Effects and related mechanisms of serotonin on malignant biological behavior of hepatocellular carcinoma via regulation of Yap

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ABSTRACT

5-hydroxytryptamine (5-HT, serotonin) and Yes-associated protein (Yap), which act as a mitogen and an oncogene, respectively, play an important role in tumors. Here, we investigated whether 5-HT could affect the hepatocarcinogenic process via promoting the activation and expression of Yap, as well as the possible underlying molecular mechanisms. We found that 5-HT promoted hepatoma cell proliferation, invasion and metastasis via regulating Yap expression in vitro and in vivo, and Yap knockdown had opposite effects. Furthermore, 5-HT activated 5-HT2B R to promote Yap expression via upregulating the pERK level. Inhibitors of 5-HT2B R and ERK attenuated the overexpression of Yap and promotional effects of 5-HT in vitro and in vivo. As a result, 5-HT affected the malignant biological behavior of hepatoma cells via the 5-HT-5-HT2B R-pERK-Yap axis. Therefore, 5-HT and Yap may be prognostic predictors and potential therapeutic targets for HCC patients in the future.

INTRODUCTION

In China, hepatocellular carcinoma (HCC) has a high incidence and mortality that are still increasing, causing a huge impact on national health. Recent studies of 5-HT have indicated the association between 5-HT and cancer [1–7]. Increasing evidence has shown that 5-HT has serum-like effects on certain types of hepatoma cells to promote proliferation, invasion and metastasis [5, 7]. Moreover, animal experiments also confirm the promoting effects of serotonin in xenograft models [5, 7].

Landmark studies have implicated that the activation of the Hippo-Yap signaling pathway influences liver cell fate [8]. Engagement of the Hippo signaling pathway results in inactivation and phosphorylation of Yap [9]. By contrast, Yap localizes to the nucleus and exerts its transcriptional activity mainly by interacting with the TEAD [10]. As an oncogene, Yap is associated with many types of tumors [11–17]. Although the Hippo-Yap signaling pathway plays a critical role in organ size regulation and tumorigenesis, rare study is available on the regulation of Yap via the Hippo-independent signaling pathway.

In this study, we investigated whether 5-HT could affect the hepatocarcinogenic process via promoting activation and expression of Yap, as well as the possible underlying molecular mechanisms. We found that 5-HT promoted hepatoma cell proliferation, invasion and metastasis via regulating Yap expression in vitro and in vivo, and Yap knockdown had opposite effects. Furthermore, 5-HT activated 5-HT2B R to promote Yap expression via upregulating the pERK level. Inhibitors of 5-HT2B R and ERK attenuated the overexpression of Yap and promotional effects of 5-HT in vitro and in vivo. As a result, 5-HT affected the malignant biological behavior of hepatoma cells via the 5-HT-5-HT2B R-pERK-Yap axis. Therefore, 5-HT and Yap may be prognostic predictors and potential therapeutic targets for HCC patients in the future.

RESULTS

5-HT promotes the malignant biological behavior of hepatoma cells via regulating Yap expression

Previous studies have shown that 5-HT exerted serum-like effects on certain types of hepatoma cells.
In our study, 5-HT elevated the viabilities of hepatoma cells, especially serum-deprived HepG2 and HHCC cells. Additionally, medium containing 100 μM serotonin exhibited the strongest promotion effect (Figure 1A). Additionally, faster wound closure (Figure 1B–1C) and more cell penetration (Figure 1D–1E) were observed in HepG2 and HHCC cells cultured with serotonin.

In order to explored the promoting effects of 5-HT on hepatoma cell, we investigated the promotion effects of 5-HT on Yap expression. Among the cells, L-O2 showed the lowest Yap expression among the selected cells (Figure 2A–2B). Additionally, the HepG2 and HHCC showed the highest Yap expression (Figure 2A–2B). To further validate the promotion effects of 5-HT on Yap expression, the expression of Yap in HepG2 and HHCC cells in the absence or presence of 100 μM serotonin was assessed. Additionally, Yap and Connective tissue growth factor (CTGF, downstream target of Yap) expression levels were higher with the administration of 5-HT than those in the control group (Figure 2C–2D).

