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## Sonic Hedgehog signaling pathway in primary liver cancer cells

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### ABSTRACT

**Objective:** To investigate clinical significance of Sonic Hedgehog (SHH) signaling pathway molecular Shh, Smo and Gli2 in primary hepatocellular carcinoma (HCC) tissue. **Methods:** A total of 30 HCC tissue samples were collected. Protein expression of SHH signaling pathway molecules Shh, Smo and Gli2 in HCC tissues and para – carcinoma tissue were detected by using immunohistochemical method. Cirrhosis and normal liver tissue specimens were observed as control to analyze the expression of SHH signaling pathway molecular Shh, Smo and Gli2 mRNA in HCC tissues and corresponding para–carcinoma tissues and its relationship with the onset of HCC. **Results:** There was no expression of Shh, Smo and Gli2 protein in normal liver tissue, while their positive rates were 63.3%, 76.7% and 66.7% in HCC tissues, respectively, with a significantly higher expression level than that in the para – carcinoma tissue ( $P < 0.05$ ). The protein expressions in HepG2 cells were slightly lower than that in Huh7 cells, with no statistical difference ( $P > 0.05$ ); Shh and Smo protein was detected in part of cirrhosis with positive expression, but Gli2 protein was not observable in cirrhosis tissues. **Conclusions:** In HCC tissues, the high expression level of SHH signaling pathway molecules signal peptide (Shh), membrane protein receptor (Smo) and nuclear transcription molecular (Gli2) can be indicators of the onset of liver cancer.

## 1. Introduction

The primary hepatocellular carcinoma (HCC) is a common malignant tumor of the digestive system, its incidence increases year by year in our country, seriously harm human health due to its complicated treatment and poor prognosis<sup>[1–3]</sup>. The pathogenesis of HCC is unclear<sup>[4]</sup>. Studies have reported<sup>[5–7]</sup>, activation of Sonic Hedgehog (SHH) pathway exists in the HCC tissue and may promote tumor growth and angiogenesis, indicating that SHH signaling pathway molecules are involved in the onset of HCC process. SHH signaling pathway is a key regulator pathway for cell proliferation in the process of embryonic development. SHH pathway inactivation are observed in both embryonic development and tumorigenesis process, which may have

been caused by the disorder expression of cancer gene<sup>[8]</sup>. Currently, there are many researches about the relationship of SHH signaling pathway and malignant tumor, but the correlation between its expression and HCC is rare reported. To observe the mechanism of SHH signaling pathway in the development of liver cancer, we detected the protein expression of molecules Shh, Smo and Gli2 in 30 HCC tissue samples using immunohistochemical method, and analyzed the relationship between the liver cancer and the expression disorder of SHH signaling pathway, as to provide experimental basis for effective prevention and treatment of liver cancer.

## 2. Materials and methods

### 2.1. Specimens collection

A total of 30 (male 22, female 8) HCC patients aged between 37–72 yr [mean (51.7±2.3) yr] who were admitted from 2011.1 to 2012.1 were recruited, the tissue specimens were

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collected after surgery. Edmondson level: I – II 17 cases, III – IV 13 cases; tumor diameter: 24 cases  $\geq 3$  cm, 6 cases  $\leq 3$  cm, 17 cases of cirrhosis, 12 cases with intrahepatic portal vein invasion; 22 cases with serum AFP  $> 25$  ng/mL. Tissue specimens (5 cm plus) of liver tumor and para-tumor were intraoperative extracted immediately from the 30 samples, then were frozen in liquid nitrogen, followed by SHH pathway molecule detection using immunohistochemical method; respective 10 specimens of cirrhosis and normal liver tissue were set for control observation.

## 2.2. Instruments and reagents

Immunohistochemical kit, DAB chromogenic agent, PBS (Golden Bridge Biotechnology co., LTD., Beijing Chinese); 1640 (powder), trypsin (GIBCO Co., USA); Rabbit polyclonal antibody against human Shh, rabbit polyclonal antibody against human Smo, rabbit polyclonal antibody against human Gli2, biotin labeled goat rabbit IgG, blocked with goat serum of 5% BSA, HRP conjugated Streptavidin were bought from Wuhan Science and Technology Co., LTD, China&United States. CO<sub>2</sub> incubator Heareus Company (Germany); BH-2 optical microscope (Olympus, Germany); Inverted microscope (OLYMPUS, Japan); Gel scan imaging system (BioRad Co., US); Image analysis system Imagepro-plus morphometric analysis software.

