Role of Hedgehog signaling pathway in proliferation and invasiveness of hepatocellular carcinoma cells

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Abstract. The Hedgehog (Hh) pathway has been confirmed a contributor to the carcinogenesis and progression of various tumor types. To investigate the Hh signaling activity in human hepatocellular carcinoma (HCC), we detected the expression levels of Hh pathway components (Shh, Ptch1 and Gli2) in 57 samples of HCC and corresponding adjacent-tumor liver tissues. The Hh pathway was overexpressed in cancer tissues compared with non-cancer tissues and correlated closely with histologic differentiation and portal venous invasion of HCC. To elucidate the relationship between Hh signaling and HCC progression, we performed a further study in vitro. First, the expression levels of the signaling were detected in a subset of hepatoma cell lines and SMMC-7721 cells selected with high level of Hh signaling expression. Next, we employed KAAD-cyclopamine (a specific inhibitor of Hedgehog pathway) to block the Hh pathway in SMMC-7721 cells and assessed the changes of their biological behaviors. The results showed that the blockade of Hh signaling pathway by KAAD-cyclopamine induced reduction of DNA synthesis leading to marked cell growth inhibition and also caused significant attenuation in invasiveness and motility of HCC cells. Collectively, our data demonstrated that the Hh pathway plays an important role in HCC development and invasion. Blockade of the Hh signaling pathway may be a potential target of new therapeutic strategy for HCC.

Introduction

Hepatocellular carcinoma (HCC) is the fifth most malignant neoplasm in the world, with approximately 1 million new cases per year worldwide and causes approximately 100,000 deaths annually in China (1,2). The etiology of HCC includes chronic viral hepatitis, alcohol abuse, environmental carcinogens or genetic disorders; hepatitis B virus (HBV) infection is the major cause of HCC in China. Although several risk factors for HCC development are known, therapeutic options for the disease are very limited. The hepatic resection remains the most effective treatment (3), but the prognosis of HCC is generally poor, with high post-operative recurrence and invasiveness of primary tumor responses (4). Decades of intense research efforts have aimed at better understanding the underlying etiologic and pathophysiological mechanisms, but its molecular mechanisms remain unclear. Therefore, it is vital to explore new molecular markers for treatment strategies have the potential to significantly improve the prognosis of this dismal disease.

The hedgehog (Hh) signaling pathway is a highly conserved system, which plays a crucial role in tissue patterning, cell differentiation and proliferation (5). In vertebrate organisms, the signaling pathway is initiated by the binding of ligands (sonic, Indian, and desert hedgehog) to the membranous receptor patched (Ptch) which in turn alleviates the suppression on smoothened (Smo), a 7-transmembranous domains protein receptor, subsequently Smo triggers a series of intra-cellular events with resultant activation of the zinc-finger transcription effectors, glioma-associated oncogenes (Gli1, Gli2, Gli3) transcription factor, which induces the expression of numerous target genes that regulate proliferation, differentiation, and extracellular matrix interactions (5-7). Apart from epidermis, myoblasts, and neural stem cells and during tissue repair, Hh signaling pathway remains inactive in most normal mature tissues including adult liver, despite the pathway being critical in the early development of the liver and contributing to differentiation between hepatic and pancreatic tissue formation (8-12).

Aberrant activation of the Hh signaling pathway, especially ectopic expression of Gli2 is responsible for the onset and progression of tumorigenesis, in cancers such as of the pancreas, colon, stomach, lung, prostate, breast and skin (13-21). Downregulation of Gli2 or inhibition of the pathway with a ligand-blocking antibody or KAAD-cyclopamine, the selective Smo antagonist, has been shown to retard the growth of tumors and to decrease tumor size. However, the role of the...
pathway in pathogenesis of HCC is still indistinct in spite of a few previous studies (22-24), particularly concerning invasiveness there is lack of information. In the current study, expression of the Hh signaling pathway in HCC was detected and the relationship with regard to clinicopathologic characteristics was investigated. In addition, the activation of Hh signaling in a subset of liver cell lines was evaluated. Finally, the response of HCC cells to inhibition of the Hh signaling by KAAD-cyclopamine was assessed by evaluating cell proliferation, motility and invasiveness.