Figure 1: 5-HT promotes the proliferation, invasion and metastasis of hepatoma cells. (A) 5-HT significantly promoted HepG2 and HHCC cell proliferation in SFM as evaluated by the MMT assay at the indicated times. (B–C) 5-HT induced faster wound closure in HepG2 and HHCC cells (B), and the results were quantified using the healing rate (C). (D–E) 5-HT induced more cell penetration at 24 h in HepG2 and HHCC cells (D), and the penetrated cells were counted (E). *P < 0.05, **P < 0.01.
Next, we ascertained whether Yap acted as a downstream target of 5-HT in hepatoma cell. After transfection with Yap-siRNA, HepG2 and HHCC cells were cultured in SFM (serum-free medium) containing 5-HT. First, we explored the efficacy of transfection of the two types of Yap-siRNA (Figure 2E–2F). Next, the cell proliferation and cell cycle of HepG2 and HHCC cells transfected with Yap-siRNA were analyzed. As a result, Yap-siRNA significantly reduced HepG2 and HHCC cell proliferation in the presence of 5-HT (Figure 2G). Moreover, the S+G2/M phase cell ratio was increased in the presence of 5-HT and was significantly reduced in Yap-siRNA-transfected hepatoma cells (Figure 2H–2I). However, cell apoptosis was not affected by 5-HT or Yap-siRNA transfection (Figure 2J). To determine the potential metastatic promotion effect of Yap affected by 5-HT, we investigated cell motility and invasion abilities. Compared with the HepG2 and HHCC cells in the presence of 5-HT, slower wound closure (Figure 3A–3B) and less cell penetration (Figure 3C–3D) were observed in Yap-siRNA-transfected cells.

### 5-HT$_{2B}$ receptor activated by 5-HT upregulates Yap expression

Accumulated studies have shown that the 5-HT$_{2B}$ receptor plays an important role in HCC [5–7]. We investigated the expression of 5-HT receptors in hepatoma cells via qRT-PCR. 5-HT$_{2B}$R expression was the highest among all the receptors (Figure 4A). Next, we investigated whether 5-HT could affect 5-HT$_{2B}$R expression and found that 5-HT$_{2B}$R expression was significantly upregulated by 5-HT administration (Figure 4B–4C).

To investigate whether Yap expression was regulated by activated 5-HT$_{2B}$R, SB204741, an inhibitor of 5-HT$_{2B}$R, was employed. As a result, SB204741 significantly reduced the cell viability, especially at the concentration of 100 μM SB204741 (Figure 4D). Moreover, HepG2 and HHCC cells treated with SB204741 lowered the S+G2/M phase cell ratio compared with the cells only in the presence of 5-HT (Figure 4E–4F). Interestingly, SB204741 did not affect the apoptosis of HepG2 and HHCC cells, a finding that was similar to Yap-siRNA effects (Figure 4G).

As SB204741 led to inhibition of proliferation, we investigated whether SB204741 was involved in invasion and metastasis. As a result, SB204741 not only induced slower wound closure but also less cell penetration at 24 h (Figure 4H–4I).

Finally, we investigated whether activated 5-HT$_{2B}$R could promote Yap expression. As a result, the expression of Yap and CTGF at the mRNA and protein levels were both inhibited by SB204741 (Figure 5A–5B). Additionally, immunofluorescence was used to detect the subcellular localization of Yap after culture in SFM containing 5-HT with or without SB204741. We found that Yap was predominantly expressed in nuclei in the presence of 5-HT, whereas it was mainly located in the cytoplasm after administration of SB204741 (Figure 5C).

### pERK, which is upregulated by activated 5-HT$_{2B}$R, promotes Yap activation

To investigate the factors involved in the 5-HT-Yap pathway, inhibitors of AKT, ERK and neutralizing antibodies to TGF-β were used. As shown in Figure 6A, the expression of Yap were all inhibited by the inhibitors, and PD0325901 (an inhibitor of ERK) had the best inhibitory effect (Figure 6A). Moreover, the pERK level was upregulated by 5-HT, indicating that pERK might be a target of 5-HT (Figure 6B).