## 2.3. Experimental methods

Immunohistochemical SP method was used to detect SHH signaling pathway molecules Shh, Smo and Gli2 protein expression. After fixation and paraffin embedding, samples were sliced and moved in baker at constant 60 °C for 30 min. After conventional dewaxing, 0.3% of Triton X-100 was added, then incubated at room temperature for 10 min, in order to dissolve the cell membrane, nuclear membrane, so that the antibodies could get into cells. 3% hydrogen peroxide was added to inactivate endogenous enzymes, washed three times with PBS. 50  $\mu$  L of primary antibodies (Shh, Smo: 1:150 dilution, Gli2: 1:100 dilution) was added at room temperature for 30 min. After PBS wash, second antibody was added to rabbit IgG, then was incubated for 30 min at room temperature. HRP conjugated Streptavidin was added then incubated for 30 min at room temperature. They were washed with PBS and DAB for coloration, redyed with hematoxylin followed by dehydration, transparency and cementing. RT-PCR reaction was conducted strictly

according to the kit instructions

## 2.4. Results determine

Five vision fields were randomly taken under the light microscopy (SP $\times$ 400), positive cells appeared blond, brown granules microscopically in cytoplasm, result determination was according to the proportion of positive cells to staining intensity<sup>[9]</sup>. Positive cell percentage of scoring criteria: 0: positive cells  $\leq 5\%$ ; 1 point: positive cells  $> 5\%$ – $25\%$ ; 2 points: positive cells  $> 25\%$ – $50\%$ ; 3 points: positive cells  $> 50\%$ – $75\%$ ; 4 points: positive cells  $> 75\%$ . Dyeing strength criteria: 0: without coloring; 1 point: light yellow; 2 points: tan; three points: brown. Product of the integral above was marked as the dyeing criteria: the negative (-): 0; weakly positive (+): 1–4 points, positive (++ – +++) :  $> 4$  points.

## 2.5. Statistics analysis

SPSS19.0 statistical software was used, all data were expressed as mean $\pm$ sd. Variance was analyzed with ANOVA and F test.  $P < 0.05$  was regarded as statistically significant difference.

## 3. Results

### 3.1. mRNA expression of SHH pathway in HCC tissues and HCC cell lines

mRNA positive expression rate of Shh, Smo and Gli2 in HCC tissue was 60%, 80%, and 70%, similar to the immunohistochemical detection results; and was 20%, 30% and 30% in para-carcinoma tissue respectively, mRNA expression significantly increased in HCC tissue compared with para-carcinoma tissue ( $P < 0.05$ ). mRNA expression of Shh, Smo and Gli2 in liver cancer cell line HepG2 was slightly lower than that in Huh7 ( $P > 0.05$ ), as shown in Table 1.

**Table 1**

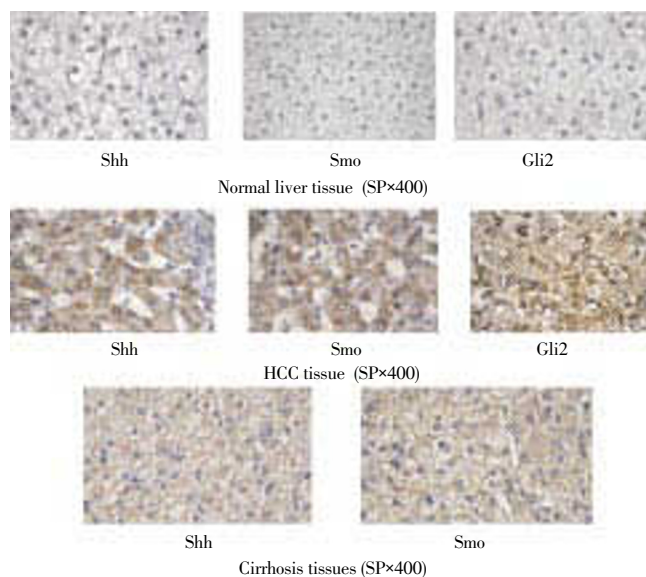
Expression of SHH pathway in HCC tissues and HCC cell lines.

