol/L Na3VO4, 1 mmol/L phenylmethylsulfonylfluoride (PMSF), 500 uM AEBSF, 150 mM Aprotinin, 1 uM E-64, 0.5 mM EDTA- Na and 1 uM leupeptin. The lysates were cleared by centrifugation at 14,000 rpm and total protein concentration was measured with bichinchoninic acid assay Kit (Bio-Rad).
Laboratories). Cell lysates were separated in 10% SDS-PAGE and transferred onto polyvinylidene difluoride membranes (Millipore). The membranes were blocked with 5% nonfat dry milk in Tris-buffered saline with 0.1% Tween 20, incubated with primary antibodies, and then with HRP-conjugated secondary antibodies. The peroxidase activity was detected by using Immobilon Western Chemiluminescent HRP Substrate (Millipore) and exposed to X-ray films. β-actin was used as loading control.

Flow cytometry
To detect the expression of cancer stem cell markers, the following antibodies were used: anti-CD44-PE, anti-CD24-APC, anti-ESA-PE-Vio770, IgG1-PE, IgG1-APC, IgG1-PE-Vio770 (Miltenyi). Cells were harvested with trypsin-EDTA to produce a single cell suspension. The cells were pelleted by centrifugation, washed twice with PBS and incubated with the antibody for 10 min in the dark at 4°C. The respective isotype control antibodies were used at the same concentrations according to the manufacturer’s instructions. After washing twice with PBS, samples were resuspended in 500 μl PBS and analyzed on a flow cytometer (FACSAriaII, USA). Cells were routinely sorted twice, and the cells were reanalyzed for the purity, which typically was >97%. Data were analyzed with BD FACS Diva software.

Cell proliferation assay
The cells were plated at a density of 2,000 cells/well in 96-well plates. After exposure to the drug for 72 h, WST-8 reagent from Cell Counting kit-8 was added (10 μl/well), incubated for 2 h, and optical absorbance at 450 nm and 630 nm were measured using a microplate reader. OD values of all experimental groups were normalized to control group. Inhibition rate = (1 - absorbance of treated cells/control cells) x 100%.

Sphere formation assay
Cells were plated in six-well ultralow attachment plates (Corning Inc.) at a density of 2,000 cells/ml in serum-free DMEM:F12 (Gibco) supplemented with 20 ng/ml EGF (PeproTech), 100 ng/ml bFGF (PeproTech), 1:50 B27 supplement (Gibco), 100 U/ml penicillin, and 100 μg/ml streptomycin at 37°C in a humidified 5% CO2 atmosphere. After 7 days of incubation, spheres were typically >75 mm. 7-day-old spheres were harvested using 40 mm cell strainers, dissociated to single cells with trypsin-EDTA, and then grown for 7 days. Differentiation was induced by culturing tumorsphere-derived cells in DMEM supplemented with 10% FBS.

Transwell invasion assay
Cells (2.0x10^4 cells/well) in DMEM were seeded into the upper chamber of a Transwell insert (Corning), which was precoated with the Matrigel Matrix (Corning). The lower chamber was filled with DMEM supplemented with 10% of FBS as the chemoattractant. The cells were then cultured for 24 h. The cell number was counted under a Fluorescence Inversion Microscope. Three independent experiments were performed.

Tumorsphere xenografts
Four-week-old nude mice (BALB/c-nu) were maintained in a barrier facility on high efficiency particulate air-filtered racks and fed by breeders from the Department of Laboratory Animal Science, Peking University Health Science Center. Single cells suspensions (1x10^6 in a total volume of 100 ul PBS) were injected subcutaneously into the right and left axilla of nude mice. Panc-1-NC CSCs were injected subcutaneously into left axilla and Panc-1-si CSCs was injected subcutaneously into right axilla.

Eight weeks later, all mice were sacrificed, and tumors were excised. A portion of the tumor tissue was collected, fixed in 4% paraform, and embedded in paraffin for hematoxylin and eosin (H&E) staining to assess tumor pathology. All the animal studies were reviewed and approved by Ethics Committee for Animal Studies at the Peking University, China.

