Expression and inhibitory effects of p53-upregulated modulator of apoptosis in gallbladder carcinoma

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Abstract. The p53-upregulated modulator of apoptosis (PUMA) has been reported to be involved in various types of cancer. However, its potential biological role in gallbladder carcinoma (GBC) has not been fully elucidated. The present study aimed to determine the expression levels of PUMA and its biological effects on GBC. The mRNA and protein expression levels of PUMA in GBC tissues and cell lines were measured using reverse transcription-quantitative PCR and western blotting, respectively. The effects of PUMA overexpression on cell viability, proliferation and invasive ability were determined in vitro using the MTT, colony formation and Transwell invasion assays, respectively. The apoptotic rates were detected using the Annexin V-FITC apoptosis detection kit. Furthermore, follow-up of patients with GBC was performed to identify the association between PUMA expression levels and GBC prognosis. The results of the present study demonstrated that the expression levels of PUMA were significantly lower in the GBC tissues and cell lines compared with those in adjacent normal gallbladder tissues and normal gallbladder cells, respectively. Further experiments indicated that overexpression of PUMA inhibited the viability, proliferation and invasive ability of GBC cells compared with those in the control group. Low PUMA expression levels were associated with a short overall survival time in patients with GBC. In conclusions, PUMA may act as a tumor suppressor in GBC and may serve as a potential novel treatment target for human GBC.

Introduction

Gallbladder carcinoma (GBC) was the fifth most common malignant tumor of the digestive system and the most common malignant tumor of the biliary tract worldwide in 2015 (1,2). Early diagnosis and radical surgical resection are the most effective and preferred treatment approaches for patients with GBC, resulting in improved long-term survival (3,4). However, due to the atypical early symptoms, and frequently rapid and asymptomatic progression, a large proportion of patients with GBC are likely to be diagnosed with late stage cancer; thus, resectable GBC accounts for only 15-47% of all cases (5,6). The aforementioned features of GBC may account for the unsatisfactory prognosis of patients for GBC. The median survival time of patients with GBC is only ~6 months, with a 5-year survival rate of <5% worldwide in 2019 (7). Eventually, the majority of patients with GBC succumb to metastatic disease (8).

Previous studies have reported that p53-upregulated modulator of apoptosis (PUMA) serves important roles in inducing apoptosis in both normal and tumor cells (9-11). In vitro experiments have demonstrated that PUMA induced esophageal and lung cancer cell apoptosis, and significantly attenuated the viability and proliferative ability of head and neck cancer cell lines; consistent with the in vitro results, the inhibitory effect of PUMA on the tumor growth was confirmed in nude mice in vivo (9-12). The aforementioned studies indicate that PUMA serves a crucial inhibitory role in tumor development. In addition, PUMA is associated with the progression, recurrence and metastasis and prognosis of colon and prostate cancers (13,14).

However, the expression profile and potential biological roles of PUMA in patients with GBC have not been fully elucidated to date. Therefore, the current study aimed to determine the expression levels of PUMA in GBC tissues and cell lines, and to further investigate its effects in GBC pathogenesis.
Materials and methods

Clinical specimen collection. A total of 18 paired GBC and adjacent normal gallbladder tissues were collected from patients who underwent surgical treatment between December 2016 and December 2018 in the Second Xiangya Hospital (Changsha, China), including 6 men and 12 women, with the mean age of 52 years (range, 42–68 years) in Nevin’s stage I–III (15). All patients enrolled in the present study had not undergone any treatment prior surgery. The fresh GBC and adjacent normal gallbladder tissues were stored at -80°C in liquid nitrogen until RNA extraction. Written informed consent for participation and publication were obtained from all patients. Follow-up was performed after surgery until December 2019, and no cases were lost to the follow-up. This study was approved by the Research Ethics Committee of the Second Xiangya Hospital (approval no. 179 in 2017).

Cell culture. The normal gallbladder cells were obtained by primary culture of cells isolated from healthy gallbladder wall tissue, which was obtained from six patients with gallbladder polyps undergoing cholecystectomy, including three men and three women, with the mean age of 50 years (range, 44–65 years). Written informed consent for participation and publication were obtained from all patients. The gallbladder wall tissue was transferred to ice-cold Ca<sup>2+</sup>-free Hanks solution with 0.5 mM EGTA and cut into 1-cm<sup>2</sup> fragments under sterile conditions. Subsequently, the tissue was placed on a 60-mm culture dish and treated for 45 min at room temperature with 0.025% trypsin (Thermo Fisher Scientific, Inc.) in PBS containing 0.02% EDTA-2Na. The tissue was agitated every 15 min using a pipette to promote the release of the cells. The gallbladder cells were collected by centrifugation at 1,200 x g for 5 min and counted in a CC-108 microcell counter (TOA Corporation).