Group	Shh mRNA	Smo mRNA	Gli2 mRNA
HCC tissues	0.67 $\pm$ 0.02*	0.73 $\pm$ 0.01*	0.29 $\pm$ 0.01*
para-carcinoma tissue	0.54 $\pm$ 0.02	0.64 $\pm$ 0.01	0.18 $\pm$ 0.02
HepG2 cell lines	0.55 $\pm$ 0.04	0.69 $\pm$ 0.01	0.22 $\pm$ 0.02
Huh7 cell lines	0.61 $\pm$ 0.03	0.71 $\pm$ 0.01	0.25 $\pm$ 0.02

note: comparison with para-carcinoma tissue \* $P < 0.05$ .

### 3.2. Protein expression of SHH signaling pathway molecules in different tissues

Shh, Smo and Gli2 protein had no expressions in normal liver tissue; Shh and Smo could be detected in part of cirrhosis tissue; Protein expression of Shh, Smo and Gli2 in HCC tissue was 63.3% (19/30), 76.7% (23/30), and 66.7% (20/30), Shh and Smo protein mainly expressed in cytoplasm, with granules in tan or brown, Smo and Gli2 protein positive rate was SHH, Gli2 protein was expressed both in nucleus and cytoplasm, with brown granules or tan granules (Figure 1).



**Figure 1.** Protein expression of SHH signaling pathway molecules in different tissues.

## 4. Discussion

HCC is a common digestive system malignant tumor, clinical diagnosis and treatment is difficult. Its pathogenesis has not been entirely clear, the onset process include multiple genes and signaling pathway<sup>[9–13]</sup>. SHH signaling pathway is involved in the process of embryonic development and tissue polarity regulation, studies have shown that<sup>[14–18]</sup>, activation of SHH signaling pathway may participate in a variety of tumor formations, closely associated with the occurrence and development of malignant tumors. This study is to observe the protein expression of Shh, Smo and Gli2 related to SHH signaling pathway in HCC tissues and HCC cell lines, analyze the function of SHH signaling pathway on the pathogenesis of HCC process, and lay a foundation for the further study of HCC pathogenesis.

Hedgehog signaling pathway was firstly found in fruit flies, adjusting the directional differentiation and embryonic

development<sup>[19]</sup>. Studies have reported<sup>[20]</sup>, SHH signaling pathway is not activated in normal liver cells in mature mice. In this study, Shh, Smo and Gli2 showed no protein positive expression in normal liver tissues, confirmed that the SHH signaling pathway is not activated in normal liver tissue, consistent with literature reports. SHH signaling pathway gene present high expression in HCC tissues<sup>[21–24]</sup>, suggesting SHH signaling pathway are highly activated. Positive expression rate of Shh, Smo and Gli2 protein in HCC tissue was 63.3%, 67.7% and 66.7%, respectively according to the results of this study, indicating that positive expression of SHH signaling pathway may participate in the occurrence and development of liver cancer<sup>[25]</sup>. In this study, HCC cell lines HepG2 showed a lower expression than Huh7, comparison between the two groups has no statistical difference ( $P>0.05$ ), which may be due to different level of activation of SHH signaling pathway in various liver cancer cell lines. SHH Mrna expression was observed in cancer cell lines of digestive tract's different parts, showing that the activation of SHH pathway in the digestive tract cell lines is closely related to the occurrence and development of tumors<sup>[26]</sup>.

This study also detected SHH signaling pathway factors in part of cirrhosis tissue, confirmed SHH signaling pathway activated in liver cirrhosis. Other studies on primary biliary cirrhosis tissue, found a high expression level of SHH signaling pathway molecules, suggesting abnormal activation of SHH are closely related to the formation and development of liver cirrhosis, blocking SHH pathway may become a new target for prevention and control of liver fibrosis<sup>[24]</sup>.

A large number of activated SHH signaling pathway molecules in HCC tissue, suggesting SHH signaling pathway may participate in the occurrence of liver cancer, and targeted therapy for SHH signaling pathway may become a new way to prevent and cure of liver cancer.

### Conflict of interest statement

We declare that we have no conflict of interest.

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