To define the relationship among 5-HT$_{2B}$R, pERK and Yap, SB204741 (an inhibitor of 5-HT$_{2B}$R), PD0325901 and Yap-siRNA were used. Firstly, the pERK and ERK expression levels were measured in cells cultured in medium containing 5-HT with or without SB204741. As a result, the expression levels of 5-HT$_{2B}$R and pERK were significantly inhibited by SB204741 (Figure 6C). However, PD0325901 only inhibited the expression of pERK, Yap and CTGF (Figure 6D). Additionally, the inhibition of Yap did not affect pERK and 5-HT$_{2B}$R expression (Figure 6E). Moreover, immunofluorescence showed the similar results and we found that Yap was predominantly expressed in nuclei in the presence of 5-HT, whereas it was mainly located in the cytoplasm after administration of the inhibition of ERK (Figure 6F).

Thus, we hypothesized that 5-HT affected the malignant biological behavior of hepatoma cells via the 5-HT-5-HT$_{2B}$R-pERK-Yap axis.

### The 5-HT-Yap axis promotes the malignant biological behavior of hepatoma cells by activating pERK and Yap in vivo

Because the in vitro study has shown that 5-HT promotes proliferation, invasion, and metastasis via the 5-HT-5-HT$_{2B}$R-pERK-Yap axis, we further investigated the function of the 5-HT-Yap axis in vivo. As a result, the tumor volume and weight of mice with 5-HT administration was larger than the control group and became smaller after treatment with SB204741 or PD0325901 (Figure 7A–7D). Additionally, SB204741 inhibited the expression levels of 5-HT$_{2B}$R, pERK, Yap and CTGF, and the expression levels of pERK, Yap and CTGF were also inhibited by PD0325901 except for the 5-HT$_{2B}$R expression level (Figure 7E). Next, immunohistochemistry of Ki67 and Yap was conducted to investigate the proliferation ability and activation of Yap in vivo. As predicted, 5-HT promoted cell proliferation in vivo, and the proliferation promotion effects induced by 5-HT were attenuated by SB204741 and PD0325901 (Figure 7F). Importantly, the nuclear expression of Yap induced by 5-HT was also inhibited after using SB204741 or PD0325901 (Figure 7G).
Figure 2: Yap, which is regulated by 5-HT, promotes proliferation. (A–B) Yap expression in hepatoma cells was higher than those in normal hepatocytes in the presence of 5-HT at the protein (A) and mRNA levels (B) and was especially higher in HepG2 and HHCC cells among the hepatoma cells. (C–D) 5-HT promoted Yap expression in HepG2 and HHCC at the protein (C) and mRNA levels (D). (E–F) Yap expression was significantly downregulated using Yap-siRNA at the protein (E) and mRNA levels (F) in HepG2 and HHCC cells. (G) Inhibition of Yap inhibited the proliferation promotion effect induced by 5-HT. (H–I) Yap inhibition reduced the S+G2/M phase cell ratio in HepG2 and HHCC cells (H), and the marker proteins of the cell cycle were also detected (I). (J) 5-HT administration and Yap inhibition did not affect cell apoptosis. Red asterisk: control group vs 5-HT group; Green asterisk: 5-HT group vs 5-HT+Yap-siRNA group; **P < 0.01.
DISCUSSION

As a neurotransmitter in the central nervous system, peripheral serotonin (95%) was shown to regulate the digestive system, respiratory system, cardiovascular system and immune system, as well as promoting the proliferation of different cell types [5, 7]. However, the mechanisms by which serotonin acted as a mitogen were still confusing. Moreover, whether serotonin acted as a mitogen through a receptor-dependent or receptor-independent pathway was worth exploring. In this study, we demonstrated that serotonin upregulated the pERK level via activating 5-HT₂B-R and revealed that Yap was a key downstream effector in the 5-HT-5-HT₂B-R-pERK-Yap axis.