Intravenous injection through vena caudalis
Four-week-old nude mice (BALB/c-nu) were maintained in a barrier facility on high efficiency particulate air-filtered racks and fed by breeders from the Department of Laboratory Animal Science, Peking University Health Science Center. Single cells suspensions (5x10^5 in a total volume of 100 ul of PBS) were injected into the lateral tail vein of nude mice.

Statistical analysis
Data were expressed as mean values and standard deviation (SD), and analyzed using SPSS v.13.0 statistical software (SPSS, Inc.). Student's t-test (for two groups) and one-way analysis of variance (ANOVA) test (for multiple groups) followed by the Student-Newman-Keuls (SNK) test were used to assess the differences. P<0.05 was considered as significant difference.

Results
Panc-1 tumorspheres exhibit stem-like properties
We isolated Panc-1 tumorspheres from human pancreatic cancer cell line Panc-1 using serum-free flating-culture system. We subsequently examined
whether Panc-1 tumorspheres have differentiation potential. ALDH1A1 and Nanog are mature markers of CSCs, and act as the differentiation markers. Real-time RT-PCR was performed to detect the expression of ALDH1A1 and Nanog. Under differentiating conditions, Panc-1 tumorspheres began to adhere and acquired epithelium-like morphology. When returned to adherent culture, Panc-1 tumorspheres redifferentiated and acquired morphology comparable to parental cells (Fig. 1A). We observed that Panc-1 tumorspheres had elevated mRNA levels for genes associated with a stem-like phenotype including ALDH1A1 and Nanog that was lost upon redifferentiation (Fig. 1B). In addition, we found high percentage of Panc-1 tumorspheres CD44+CD24+ESA+ population and the percentage declined after redifferentiation (Fig. 1C and D). These results suggest that Panc-1 tumorspheres have differentiation ability and stem cell-like properties.

Figure 1. Differentiation ability and stem cell-like properties of Panc-1 tumorspheres. (A) The morphology of Panc-1 tumorspheres and Panc-1 re-differentiated tumorsphere. (B) The expression of Nanog and ALDH1A1 was measured by Real-time RT-PCR in Panc-1, Panc-1 tumorsphere and Panc-1 re-differentiated tumorsphere. (C) The percentage of CD44+CD24+ESA+ population in Panc-1, Panc-1 tumorsphere and re-differentiated tumorsphere. (D) Flow cytometry analysis of CD24, CD44, ESA as cancer stem cell markers in Panc-1, Panc-1 tumorsphere and re-differentiated tumorsphere. Data represent mean ± SD. *P < 0.05, ****P < 0.0001.
EMT and activated Hh signaling in Panc-1 tumorspheres

Next we examined the expression of SMO, Gli1, Shh, Snail, E-Cadherin and N-Cadherin in Panc-1 tumorspheres. Real-time RT-PCR showed the high expression of SMO, Gli1 and Shh, and the activated Hh signaling in Panc-1 tumorspheres. In addition, we found that the expression of Snail and N-cadherin was upregulated, while the expression of E-cadherin was down-regulation at mRNA levels in Panc-1 tumorspheres. However, after redifferentiation of Panc-1 tumorspheres SMO, Snail and N-Cadherin were downregulated while E-Cadherin was upregulated (Fig. 2).

Figure 2 The expression of SMO and EMT markers in Panc-1 tumorspheres. Real-time RT-PCR analysis of the expression of SMO, Gli, Shh, Snail, E-Cadherin and N-Cadherin in Panc-1, Panc-1 tumorsphere and Panc-1 redifferentiated tumorsphere. Data represent mean ± SD. * P < 0.05, ** P < 0.01, ***P < 0.001, ****P < 0.0001.