Human GBC cell lines GBC-SD, SGC-996, NOZ and EHGB-1 obtained from the Institute of Biochemistry and Cell Biology of the Chinese Academy of Sciences. The GBC cells were cultured in Dulbecco's modified Eagle's medium (Gibco; Thermo Fisher Scientific, Inc.) containing 10% fetal bovine serum and 1% penicillin-streptomycin (Invitrogen; Thermo Fisher Scientific, Inc.) at 37°C with 5% CO<sub>2</sub>.

RNA extraction and reverse transcription-quantitative (RT-qPCR). The TRIzol<sup>®</sup> Reagent (Invitrogen; Thermo Fisher Scientific, Inc.) was used to extract RNA from clinical tissues and cell lines, which was subsequently treated with DNase I (Invitrogen; Thermo Fisher Scientific, Inc.) for purification according to the manufacturer’s instructions. The miRoute miRNA First-Strand cDNA Synthesis kit (Takara Bio, Inc.) was used to perform reverse transcription according to the manufacturer's instructions. qPCR was performed using the SYBR<sup>®</sup> Green PCR detection kit (Takara Bio, Inc.) in a 7500 Real-Time PCR system (Applied Biosystems; Thermo Fisher Scientific, Inc.) according to the manufacturer's instruction with the following primers: PUMA forward, 5'-TGAAGA GCAAATGACGCAAACG-3' and reverse, 5'-CAGACGACA GGATTCCACGTCT-3'; and GAPDH forward, 5'-CTGACTT CAACAGCGACACCCA-3' and reverse, 5'-CACCCTGTT GCTGTAGCCCAA-3'. The expression levels of GAPDH were used to normalize the results and the 2<sup>ΔΔCT</sup> method was used to calculate the relative expression levels of target gene (16).

Western blotting. Total protein was extracted from clinical tissues and cell lines using TRIzol lysis buffer (50 mM Tris pH 8.0, 120 mM NaCl, 0.5% sodium deoxycholate, 0.5% NP-40, 0.1% SDS, 1 mM EDTA, 50 mM NaF, 1 mM Na<sub>3</sub>VO<sub>4</sub>, 1 mM PMSF and 2 µg/ml aprotinin) on ice for 30 min. The protein concentration was determined by BCA assay, and 50 µg protein/lane was separated by 10% SDS-PAGE at 120 V for 60 min and transferred to PVDF membranes. The membranes were blocked with 5% non-fat milk in TBS (10 mM Tris pH 7.4 and 150 mM NaCl) and 0.1% Tween-20 (TBST) at room temperature for 60 min, followed by incubation with primary antibodies against PUMA (1:1,000; cat. no. ab39306), E-cadherin (1:10,000; cat. no. ab40772), vimentin (1:1,000; cat. no. ab92547), Bax (1:1,000; cat. no. ab32503), Bcl-2 (1:2,000; cat. no. ab18288) (all from Abcam) and β-actin (1:1,000; cat. no. A2228; Sigma-Aldrich; Merck KGaA) at 4°C for 12 h. Subsequently, the membranes were washed three times with TBST for 5 min and incubated with HRP-conjugated goat anti-rabbit IgG (1:2,000; cat. no. sc-2004) or anti-mouse IgG (1:2,000; cat. no. sc-2005) (both from Santa Cruz Biotechnology, Inc.) secondary antibody at 37°C for 60 min. The protein bands were visualized with an enhanced chemiluminescence system (Thermo Fisher Scientific, Inc.) according to the manufacturer's instructions. The Gel Doc 2000 imaging system (Bio-Rad Laboratories, Inc.) was used for densitometric analysis of the protein bands with ImageJ software (version 1.8.0.112; National Institutes of Health) according to the manufacturer's instructions. β-actin was used as the internal control.