Patients and methods

Patients and specimens. A total of 57 HCC patients who had undergone liver resection without preoperative treatment at Tongji Hospital of Tongji Medical College of Huazhong University of Science and Technology between June 2004 and December 2007 were examined. All tumor specimens were pathologically diagnosed as HCC. The age of patients ranged from 24 to 70 years, with an average age of 46.4 years, and there were 43 males and 14 females. The number of patients positive for HBV and HCV was 47 and 3 respectively. There were 27 well-, 16 moderately- and 14 poorly-differentiated HCC. Portal vein tumor thrombus was observed in 26 patients.

Immunohistochemistry. Immunohistochemical stains were performed on formalin-fixed and paraffin-embedded tissue sections (4 μm). The sections were prepared according to classical methods and treated with blocking solution before being sequentially incubated with primary Ab against Shh (1:100, Santa Cruz, USA), Ptc1 (1:100, Santa Cruz, USA), Gli2 (1:200, Abcam, UK) overnight at 4˚C, followed by biotinylated Ab and HRP-conjugated streptavidin (each for 30 min at room temperature). Detection was achieved with dianinobenzidine and hematoxylin counter staining. No significant staining was observed in the negative controls, which were treated with the same species normal IgG in place of primary Ab. The expression was evaluated independently by two pathologists; at least twice for each tissue sample which were treated with the same species normal IgG in place of primary Ab. The expression was evaluated independently by two pathologists; at least twice for each tissue sample with similar results.

Cell culture. Cell lines Hep3B and Huh-7 were obtained from the Chinese Type Culture Collection (Wuhan), L02, HepG2, and SMMC-7721 were generously provided by Drs Liming Xia and Wei Yan. All cell lines were grown in Dulbecco's modified essential medium (DMEM, Thermo, USA) supplemented with 10% fetal bovine serum (FBS, Invitrogen, USA). The cells were grown at 37˚C, 5% CO2 in fully humidified air. Cells were treated with KAAD-cyclopamine (Toronto Research Chemicals, Canada), the special antagonist of smoothened (Smo), control medium containing tomatidine (Sigma, USA), the structurally similar compound with non-specific inhibition on hedgehog signaling, or DMSO alone.

RNA extraction and RT-PCR. Total RNA was extracted from all cell lines and typical tissue samples using TRIzol reagent (Invitrogen), and lysates were treated with DNA-free DNase treatment and removal reagents (Ambion, USA). The house-keeping gene GAPDH was used as a loading control. RT-PCR was performed using ThermoScript RT-PCR system (Invitrogen). In brief, 1 μg of total RNA was used in a 20 μl reverse transcription assay. Subsequently, 2 μl of the reverse transcription product was used in a 25 μl PCR reaction. All PCR reactions were performed at 5-55˚C annealing temperature for 25-32 cycles. All primers used in this study are listed in Table I.

MTT assay. The antiproliferative effects of KAAD-cyclopamine with different concentration on hepatoma cells were detected by the methyl thiazolyl tetrazolium (MTT) assay. The cells were cultured in 96-well plates at 2x10^4 cells per well and treated with 10% FBS medium containing KAAD-cyclopamine at various concentrations (1, 5, 10 μM) for 12, 24, 48, 60, 72 h. All experiments were repeated three times. 20 ml MTT (10 g/l) was added to each well, medium was discarded after 4 h, followed by the addition of 150 ml DMSO. The 490 nm absorbance was measured by a Bio-Tek microplate reader, the inhibitory ratios (IR) were calculated as follows: IR (%) = [(1-absorbance of the treated wells)/absorbance of the control wells] x 100.

BrdU incorporation assay. To measure DNA synthesis rates, following treatments for 72 h, the cells were chased by 10 μmol/l 5-bromodeoxyuridine (BrdU), for 2 h. BrdU was detected with mouse anti-BrdU (1:200, Santa Cruz, USA) for 2 h followed by fluorescein isothiocyanate-conjugated antibody incubated for 45 min, protected from direct light. Total DNA was stained with PI. BrdU incorporation into DNA and DNA content in nuclei were detected and quantified on a FACSscan (BD, USA). Shown here are the representative data seen in three independent experiments. Note that: % cells in active DNA synthesis is defined as BrdU^+ cells/total cells (the events in R2 area/total events) and the average BrdU incorporation into DNA (mean BrdU signal intensities in R2 events) is shown.

Wound healing assay. Confluent cell monolayers in 24-well plates were scrape-wounded using a micropipette tip. After being washed, cells were cultured in media with 0.5% FCS. Migration rate (average velocity) and degree of wound closure were assessed by measuring the distance between wound edges at time intervals of 3 h, representative wells were observed by inverted microscope. The cell movement velocity was determined by tracing single cells at different time-points, using MetaMorph software (Metamorph Imaging System, USA). All samples were tested in triplicate and were scored in a double-blinded manner by three investigators.