SMO regulates EMT and the expression of cancer stem cell markers in pancreatic cancer cells

Panc-1 cells were transfected with lentiviruses for the delivery of two different siRNAs for SMO (Panc-1-si-1 and Panc-1-si-2) or negative control vectors (Panc-1-NC). Real-time RT-PCR and Western blot analysis confirmed that SMO expression was significantly downregulated in Panc-1-si cells (Fig. 3A-C). Moreover, lentivirus mediated delivery of two different siRNAs for SMO led to reduced expression of Nanog and ALDH1A1 in Panc-1 cells (Fig. 3A and C). In addition, SMO knockdown caused increased expression of E-cadherin and decreased expression of N-cadherin and Snail (Fig. 3A-C). Finally, we detected the expression levels of cancer stem cell markers CD24+CD44+ESA+ in pancreatic cancer cells. The results showed that SMO knockdown resulted in reduced CD24+CD44+ESA+ subfraction in Panc-1-si-1 (Fig. 3D and E). Collectively, these data indicate that Hh signaling could regulate EMT and the expression of cancer stem cell markers in pancreatic cancer cells.

SMO regulates EMT and the expression of cancer stem cell markers in pancreatic CSCs

Panc-1-si cells were plated in six-well ultralow attachment plates in serum-free medium. Five days later, Panc-1-si tumorspheres began to form in culture. After 14 days, we found that Panc-si tumorspheres were less in number and smaller in size than Panc-1-NC tumorspheres (Fig.4A). In addition, SMO knockdown resulted in reduced percentage of CD24+CD44+ESA+ subfraction in Panc-1-si tumorsphere (Fig. 4B and C).

Next we used Fluorescence-activated cell sorting (FACS) to acquire the relatively pure CD24+CD44+ESA+ Panc-1-si CSCs from Panc-1-si tumorspheres. Real-time RT-PCR and Western blot analysis confirmed that SMO expression was significantly downregulated in Panc-1-si CSCs (Fig. 4D-F). As expected, the expression of CSCs markers ALDH1A1 and Nanog was significantly downregulated in Panc-1-si CSCs (Fig. 4D-F). Furthermore, SMO knockdown in the CSCs resulted in the down-regulation of epithelial marker E-cadherin and up-regulation of mesenchymal markers Snail and N-cadherin at mRNA and protein levels (Fig. 4D-F). Taken together, these results suggest that SMO is critical for the acquisition of EMT characteristics of CSCs.

SMO regulates malignant behaviors of pancreatic CSCs in vitro and in vivo

To explore the potential role of Hh signaling in the invasive capacity of pancreatic CSCs, we employed two different siRNAs to knockdown SMO and performed Transwell invasion assay. We found that the number of Panc-1-si CSCs that migrated through the upper chamber of the Costar wells was less than that of Panc-1-NC CSCs (Fig. 5A), indicating that SMO knockdown inhibits the invasion capacity of CSCs.

Next we performed chemosensitivity assay to detect the chemotherapy resistance of Panc-1-si CSCs and Panc-1-si CSCs. Panc-1-NC CSCs were more resistant to gemcitabine than Panc-1-si CSCs (Fig. 5B). IC50 of gemcitabine was 322.33±54.72 µg/ml in Panc-1-NC CSCs, significantly higher than that of Panc-1-si CSCs (P<0.05, Fig. 5C).