Plasmid generation and cell transfection. The PUMA sequence was synthesized and subcloned into the LV-BBC3 vector (22944-1) (Shanghai GeneChem Co., Ltd.). Ectopic expression of PUMA was achieved by the PUMA lentivirus transfection, and an empty vector (KL8781-1) (Shanghai GeneChem Co., Ltd.) was used as a negative control. Prior to transfection, human GBC cell lines were cultured in complete medium without antibiotics for at least 24 h. When the GBC-SD and EHGB-1 cell confluence reached 30-50%, the cells were washed with 1X PBS once and transfected with the LV-BBC3 (22944-1) (5x10<sup>4</sup> TU/ml; 40 µl; MOI, 20) and empty (KL8781-1) vector at 37°C for 72 h, in the presence of Lipofectamine<sup>®</sup> 2000 (Invitrogen, Thermo Fisher Scientific, Inc.) according to the manufacturer’s instructions. At 48 h post-transfection, the cells were harvested, and the expression levels of PUMA were determined by RT-qPCR.

MTT assay. At 48 h post-transfection, the cells were seeded into 96-well plates (2x10<sup>4</sup> cells/well). Following culture for 4 h at 37°C, the supernatant was discarded, and 150 µl DMSO was added to each well. After shaking for 10 min, 100 µl cell suspension was placed into another 96-well plate with 100 µl DMSO solution as a control. The Cell Proliferation Reagent Kit I (MTT; Roche Applied Science) was used at 37°C for 4 h to determine the cell viability. The absorbance values were determined at 450 nm using a spectrophotometer (Omega Bio-Tek, Inc.).
Colony formation assay. At 48 h post-transfection, the cells were harvested, trypsinized and counted. A total of 500 cells/well were placed into a 6-well plate and cultured at 37°C. The culture medium was refreshed every 3 days. Following 14-day culture, the cells were washed with 1X PBS, fixed with 3.7% methanol at 37°C for 15 min and stained with 0.1% crystal violet at 37°C for 10 min (Sigma-Aldrich; Merck KGaA). Visible colonies ≥50 cells were manually counted using an optical microscope (Olympus Corporation).

Transwell invasion assay. The invasive ability of the cells was evaluated using an 8-µm pore Transwell chamber. The upper chamber was pre-coated with Matrigel at 37°C overnight. At 48 h post-transfection, the cells (1x10^5) were suspended in serum-free DMEM and plated into the upper chamber. The lower chamber was filled with DMEM containing 10% FBS. Following 48-h incubation at 37°C with 5% CO₂, the chamber was removed, and the cells remaining in the upper chamber were cleaned with cotton wool. The invasive cells were washed with 1X PBS, fixed with 100% methanol at 37°C for 30 min and stained with 0.1% crystal violet at 37°C for 20 min. Finally, images of five random fields were captured using an optical microscope (x400 magnification; Olympus Corporation) for quantification analysis. Each experiment was repeated three times.

Flow cytometry. The cells were inoculated in a 6-well plate (1x10^5 cells/well) and divided into four groups: GBC-SD control, GBC-SD PUMA overexpression, EHGB-1 control and EHGB-1 PUMA overexpression. The cells were collected, washed and resuspended in PBS. The Annexin V-FITC apoptosis detection kit (Beckman Coulter, Inc., USA) was used to detect apoptotic cells with Annexin V-FITC and PI double staining according to the manufacturer's instructions. The analysis was performed using a BD FACSARia™ II flow cytometer (Becton, Dickinson and Company), and the data were analyzed using CellQuest Pro software (version 5.1; Becton, Dickinson and Company).

Statistical analysis. Data are presented as the mean ± standard deviation. SPSS 17.0 software (SPSS, Inc.) was used for statistical analyses. The differences were analyzed using
paired-samples Student’s t-test for two paired groups, independent sample Student’s t-test for two independent groups and one-way ANOVA with the Bonferroni correction for multiple groups. P<0.05 was considered to indicate a statistically significant difference.

Results

PUMA is expressed at low levels in GBC tissues and cell lines. RT-qPCR and western blotting were performed to determine the levels of PUMA expression in GBC and adjacent normal gallbladder tissues, as well as in GBC cell lines. As demonstrated in Fig. 1A and B, 77.8% (14/18) of GBC tissues exhibited decreased mRNA expression levels of PUMA compared with those in the adjacent tissues. Compared with those in the adjacent normal gallbladder tissues, the expression levels of PUMA were significantly lower in GBC tissues. Furthermore, the expression levels of PUMA were also significantly lower in GBC cell lines compared with those in primary normal gallbladder cells. As PUMA expression levels were lower in GBC-SD and EHGB-1 cell lines compared with those in the other two cell lines, the GBC-SD and EHGB-1 cell lines were used in the subsequent experiments. Following PUMA overexpression vector transfection, the levels of PUMA protein expression were detected; as presented in Fig. 1C, the expression levels of PUMA protein were significantly higher in PUMA-overexpressing EHGB-1 and GBC-SD cells compared with those in the corresponding control groups.