Time-lapse video. For time-lapse observation of cell movement, the cells plated in complete medium were monitored by inverted microscope with a charge coupled-device camera at planned intervals. Phase-contrast images were captured and analyzed with Metamorph software (Metamorph Imaging System). The motility of individual cells was evaluated by tracking their movement from images recorded every 10 min. The average speed (μm/h) of locomotion was calculated as the total track length divided by the number of hours recorded. For each experimental condition, 30-50 cells were analyzed.
Modified Boyden chamber invasion assays. The effect of KAAD-cyclopamine on hepatocellular carcinoma was determined using modified Boyden chamber assays as previously described (25). Briefly, 24-well transwell units with polycarbonate membrane filters (6.5 mm diameter, 8 μm pore size, Costar, Corning, USA) were coated with Matrigel (100 μg/cm²/100 μl PBS). Cells (1x10⁵/200 μl DMEM with 0.5% FBS) were added to the upper compartment and incubated for 48 h in the absence or presence of KAAD-cyclopamine (10 μmol/l) at 37˚C. The lower chambers were filled with 2 ml DMEM supplemented with 5% FBS as chemoattractant. After removal of nonmigratory cells from the upper surface of the filter, invasive cells that had passed through to the lower surface of the filter were fixed and stained with crystal violet. Invasive cells were scored by counting at least five fields per filter. Counting accuracy was controlled by optical density OD570 quantification of methanol-solubilized dye.

Statistical analyses. Associations between gene expression and clinicopathologic variables were analyzed by Fisher’s exact test. The variables such as the data using cell lines were compared by Mann-Whitney’s U test, One- or Two-way ANOVA according to the experimental design applied. All statistical analyses were performed using the SPSS 11.0 software program (SPSS, USA). A two-tailed P<0.05 was considered statistically significant.

Results

Overexpression of Hh pathway in HCC comparing with its corresponding adjacent-tumor liver tissue. We detected Shh, Pch1 and Gli2 expressions in 57 cases of HCC and the corresponding tumor-adjacent liver tissues for the purpose of evaluating the hedgehog signaling pathway activation. By immunohistochemical staining, Shh and Pch1 expressions were defined as cytoplasmic staining with either stippled or granular pattern, and Gli2 located in cytoplasm and/or nucleus, with predominance in the cytoplasm. The overexpression of the three molecules in carcinoma presented a striking contrast to non-tumor liver tissues.

Relationship between immunohistochemical results and clinicopathologic features. Table II shows the expression distinction of the Hh components in HCC varied clinicopathologic factor groups. Overexpression of three components, Shh, Pch1 and Gli2, was significantly associated with portal venous invasion (P<0.05); meanwhile, Shh and Gli2 were notably correlated to HCC histologic differentiation (P<0.05). But the results did not demonstrate any association between the Hh pathway and gender, tumor size and virus infection.

Different expressions of Hh pathway components in HCC cell lines. Five cell lines including L02, HepG2, Hep3B, SMMC-7721 and Huh7 as well as representative HCC tissue and adjacent-tumor tissue were investigated for expression
levels of the Hh pathway component gene transcription by RT-PCR. All of the components were expressed in HCC cell lines to different extent (Fig. 2). Expression of Shh, Ptch, Smo and Gli2 mRNAs was robustly observed in hepatoma cell lines, with predominance in SMMC-7721, but weaker expression in normal liver cell lines (L02) was noted, especially Gli2 was significantly overexpressed in SMMC-7721 but almost undetectable in L02. The enhanced expression of Gli1 was only observed in Hep3B and SMMC-7721. However, Ihh and Gli3 were expressed at similarly low levels in all cell lines. In addition, the different expressions of the Hh pathway components between HCC tissue and adjacent-tumor liver tissue were further confirmed. Among the 5 cell lines, Hh pathway was significantly activated in SMMC-7721, which is characterized by poor differentiation and better invasiveness. Therefore, we used it for the subsequent experiments.