Then we implanted subcutaneously equal numbers of Panc-1-NC CSC and Panc-1-si-1 CSC in nude mice and measured tumor growth. Panc-1-NC CSC had 90% (9/10) tumor formation while only 4/10 mice injected with Panc-1-si-1 CSC showed pancreatic tumor engraftment (Fig. 5D).
Figure 3. Changes of EMT and cancer stem cell makers after SMO silencing in Panc-1 cells. (A) Real-time RT-PCR analysis of the expression of SMO, EMT and cancer stem cell makers in Panc-1-NC and Panc-1-si-1. (B) Western blot analysis of the expression of SMO, EMT and cancer stem cell makers in Panc-1-NC and Panc-1-si-1. (C) Real-time RT-PCR analysis of the expression of SMO, EMT and cancer stem cell makers in Panc-1-NC and Panc-1-si-2. (D) The percentage of CD44+CD24+ESA+ population in Panc-1-NC and Panc-1-si-1. (E) Flow cytometry analysis of CD24, CD44, ESA as cancer stem cell markers in Panc-1-NC and Panc-1-si-1. Data represent mean ± SD. * P < 0.05, ** P < 0.01, ****P < 0.0001.
Figure 4. Changes of EMT and cancer stem cell makers after SMO silencing in Panc-1 CSCs. (A) The morphology of Panc-1-NC and Panc-1-si-1 tumorspheres. (B) The percentage of CD44+CD24+ESA+ population in Panc-1-NC and Panc-1-si-1 tumorspheres. (C) Flow cytometry analysis of CD24, CD44, ESA in Panc-1-NC and Panc-1-si-1 tumorspheres. (D) RT-PCR analysis of SMO, EMT and cancer stem cell makers in Panc-1-NC CSCs and Panc-1-si-1 CSCs. (E) Western blot analysis of SMO, EMT and cancer stem cell makers in Panc-1-NC CSCs and Panc-1-si-1 CSCs. (F) RT-PCR analysis of SMO, EMT and cancer stem cell markers in Panc-1-NC CSCs and Panc-1-si-2 CSCs. Data represent mean ± SD. * P < 0.05, ** P < 0.01, ***P < 0.001.
Figure 5. SMO regulates malignant behaviors of Panc-1 CSCs. (A) Effect of SMO knockdown on the invasion of Panc-1-NC CSC and Panc-1-si CSC. (B) Inhibition curve of Panc-1-NC CSC and Panc-1-si CSC at various dose of gemcitabine (GEM). (C) IC50 of GEM in Panc-1-NC CSC and Panc-1-si CSC. (D) Tumorigenicity assay using Panc-1-NC CSC and Panc-1-si CSC. (E) The pulmonary metastasis capability of Panc-1-si CSC compared with Panc-1-NC CSC. (F) The pulmonary morphology of Panc-1-NC CSCs and Panc-1-si CSCs after intravenous injection through vena caudalis. (G) Lung metastatic lesion of Panc-1-NC CSCs and Panc-1-si CSCs. Data represent mean ± SD. ***P < 0.001, ****P < 0.0001.
Panc-1-NC and Panc-1-si were transfected with recombinant lentiviral vectors and we used Fluorescence-activated cell sorting (FACS) to acquire the relatively pure CD24+CD44+ESA+ Panc-1-si-luci CSCs from Panc-1-si-luci tumorspheres. Single cell suspensions of Panc-1-NC-luci CSCs or Panc-1-si-luci CSCs were injected into the lateral tail vein of nude mice. We observed that Panc-1-si-luci CSCs showed reduced capability of pulmonary metastasis compared with Panc-1-NC-luci CSCs (Fig. 5E). The nude mice were killed following the lung-taken operation. Compared with control group, pulmonary metastasis was less in the test group with Panc-si-luci CSC intravenous injection through vena caudalis (Fig. 5F). Photomicrographs showed that tumor cells were found in 5/5 mice injected with Panc-1-NC CSCs and 0/5 mice injected with Panc-si-1 CSCs (Fig. 5G).

Discussion

Accumulating data indicate the existence of tumor-initiating or cancer stem cells in human pancreatic tumor that possess the capacity to self-renew and differentiate into bulk tumor cells. However, the molecular mechanisms that drive the enhanced tumorigenic potential and stem-like behavior of tumor-initiating cells are poorly understood. In this study, we provide evidence that SMO plays a vital and previously unrecognized role in the maintenance of stem-like cells isolated from human pancreatic cancer cells.

Currently, CSCs from pancreatic cancer are identified based on cell surface antigens such as CD44, CD24, epithelial-specific antigen (ESA) and CD133 [11]. There are still controversies to use which types of markers to enrich pancreatic CSCs. However, cell populations enriched for high ALDH1A1 activity were shown to be sufficient for efficient tumor-initiation with enhanced tumorigenic potential [12]. Therefore, in the present study we chose CD24CD44ESA as the surface markers of pancreatic CSCs, and the expression of ALDH1A1 and Nanog as the markers of cancer stem cell-like properties of pancreatic CSCs.