Yap was a co-activator of the Hippo signaling pathway. Overexpression of Yap could aberrantly activate different target genes, including CTGF, AREG, and Gli, to promote proliferation, migration and survival [18, 19]. Additionally, Yap was related to diverse tumors [11, 20, 21]. However, the relationship between serotonin and Yap was poorly understood. Interestingly, we found serotonin promoted the Yap expression and cancer process and the results were further confirmed by animal experiments. Yap expression was, no doubt, higher in hepatoma cells in the presence of serotonin. Additionally, Yap was required for proliferation, invasion and metastasis of hepatoma cells. Our data suggested that inhibition of Yap via Yap-siRNA induced inhibition of proliferation, G0/G1 phase cell arrest, slower wound closure and less cell penetration. However, cell apoptosis was not affected by Yap-siRNA or SB204741, possibly because serotonin mainly activated CTGF, which was only associated with proliferation. Additionally, Yap

Figure 3: Yap promoted invasion and metastasis of HepG2 and HHCC cells. (A–B) Inhibition of Yap induced slower wound closure in HepG2 and HHCC cells (A), and the healing rate was used for quantification (B). (C–D) Inhibition of Yap induced less cell penetration (C), and the penetrated cells were counted for quantification (D). ns: no statistical significance, **P < 0.01.
Figure 4: 5-HT$_{2B}$R promoted the proliferation, invasion and metastasis of HepG2 and HHCC cells. (A) The expression of 5-HT receptors in different types of hepatoma cells. The 5-HT receptors expression of SMMC-7721 were considered as 1. (B–C) 5-HT promoted the expression of 5-HT$_{2B}$R in HepG2 and HHCC at the protein (B) and mRNA levels (C). (D) Inhibition of 5-HT$_{2B}$R inhibited the promotion effect on proliferation induced by 5-HT. (E–F) Inhibition of 5-HT$_{2B}$R reduced the S+G2/M phase cell ratio in HepG2 and HHCC cells (E), and cell cycle marker proteins were also detected (F). (G) 5-HT administration and 5-HT$_{2B}$R inhibition did not affect cell apoptosis. (H) Inhibition of 5-HT$_{2B}$R induced slower wound closure in HepG2 and HHCC cells, quantified by the healing rate. (I) Inhibition of 5-HT$_{2B}$R induced less cell penetration, quantified by the number of penetrated cells. Blue asterisk: 5-HT group vs 5-HT+SB204 10 μM; Green asterisk: 5-HT group vs 5-HT+SB204 100 μM; ns: no statistical significance, *$P$ < 0.05, **$P$ < 0.01.
affected cell apoptosis by interacting with p73; however, the expression of p73 was not altered by using Yap-siRNA or SB204741. In agreement with our findings, Perra et al. reported that Yap was accumulated in the foci of preneoplastic hepatocytes after DENA treatment, and Yap accumulation was paralleled by its target gene upregulation [20]. Additionally, a study conducted by Zhou et al. demonstrated that approximately 30% of HCC patients showed lower Yap phosphorylation [22]. Surprisingly, Yap activation was also regulated by the Hepatitis B virus X protein via CREB [23].

Accumulated studies have indicated that serotonin worked by binding to serotonin receptors. A study conducted by Soll et al. indicated that serotonin promoted hepatocyte proliferation via upregulating the expression of 5-HT$_{2B}$R [5]. Similarly, 5-HT$_{2B}$R was also overexpressed in liver regeneration models [24]. Additionally, Soll et al. found that the expression levels of 5-HT$_{1B}$R and 5-HT$_{2B}$R in HCC patients were 32% and 35%, respectively, and were associated with an increased proliferation index [6]. Moreover, a study conducted by Liang et al. demonstrated that serotonin downregulated FoxO3a to promote

![Figure 5: 5-HT$_{2B}$R promoted the activation and expression of Yap.](image)