Blockade of the Hh pathway reduced SMMC-7721 cell proliferation. In order to evaluate the effects of Hh pathway on cell growth, we detected proliferation of SMMC-7721 cells at interval of 12 h for up to 72 h, which was incubated with KAAD-cyclopamine, a specific antagonist of Hh signaling. The antagonist reduced the cell proliferation in a dose- and time-dependent pattern (Fig. 3A). For each concentration, IR increased gradually over time (P<0.05); at

<table>
<thead>
<tr>
<th>HCC subgroup</th>
<th>Shh pos</th>
<th>P-value</th>
<th>Ptch1 pos</th>
<th>P-value</th>
<th>Gli2 pos</th>
<th>P-value</th>
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<tr>
<td>Hepatocellular carcinoma</td>
<td>36 (63.2)</td>
<td>0.039*</td>
<td>30 (52.6)</td>
<td>0.013*</td>
<td>40 (70.2)</td>
<td>0.002*</td>
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<td>Adjacent-tumor liver tissues</td>
<td>25 (43.9)</td>
<td>0.039*</td>
<td>16 (28.1)</td>
<td>0.013*</td>
<td>24 (42.1)</td>
<td>0.002*</td>
</tr>
<tr>
<td>Sex</td>
<td></td>
<td></td>
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<tr>
<td>Male</td>
<td>26 (60.5)</td>
<td>0.46</td>
<td>24 (55.8)</td>
<td>0.399</td>
<td>30 (69.8)</td>
<td>0.906</td>
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<tr>
<td>Female</td>
<td>10 (71.4)</td>
<td>0.46</td>
<td>6 (42.9)</td>
<td>0.399</td>
<td>10 (71.4)</td>
<td>0.906</td>
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<td>Tumor size (cm)</td>
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<td>&gt;3</td>
<td>30 (66.7)</td>
<td>0.288</td>
<td>26 (57.8)</td>
<td>0.132</td>
<td>33 (73.3)</td>
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<td>4 (33.3)</td>
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<td>7 (58.3)</td>
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<td>0.039*</td>
<td>25 (53.2)</td>
<td>0.013*</td>
<td>34 (72.3)</td>
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<td>HCV*</td>
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<td>0.039*</td>
<td>1 (33.3)</td>
<td>0.013*</td>
<td>2 (66.7)</td>
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<td>HBV- HCV*</td>
<td>5 (71.4)</td>
<td>0.876</td>
<td>3 (42.9)</td>
<td>0.774</td>
<td>4 (57.1)</td>
<td>0.138</td>
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<td>Well</td>
<td>13 (48.1)</td>
<td>0.019*</td>
<td>12 (44.4)</td>
<td>0.252</td>
<td>15 (55.6)</td>
<td>0.002*</td>
</tr>
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<td>Moderately</td>
<td>10 (62.5)</td>
<td>0.019*</td>
<td>8 (50)</td>
<td>0.252</td>
<td>11 (68.8)</td>
<td>0.002*</td>
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<tr>
<td>Poorly</td>
<td>13 (92.9)</td>
<td>0.019*</td>
<td>10 (71.4)</td>
<td>0.252</td>
<td>14 (100)</td>
<td>0.002*</td>
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<td>Positive</td>
<td>20 (76.9)</td>
<td>0.049*</td>
<td>18 (69.2)</td>
<td>0.022*</td>
<td>23 (88.5)</td>
<td>0.006*</td>
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<td>Negative</td>
<td>16 (51.6)</td>
<td>0.049*</td>
<td>12 (38.7)</td>
<td>0.022*</td>
<td>17 (54.8)</td>
<td>0.006*</td>
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</tbody>
</table>

Abbreviations: pos, positive; PVTT, portal vein tumor thrombus. *P<0.05; **P<0.01.
each time-point, the IR rose with the increase of treatment concentration, except on the first day when no concentration treatment appeared to have a distinct inhibitory effect (P>0.05). The maximum IR by 1, 5 and 10 μmol/l KAAD-cyclopamine treatment was 10.9±2.35%, 30.24±4.21%, 67.34±4.18% at 72 h, respectively. However, cell proliferation was weakly affected by the control homologue tomatidine compared with KAAD-cyclopamine (P>0.05) (Fig. 3B). For validation of the role of Hh pathway in cell proliferation, we measured the DNA synthesis by more sensitive BrdU labeling assay further (Fig. 4). The result showed that KAAD-cyclopamine induced reduction of cell number in active DNA synthesis, the proportion of R2 (BrdU positive cells/total cells) was 27.41±2.2% in treated cells, remarkably less than 40.7±3.1% in controls (P<0.01). These findings implied that inhibition of Hh pathway can significantly decrease proliferation of SMMC-7721.