Serum-free floating culture system is an ordinary technique used to isolate CSCs from several cancer types such as brain, breast and colon cancers [13-15]. Panc-1 is a common pancreatic adenocarcinoma cell line used for the study of CSCs [16-18]. In our study, compared to control Panc-1, SMO, Snail, N-Cadherin, ALDH1A1 and Nanog were highly expressed in Panc-1 tumorsphere with decreased expression of E-cadherin and increased CD24+CD44ESA+ cell population. After the induction of differentiation, the expression levels of SMO, Snail, N-Cadherin, E-cadherin and the CD24+CD44ESA+ cell population were similar to Panc-1. Taken together, Panc-1 tumorspheres display the potentials of self-renewal and differentiation, which are the characteristics of CSCs. Morel et al. claimed that EMT process could stimulate the stem-like capability of human breast cancer cells [19]. In this study, compared with the highly differentiated CD44-/CD24+ cells, the stem cell-like CD44+/CD24- cells had high expression of EMT-inducing transcription factor Snail and mesenchyme cell marker [20]. These data suggest that Hh signaling and EMT process are active in Panc-1 tumorspheres.

Hh signaling has been associated with the self-renewal process of CSCs [21]. Previous study demonstrated that Hh signaling receptor SMO was highly expressed in normal human mammary stem/progenitor cells and implicated in malignant human mammary stem cells [22]. In our study, SMO knockdown resulted in the down-regulation of ALDH1A1 and Nanog and the reduction of CD24+CD44+ESA+ cell population, and reduced the size of the tumorsphere. Therefore, SMO plays a vital role in the regulation of pancreatic cancer stem cell properties and the maintenance of the self-renewal of pancreatic CSCs.

EMT is a vital process in which malignant tumor cells stem from the epithelium to obtain the capability of migration and invasion, including pancreatic tumor [23]. In this study we investigated the role of Hh signaling in EMT of pancreatic CSCs. We found that SMO knockdown resulted in declined invasion capability of Panc-1 CSCs in vitro. SMO knockdown also caused the decreases in the incidence of subcutaneous tumorigenesis and pulmonary metastasis in vivo. These data demonstrate that SMO could regulate EMT process in human pancreatic CSCs and subsequently impact the capability of invasion and migration of pancreatic CSCs.

Furthermore, EMT is implicated in the drug resistance of pancreatic tumor. Li et al. found that gemcitabine-sensitive PC cells showed strong expression of epithelial marker E-cadherin while gemcitabine-resistant PC cells exhibited strong expression of mesenchymal makers at mRNA and protein levels [24]. In addition, chemoradiation-resistant pancreatic cancer cells exhibited the properties similar to CSCs and also underwent EMT, suggesting that chemoradiation resistance-induced EMT is associated to the generation of pancreatic CSCs [25]. Thus, we wondered whether SMO could regulate the chemoresistance of pancreatic CSCs. We found that Panc-1-NC CSCs were more resistant to gemcitabine compared to Panc-1-si CSCs. These results indicate that pancreatic CSCs demonstrate declined capacity of chemoresistance with a decreased expression of SMO.

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We proposed that Hh signaling is implicated in the chemoresistance of pancreatic cancer by regulating EMT process of CSCs. Further studies in stem-like tumorspheres derived from other pancreatic cancer cell lines such as MiaPaCa-2 and L3.6pl are needed to confirm our conclusion.

In conclusion, we demonstrate that tumorspheres derived from pancreatic cancer cell line Panc-1 possess self-renewal, differentiation and stemness properties. Panc-1 tumorspheres underwent EMT and exhibited activated Hh signaling. The inhibition of Hh signaling by SMO knockdown represents an effective method to inhibit the self-renewal, EMT, invasion, chemoresistance, tumorigenesis and pulmonary metastasis of pancreatic CSCs. These data suggest that targeted therapy against Hh signaling may be beneficial in the treatment of pancreatic cancer.

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Competing Interests

The authors have declared that no competing interest exists.

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