(A, B) The expression levels of Yap and CTGF (a Yap target) were downregulated by SB204741 at the protein (A) and mRNA levels (B). (C) 5-HT induced the entry of Yap into the nucleus, and the inhibition of 5-HT$_{2B}$R led to the retention of Yap in the cytoplasm and inactivation of Yap. **P < 0.01.
Figure 6: pERK upregulated by activated 5-HT_{2B}R promotes Yap activation and expression. (A) PD0325901, an inhibitor of ERK, showed the greatest inhibition of Yap expression induced by 5-HT among the inhibitors. (B) pERK expression was promoted by 5-HT in HepG2 and HHCC cells. (C) The expression levels of pERK and 5-HT_{2B}R were inhibited by SB204741 in HepG2 and HHCC cells in the presence of 5-HT. (D) The Yap and CTGF expression levels were inhibited by PD0325901; however, the expression of 5-HT_{2B}R was not affected. (E) Only Yap expression was inhibited by Yap-siRNA in the HepG2 and HHCC cells. (F) 5-HT induced the entry of Yap into the nucleus, and the inhibition of ERK led to the retention of Yap in the cytoplasm and inactivation of Yap.

MATERIALS AND METHODS

Cell lines and cell culture

The human immortalized normal hepatic cell line (L-O2) and HCC cell line (HepG2, Hep3B, Huh-7, HHCC, SMMC-7721 and MHCC-97H) were obtained from the cell bank of Chinese Academy of Sciences in Shanghai. The cells were cultured in Dulbecco’s Modified Eagle’s Medium (DMEM; Hyclone, South Logan, UT, USA) containing 10% fetal bovine serum (Gibco, Grand Island, NY, USA) 1% penicillin/ streptomycin at 37°C with 5% CO₂ in incubator. During the administration, the cells were cultured only in DMEM without FBS. The 5-HT_{2B} receptor inhibitor SB204741 (Santa Cruz Biotechnology, Santa Cruz, CA, USA) and ERK inhibitor PD0325901 (Selleck Chemicals, Houston, TX, USA) were used to treat cells at concentration of 100 mM and 1 mM respectively according to the manufacturer’s instructions.

The HCC cells were harvested, re-plated and cultured in incubator overnight to allow adhesion of cells. Before administration, the medium of cells was replaced by DMEM without FBS for synchronization. The groups of the study included control group (serum free medium, SFM), 5-HT group (SFM+5HT), 5-HT+SB204 group (SFM+5HT+SB204741), 5-HT+PD032 group (SFM+5HT+PD0325901) and 5-HT+Yap-siRNA group (SFM+5HT+Yap-siRNA). During the administration, the cells were pretreated in the medium containing SB204741 or PD0325901 for 30 min before addition of 5-HT.

Small interfering RNA (siRNA) transfection

The human Yap-siRNA and control siRNA were synthesized by GenePharma (Shanghai, China). The siRNA sequences were shown in Supplementary Table 1. The cells were transfected with a mixture of siRNA using Lipofectamine 2000 (Invitrogen, Carlsbad, CA, USA) according to the manufacturer’s instructions. After cultured for 8 h, the medium was replaced. And the knockdown efficiency of Yap was detected by qRT-PCR and Western blot assays after replacement of medium for 24 h.

RNA isolation and quantitative real-time PCR (qRT-PCR)

Total RNA from cells and tissue samples was extracted using TRIzol reagent (Invitrogen). qRT-PCR was performed using the SYBR Premix Ex Taq Kit (Takara,
Tokyo, Japan) and TaqMan microRNA assays (Applied Biosystems, Foster City, CA, USA). The 2^{-ΔΔCt} method was selected to calculate the relative expression level of target genes. The primers are shown in Supplementary Table 2.

**Western blot analysis**

Proteins were extracted from cell and tissue lysates according to the manufacturer’s instructions. Western blotting was performed as previously described [30].

**MTT assays**

Cells were seeded into 96-well plates (1 × 10^4/well). After administration of 5-HT, SB204741 or Yap-siRNA transfection, MTT (Sigma-Aldrich, St. Louis, MO, USA) was added to the platelets for staining at 37°C for 4 h, and 150 μl of DMSO (Sigma) was subsequently added after discarding the culture medium. The absorption value was measured at OD490.