Overexpression of PUMA inhibits GBC cell proliferation and invasion. MTT and colony formation assays were performed to verify the function of PUMA in GBC. The expression levels of PUMA in GBC-SD and EHGB-1 cells were upregulated by transfection with LV-BBC3 (22944-1) vector, and the empty (KL8781-1) vector was used as a negative control (Fig. 2A). As presented in Fig. 2B, PUMA mRNA expression levels were significantly upregulated in GBC-SD and EHGB-1 overexpression groups compared with those in the negative control groups. The proliferation of GBC-SD and EHGB-1 cell lines was significantly inhibited in the PUMA overexpression groups compared with that in the control groups, and the inhibitory effect appeared to become stronger with the increasing time post-transfection (Fig. 2C and D). Furthermore, the colony formation ability of the GBC-SD and EHGB-1 cells was detected by colony formation assay; as demonstrated in Fig. 3A, compared with that in the negative control group, the colony formation ability was significantly suppressed in the PUMA overexpression groups.

Transwell invasion assay was used to evaluate the effects of PUMA on the cell invasive ability in vitro. The results demonstrated that compared with those in the negative control groups, the numbers of invasive cells were significantly lower in the PUMA overexpression groups (Fig. 3B). These results...
suggested that PUMA may act as a negative regulator of the invasive ability of GBC cells.

PUMA inhibits the epithelial-mesenchymal transition (EMT), and promotes the upregulation of Bax and the downregulation of Bcl-2. As the EMT process serves a critical role in cancer cell migration and invasion, the expression levels of EMT-induced markers were detected in GBC cells. As demonstrated in Fig. 4A, the protein levels of E-cadherin were upregulated, whereas the levels of vimentin were downregulated in the PUMA overexpression groups compared with those in the negative control groups. In addition, the protein expression levels of Bax were upregulated, and the levels of Bcl-2 were downregulated in the PUMA overexpression groups compared with those in the negative control groups (Fig. 4B). These results suggested that PUMA may regulate the Bax/Bcl-2 signaling pathway and the EMT to inhibit the invasive ability of GBC cells.

PUMA promotes apoptosis in GBC cells. Following overexpression of PUMA, the apoptotic rates of GBC-SD and
EHGB-1 cells were detected by flow cytometry. As demonstrated in Fig. 5, the apoptotic rates were 5.0±1.7 and 5.4±1.8% in the GBC-SD and EHGB-1 control groups, respectively, but sharply increased to 35.0±4.5 and 28.6±3.9% in the GBC-SD and EHGB-1 PUMA overexpression groups, respectively (both P<0.01).

**PUMA is a predictor of a favorable prognosis in patients with GBC.** Patients with GBC included in the present study were followed up. According to the relative PUMA mRNA expression level and the logarithm of 2^ΔΔCt value was zero as a cutoff point, the patients were divided into PUMA high (n=4) and low (n=14) groups. No deaths occurred in the two groups in first three months after surgery. As demonstrated in Fig. 6, patients in the PUMA high group exhibited a longer overall survival time compared with those in the PUMA low group (P<0.05).

**Discussion**

Emerging evidence has suggested that alterations in a number of tumor-suppressor genes and oncogenes are associated with
the oncogenesis and progression of GBC (17,18); however, the specific molecular mechanism of GBC remains largely unclear. Due to the poor prognosis of patients with GBC, it is necessary to urgently identify the exact mechanisms of occurrence and progression of GBC.

Previous studies have reported that PUMA induces apoptosis in normal and tumor cells (19‑22); however, a limited number of studies are currently available on the role of PUMA in GBC (23‑25). Our previous study has demonstrated the expression of PUMA and its pathological significance in GBC, adjacent tissues, adenomatous gallbladder polyps and chronic cholecystitis (23). Another study has investigated the expression of PUMA and its clinicopathological significance in benign and malignant lesions of the gallbladder, concluding that PUMA is associated with the carcinogenesis, progression, metastasis and high invasiveness of gallbladder adenocarcinoma (24). Additionally, the expression of WW domain‑containing oxidoreductase and its effects on inducing apoptosis by upregulating the expression of PUMA in GBC cells have been demonstrated (25). However, in the three aforementioned studies, the expression levels of PUMA in GBC cell lines have not been reported. In the present study, the expression levels of PUMA in GBC, adjacent tissues and GBC cell lines were investigated, and a series of functional experiments were performed to study the effects of PUMA on GBC for the first time.

