JTC-801 inhibits the proliferation and metastasis of the Hep G2 hepatoblastoma cell line by regulating the phosphatidylinositol 3-kinase/protein kinase B signalling pathway

BUFEI ZHAO¹* and TING HU²*

¹Department of Hepatopancreatobiliary Surgery, Affiliated Hospital of Beihua University, Jilin 132001; ²Department of Oncology, The First Affiliated Hospital of Changchun University of Chinese Medicine, Changchun, Jilin 130021, P.R. China

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Abstract. The increased worldwide mortality rate due to liver cancer may be attributed to the aggressive nature of the disease. Signal transduction through G-protein-coupled receptors (GPCRs) can affect a number of aspects of cancer biology, including invasion, migration and vascular remodelling. JTC-801, a novel GPCR antagonist, has demonstrated promising anticancer effects in adenocarcinoma and osteosarcoma cells. In the present study, the effect of JTC-801 on the proliferation and migration of hepatoblastoma Hep G2 cells was investigated. The Cell Counting Kit-8 assay revealed that JTC-801 markedly suppressed the growth of the Hep G2 cells. Additionally, JTC-801 significantly inhibited cell invasion and migration in a Transwell assay. Furthermore, the expression of anti-apoptotic protein B-cell lymphoma 2 decreased and the expression of the pro-apoptotic proteins active caspase-3 and apoptosis regulator BAX increased in the Hep G2 cells following JTC-801 treatment. Additionally, JTC-801 suppressed the phosphatidylinositol 3-kinase (PI3K)/protein kinase B (AKT) signalling pathway in the Hep G2 cells. Therefore, the present study revealed that JTC-801 can induce the apoptosis of Hep G2 cells by regulating the PI3K/AKT signalling pathway, which suggests that JTC-801 may be a potential novel drug target for clinical liver cancer treatment.

Introduction

The increased global mortality rate due to liver cancer may be attributed to the aggressive nature of the disease (1). Liver cancer accounts for >1 million newly diagnosed cases annually, making it a global health care problem with increasing incidence (2). The incidence of liver cancer is markedly high in developing countries and is steadily rising in developed countries (3). Although there is a continuous rise in the 5-year survival rates for liver and intrahepatic bile duct cancer, the numbers of new cases and estimated mortalities are increasing (4,5). Liver cancer is often not detectable until late in disease progression (6), and chemotherapy remains the primary treatment for numerous patients with cancer of advanced stage. However, chemotherapy has side effects, particularly in normal tissues with potent proliferative activity (7), and this therapy does not always lead to a better prognosis. Therefore, finding novel therapeutic entities against longer cancer cell survival and growth is required.

JTC-801 [N-(4-amino-2-methylquinolin-6-yl)-2-(4-ethylphenoxymethyl)benzamide monohydrochloride] is a high-affinity and selective opioid receptor-like 1 (ORL1) receptor antagonist, which belongs to the family of G-protein-coupled receptors (GPCRs) (8). Studies have demonstrated that signal transduction through GPCRs can affect numerous aspects of cancer biology, including invasion, migration and vascular remodelling (9,10). Furthermore, ORL1 may antagonize lipopolysaccharide-stimulated proliferation, migration and inflammatory signalling in human glioblastoma U87 cells (11). In a screening of agents that interact with GPCR pathways, Song et al (12) revealed that JTC-801 induces pH-dependent cell death (alkaliplosis) specifically in cancer cells, including pancreatic ductal adenocarcinoma cells, by reducing the expression of carbohydrate antigen 9, which is increased in human cancer tissues. JTC-801 may be used in the development of treatment for pancreatic cancer. Zheng et al (13) described the antitumour effects of JTC-801...
on human osteosarcoma cells. Therefore, JTC-801 may be used for the treatment of other cancer types, including liver cancer.

In the present study, the effect of JTC-801 on the Hep G2 hepatoblastoma cell line was investigated. First, the Cell Counting Kit-8 (CCK-8) assay was used to detect the proliferation of Hep G2 cells treated with JTC-801. As cancer is characterized by increased migratory/invasive capacity, the migratory and invasive abilities of Hep G2 cells following JTC-801 treatment were investigated. Furthermore, the expression of apoptotic proteins was assayed in the Hep G2 cells following JTC-801 treatment. Additionally, the phosphatidylinositol 3-kinase (PI3K)/protein kinase B (AKT) signalling pathway may have a function in inducing the apoptosis of Hep G2 cells.