**Cell cycle and cell apoptosis assays**

To study the effects of the 5-HT-Yap pathway on the cell cycle, cells were seeded in 6-well plates with treatments. Next, cells were collected and fixed in 70% ethanol at −20°C overnight. Subsequently, the cells were incubated with PI staining solution (50 μg/mL, Sigma) containing RNase A (100 μg/mL, Sigma) for 20 min at room temperature in the dark. Finally, the cells were analyzed by FACSCalibur (BD Biosciences, Bedford, MA, USA).

For cell apoptosis analysis, the cells treated as presented above were harvested and stained with Annexin V-FITC/PI or Annexin V-PE/7-AAD Apoptosis Detection Kit (KeyGEN BioTECH, Nanjing, China) following the manufacturer’s instructions. Stained cells were detected using the FACSCalibur instrument (BD Biosciences).

**Wound healing and cell migration assays**

Cells were plated in 6-well plates (5 × 10^5 cells/well) and then were cultured with 10% FBS to achieve a nearly confluent cell monolayer. A scratch was made carefully by a 10-μl sterile micropipette tip on the cell layer. Next, the cells were washed and the wounded monolayers were photographed to determine the wound width at time 0 h. After administration, the cells were photographed again at 24 h and 48 h.

For the transwell migration assay, the control cells or transfected cells (1 × 10^5) were added to the top
chamber, and the bottom chamber was filled with DMEM containing 5-HT and SB204741 or PD0325901. After fixing, the transwell chambers were stained with 1% crystal violet for 10 min and the cells were counted under a light microscope for quantitation.

**Immunofluorescence assays**

To explore the subcellular localization of Yap, immunofluorescence was conducted after treatment with 5-HT, SB204741 or PD0325901. Immunofluorescence was performed as previously described [31].

**Immunohistochemical staining**

Immunohistochemical staining was performed using a streptavidin peroxidase-conjugated (SP-IHC) method with Yap or Ki-67 antibodies. Evaluation of the immunohistochemical staining was conducted by two independent pathologists who were blinded to the administration. The assessment criteria were based on a previous study [32].

**In vivo experiments**

Sixteen female BALB/c nude mice (4–5 weeks old, 14–18 g) were purchased from Animal Feeding Center of Xi’an Jiaotong University Health Science Center. All mice were housed in an IVC system with a controlled environment and temperature (24 ± 1°C). All of the mice were fed standard rodent chow and water ad libitum and were adapted to the environment for 7 days before use. All mice were inoculated subcutaneously by HepG2 cell (1 × 10⁵) suspended in DMEM on the left flank. After subcutaneous injection for 5 days (diameter of xenograft tumor was about 3mm), sixteen nude mice were randomly allocated into 4 groups as follows (4 mice per group): (1) control group: the mice were treated with physiological saline. (2) 5-HT group: the mice were treated with 5-HTP (50 mg/kg) every other day via subcutaneous injection. (3) 5-HT+SB204 group: the mice were treated with 5-HTP (50 mg/kg) every other day and SB204741 (10 mg/kg) twice a day via subcutaneous injection, synchronously. (4) 5-HT+PD03 group: the mice were treated with 5-HTP (50 mg/kg) every other day via subcutaneous injection, and PD0325901 (10 mg/kg) once a day via intragastric gavage. Tumor volume was measured twice a week and calculated as (π×length×width×height)/6. All mice were sacrificed after subcutaneous injection for 4 weeks and the xenograft tumors were explanted for weighing and further experiments. Animal care was in compliance with criteria outlined in the Guide for the Care and Use of Laboratory Animals established by the US National Institutes of Health. The study was approved by the Animal Research Committee of Xi’an Jiaotong University Health Science Center.

**Statistical analysis**

The data were expressed as means ± SD. The correlation differences between the respective groups were evaluated by either ANOVA or nonparametric test. A P-value less than 0.05 was considered to be statistically significant.

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**CONFLICTS OF INTEREST**

The authors declare no conflicts of interest.

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