The biological functions of the PUMA gene have been introduced in detail in a previous study (23). The p53-dependent and non-dependent apoptotic pathways serve key roles in PUMA-induced apoptosis; previous studies have reported that the mechanism of PUMA-induced apoptosis may include the following: i) PUMA and Bcl-2/Bcl2xl combine to inhibit the Bax/Bak signaling pathway in normal or tumor cells (9); ii) PUMA combines with p53/Bcl2xl to release p53, which activates Bax to induce apoptosis (9,20,21); iii) PUMA directly binds to Bax/Bak to change its conformational, and then Bax is activated to induce cell apoptosis (21,22). However,
further studies are needed to elucidate the precise underlying mechanism of PUMA-induced apoptosis.

Our previous study has reported that PUMA is a marker that may be associated with the clinical behavior of GBC and may reflect the occurrence, development and prognosis of GBC in patient tissues (23). In the present study, the expression levels of PUMA were detected in GBC tissues and cell lines, and its functional roles on the biological behaviors of GBC cell lines were further investigated. The results demonstrated that the expression levels of PUMA were significantly lower in GBC tissues compared with those in the adjacent tissues, and were low in GBC cell lines. Further experiments were performed in vitro to demonstrate the functions of PUMA in the biological behavior of GBC; the results demonstrated that following overexpression, PUMA significantly inhibited the proliferative and invasive abilities of GBC cells compared with those of the control groups, which suggested that PUMA may serve as a tumor suppressor in GBC.

The potential mechanisms of PUMA-inhibited invasion were further investigated in the present study. The EMT is a process during which epithelial cells trans-differentiate into mesenchymal cells (26,27). The EMT endows tumor cells with the traits of migration and invasion and induces cancer stem cell properties (27,28). However, the effects of the EMT in PUMA-induced GBC cell apoptosis remain unknown. The results of the present study demonstrated that PUMA overexpression in GBC cell lines increased the protein expression levels of E-cadherin and decreased the levels of vimentin compared with those in the negative control group. These results suggested that PUMA-induced EMT reversal may account for the inhibitory effects of PUMA on GBC cell invasion. However, further studies are needed to unravel the precise signaling pathways involved in the PUMA-mediated inhibition of the EMT.

The results of the present study also demonstrated that the protein expression levels of Bax were upregulated, whereas the levels of Bcl-2 were downregulated following PUMA overexpression compared with those in the negative control groups. The Bcl-2 family serves a vital role in apoptosis (29). As a member of the Bcl-2 family, Bax promotes apoptosis; the protein expression levels of Bax are associated with the regulation of apoptosis, since high levels of the Bax protein induce apoptosis (30,31). As a suppressor of apoptosis, Bcl-2 negatively regulates apoptosis by regulating mitochondrial cytochrome c release (32). In addition, Bcl-2 binds to Bax to form a Bax/Bcl-2 polymer to inhibit the apoptotic process (33). High expression levels of Bcl-2 also maintain cell stability by inducing glutathione aggregation in the cell nucleus to reduce caspase activity (34). The results of the present study suggested that PUMA may inhibit GBC invasion at least partially by regulating the EMT. These results suggested that PUMA may serve as a potential tumor maker for the diagnosis and treatment of GBC in clinical practice. In the last decade, studies on circulating tumor cells (CTCs) have enriched the methods of detecting tumor cells (35,36); in our future work, the potential functions of PUMA in CTCs will be investigated.

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Availability of data and materials

The datasets used and/or analyzed during the current study are available from the corresponding author on reasonable request.

Authors' contributions

QL and WC conceived and designed the study. ZL and CY obtained the clinical samples and were the major contributors in acquiring the data. WZ and LW performed the cell experiments. WC and YX analyzed and interpreted the data, and drafted the manuscript. QL and WC revised the manuscript. All authors read and approved the final manuscript.

Ethics approval and consent to participate

This study was approved by the Research Ethics Committee of Second Xiangya Hospital (approval no. 179 in 2017; Changsha, China). Informed consent was obtained from all included patients.

Patient consent for publication

Not applicable.
Competing interests

The authors declare that they have no competing interests.

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