Materials and methods

Cell lines and cell culture. The human Hep G2 cell line was purchased from the Shanghai Cell Bank (Shanghai Institute for Biological Science, Chinese Academy of Science, Shanghai, China). Cells were cultured in RPMI-1640 medium (GE Healthcare Lifesciences, Logan, UT, USA) supplemented with 10% fetal bovine serum (FBS; Gibco; Thermo Fisher Scientific, Inc., Waltham, MA, USA), 100 U/ml penicillin and 0.1 mg/ml streptomycin (both Sigma-Aldrich; Merck KGaA, Darmstadt, Germany) at 37°C in an incubator with 5% CO₂. When the cells entered the logarithmic growth phase they were washed 3 times with PBS and digested with 0.25% trypsin-EDTA (Beijing Solarbio Science & Technology Co., Ltd., Beijing, China). Cells were resuspended in RPMI-1640 medium containing 10% FBS to form a single-cell suspension. The cells were seeded in a 6-well plate for subsequent experiments. When the cell density reached ~80%, the cells were treated with 20 µM JTC-801 (MedChem Express, Monmouth Junction, NJ, USA), whereas the negative control group was treated with 0.1% dimethylsulfoxide (DMSO) (Amresco, LLC, Solon, OH, USA), for 24 h at room temperature.

Western blot analysis. Following the treatment of the cells with DMSO and JTC-801 for 24 h, protein was extracted with radioimmunoprecipitation assay lysis buffer with the protease inhibitor phenylmethylsulfonyl fluoride (CWBio, Beijing, China). The concentration was determined using the BCA method (CWBio). The protein was heated at 95°C for 5 min. Overall, ~20 µg protein per group was added to each well in the vertical electrophoresis tank, separated by 10% SDS-PAGE and transferred onto a polyvinylidene fluoride membrane. The membrane was blocked with 5% non-fat milk in Tris-buffered saline containing 0.1% Tween-20 (TBST; Beijing Solarbio Science & Technology Co., Ltd.) for 1 h at room temperature and incubated with primary antibodies at 4°C overnight. On the second day, the membrane was washed 3 times in TBST for 5 min and incubated with secondary antibodies at room temperature for 1 h. An enhanced chemiluminescence chromatographic substrate (ProteinTech Group, Inc., Chicago, IL, USA) was added to visualize the bands, following the washing of the membrane. The grey value was scanned by Quantity One software (version 4.6.9; Bio-Rad Laboratories, Inc., Hercules, CA, USA). β-tubulin served as the internal control. The relative expression of each protein was calculated against that of β-tubulin. Western blotting was performed with the following antibodies: Rabbit anti-human AKT (cat. no. 4691; 1:1,000 dilution), rabbit anti-human p-AKT (cat. no. 4060; 1:1,000 dilution), rabbit anti-human mechanistic target of rapamycin (mTOR; cat. no. 2983; 1:1,000 dilution), rabbit anti-human p-mTOR (cat. no. 2971; 1:1,000 dilution) (all Cell Signaling Technology, Inc., Danvers, MA, USA), rabbit anti-human B-cell lymphoma 2 (Bcl-2; cat. no. 12789-1-AP; 1:1,000 dilution), rabbit anti-human apoptosis regulator BAX (Bax; cat. no. 50599-2-1g; 1:1,000 dilution), rabbit anti-human active caspase-3 (cat. no. 19677-1-AP; 1:1,000 dilution), rabbit anti-human cyclin D1 (cat. no. 60186-1-Ig; 1:1,000 dilution), rabbit anti-human p70S6 kinase (p70S6K; cat. no. 14485-1-AP; 1:1,000 dilution), rabbit anti-human tubulin (cat. no. 10068-1-AP; 1:5,000 dilution) and the secondary horseradish peroxidase-labelled goat anti-rabbit/goat anti-mouse antibody (cat. no. 10545-2-AP; 1:5,000 dilution) (all ProteinTech Group, Inc.).

Cell Counting Kit-8 (CCK-8) proliferation test. Hep G2 cells from the conventional culture were digested with 0.25% trypsin-EDTA solution for 2 min at room temperature and counted in preparation of the cell suspension. Subsequently, 100 µl of the cell suspension was seeded onto a 96-well plate with 1,000 cells per well and 0.1% DMSO was added to the negative control group, while 0.02, 0.2, 2, 20 or 200 µM JTC-801 was added to the experimental groups. The cells were cultured in a 5% CO₂ incubator at room temperature to detect cell viability once every 24 h. Prior to detection, 10 µl CCK-8 solution (CWBio) was added to each well and incubated at 37°C for 1.5 h. The optical density (OD) was measured with a microplate reader at 450 nm and the growth curve was plotted.

Cell invasion and migration assessed by Transwell assay. Matrigel (BD Biosciences, Franklin Lakes, NJ, USA) was dissolved overnight (serum-free RPMI-1640 medium, 1:6 dilution) and 100 µl was added to the upper chamber of the 24-well Transwell insert (Merck KGaA). Once the Matrigel was evenly distributed, it was placed into a CO₂ incubator at room temperature to detect cell viability once every 24 h. Prior to the Transwell insert, 10 µl CCK-8 solution (CWBio) was added to each well and incubated at 37°C for 1.5 h. The optical density (OD) was measured with a microplate reader at 450 nm and the growth curve was plotted.
Cell apoptosis assay. The Hep G2 cells were treated with JTC-801 for 24 h, as aforementioned, and the medium was removed and replaced with 500 µl serum-free medium. Following starvation for 24 h, the cells were harvested,
digested with trypsin without EDTA, collected in a centrifuge tube, centrifuged at 200 x g for 5 min at room temperature, resuspended in pre-cooled PBS at 4˚C and centrifuged again as aforementioned, and then the supernatant was carefully aspirated. A total of 200 µl Annexin V-FITC binding solution (contained within the kit) was added to resuspend the cells and the cell density was adjusted to 1-5x10^6 cells/ml. Overall, ~100 µl cell suspension was transferred into a 5‑ml flow tube following staining with 5 µl Annexin V-fluorescein isothiocyanate (FITC) (FXP018‑100; 4A Biotech Co., Ltd., Beijing, China) for 5 min at room temperature and kept in the dark. The cells were stained with 10 µl propidium iodide (PI) in 400 µl PBS for 10 min at room temperature and kept in the dark, prior to collection and detection using flow cytometry. The results were analysed with FlowJo software (version 7.6.5; Tree Star, Inc., Ashland, OR, USA).