**KAAD-cyclopamine induced-inhibition in invasiveness of SMMC-7721 cells.** To investigate the putative role of Hh pathway in the invasive potential of HCC, we performed wound assays and invasion assays. As shown in Fig. 4, the motility and invasive capacity of SMMC-7721 cells were greatly suppressed by KAAD-cyclopamine. Wound assay revealed significant reductions in wound closure by >65% for SMMC-7721 cells with blockade of the Hh pathway compared with those of control cells (Fig. 4A and B). Moreover, mean velocities of treated cells were significantly lower (~40%) than those of control cells (24.9±3.8 vs. 41.3±7 μm/h; P<0.01; Fig. 4C). Invasion assay displayed that SMMC-7721 exhibited the strongest invasive potential (71.1±11.5 cells/hpf) but to a significantly lesser extent after application of KAAD-cyclopamine (36.1±6.4 cells/hpf; P<0.01; Fig. 4D and E). Collectively, these studies demonstrate that the Hh pathway could play some facilitatory role in invasion of HCC.

**Discussion**

HCC is a common malignant disease with inadequate therapeutic methods, studies of the underlying molecular mechanism provide potential to identify new therapeutic target. Previous evidence has demonstrated that the Hh signaling plays an important role in multiple tumor types, for example, pancreatic cancer, colon carcinoma, gastric cancer, small cell lung cancer, prostate cancer, breast cancer and basal cell carcinoma (13-20). Recent studies revealed that the Hh pathway is abnormally activated in human HCC (22-24).
However, to date no report has been presented to give a detailed description on the functional relevance of the Hh pathway to HCC progression.

In the present study, we found overexpression of Hh signaling pathway, represented by Shh, Ptc1 and Gli2, in HCC compared with corresponding adjacent-tumor liver tissues, which was in line with previous results in other cancers (16,19). Moreover, the ectopic expression of Shh and Gli2 was associated with histologic differentiation of HCC, which was inconsistent with the results of Huang et al (23), who did not detect Gli2 expression and divided the differentiation degree into two groups (well-differentiated, and moderately
to poorly-differentiated). Noticeably, we observed for the first time that Hh pathway was associated significantly with portal venous invasion of HCC. Poor differentiation and intrahepatic tumor spread through the portal vein system are the crucial histologic features of HCC aggressiveness (26). In addition, we detected the expression of Hh signaling in hepatoma cell lines and normal liver cell lines, and found a higher expression level in hepatoma cell lines especially in SMMC-7721 cells, which were regarded as poorly differentiated. These results indicated that continuous activation of the Hh signaling pathway is associated with histologic differentiation and invasiveness and may promote the progression of HCC.

Although it has been confirmed that the Hh pathway participates in cell migration and angiogenesis processes and is correlated with the severity of the associated tumor (27-29), additionally, disruption of Hh pathway has been shown to inhibit metastasis in pancreatic cancer and gastric cancer (29-31), the relationship between the Hh signaling pathway and aggressiveness of HCC cells remains unclear. It is necessary to investigate whether the signaling influences proliferative and invasive potential of HCC cells. It has been reported that cyclopamine is a specific antagonist of the Hh pathway by binding to Smo and induces inactive Hh signaling, KAAD-cyclopamine is a synthetic cyclopamine-derivative (32). We employed KAAD-cyclopamine to block the Hh signaling pathway in SMMC-7721 cells and to determine the effect of the pathway on proliferation and invasiveness of HCC cells. The outcomes demonstrated that KAAD-cyclopamine reduced the DNA synthesis leading to marked cell growth inhibition, moreover, the antagonist inhibited the proliferation in a dose- and time-dependent manner, the maximum inhibitory ratio close to 70%. In addition to attenuation the proliferative program, the abrogation of the Hh signaling with KAAD-cyclopamine resulted in reduction in motility and invasiveness of SMMC-7721 cells, the capacity of migration and invasion decreased by ~40 and 50% respectively.

It is notable that the Gli2 oncogene RNA expression level was abnormally high in HCC cell lines, significantly in SMMC-7721 cells, but it was almost undetectable in L02 cells, the normal liver cell lines. This different expression of Gli2 agrees with its different immunoreactivity between cancerous and noncancerous samples from one HCC patients. Gli2 was overexpressed in HCC with portal venous invasion or poorly differentiated samples showed positive correlation between Gli2 expression and malignant phenotype of SMMC-7721 cells, the capacity of migration and invasion decreased by ~40 and 50% respectively.

Blocking the Hh signaling pathway remarkably decreased HCC cell growth and motility, indicating that it could provide a new therapeutic option for HCC to improve prognosis.

References