Statistical analysis. The experimental data were analysed by SPSS statistical analysis software (version 18.0; SPSS Inc., Chicago, IL, USA). The results were expressed as the mean ± standard deviation. The comparison of 2 groups was performed using the Student’s t-test and ≥2 groups were compared using one-way analysis of variance. Multiple comparisons between the groups were performed using a Student-Newman-Keuls test. P<0.05 was considered to indicate a statistically significant difference.

Results

**JTC-801 inhibits Hep G2 cell proliferation.** To evaluate the effect of JTC-801 on liver cancer, the Hep G2 cell line was selected and treated with various doses of JTC-801 (0.02, 0.2, 2, 20 and 200 µM) for 24, 48 and 72 h to detect cell viability. As presented in Fig. 1, higher concentrations of JTC-801 (≥20 µM) led to a statistically significant decrease in the proliferation of the Hep G2 cells compared with the control (P<0.05). In the following experiment, 20 µM JTC-801 was used. Furthermore, the OD value decreased significantly (P<0.05; Fig. 1) at 72 h. This suggests that JTC-801 may effectively inhibit liver cancer Hep G2 cell proliferation.

**JTC-801 inhibits Hep G2 cell invasion and migration.** The effects of JTC-801 on the invasion and migration ability of the Hep G2 cells were investigated using a Transwell assay (Fig. 2A). In the invasion experiment, the number of crystal violet-stained cells was decreased following the JTC-801 treatment. Similarly, in the migration experiment, cell migration was also decreased. These results suggest that JTC-801 may inhibit the metastasis of Hep G2 cells. There were fewer invaded cells in the JTC-801-treated group than in the control group (92±2 vs. 131±4 cells, respectively), indicating a negative effect of JTC-801 on the invasion ability of the Hep G2 cells (P<0.05; Fig. 2B). The capacity for migration was also
inhibited in the JTC-801-treated cells as compared with that in the negative control group (117±3 vs. 153±5 cells, respectively), and the results were significantly different (P<0.05; Fig. 2C).

JTC-801 promotes Hep G2 cell apoptosis. The impact of JTC-801 on Hep G2 cell apoptosis was determined by an Annexin V-FITC and PI double-staining assay. The apoptosis rate in the JTC-801-treated group was 14.86%, while the apoptosis rate in the control group was 7.05%. The statistical analysis suggested that the apoptosis rate in the JTC-801-treated group was significantly increased compared with that in the control group (P<0.05; Fig. 3B). Furthermore, the apoptosis regulators, including anti-apoptotic protein Bcl-2, and pro-apoptotic proteins active caspase-3 and Bax, were analysed by western blotting (Fig. 3C). In concordance with the flow cytometry analysis, the western blotting revealed that the expression of the anti-apoptotic protein Bcl-2 was decreased, and the expression of the pro-apoptotic proteins active caspase-3 and Bax was increased in the JTC-801-treated group compared with that in the control group (P<0.05; Fig. 3D). These results indicate that JTC-801 promotes Hep G2 cell apoptosis.

JTC-801 suppresses the PI3K/AKT pathway in Hep G2 cells. The PI3K/AKT signalling pathway is important in tumour progression. Proteins mTOR, p70S6K and cyclin D1 were selected as indicators to evaluate the activity of the PI3K/AKT signalling pathway following the JTC-801 treatment. The results of the western blotting revealed that the phosphorylation levels of AKT and mTOR were significantly decreased in the JTC-801-treated Hep G2 cells, as were the expression levels of p70S6K and cyclin D1 (P<0.05; Fig. 4). These results suggest that JTC-801-induced Hep G2 cell growth inhibition functions via the PI3K/AKT pathway.

Discussion

In the present study, the inhibition of hepatoblastoma Hep G2 cell proliferation, invasion and migration by the JTC-801 antagonist was demonstrated, along with the promotion of cell apoptosis. Finally, the effects of JTC-801 on Hep G2 cells were indicated to be associated with the inhibition of PI3K/AKT signalling.

JTC-801 is an opioid analgesic drug used in scientific research. However, the antitumour activity of JTC-801 is not dependent on its known analgesic function (14). JTC-801 contributes to this process by inducing a unique pH-dependent form of regulated cell death known as alkalipoptosis (14,15). In U20S osteosarcoma cells, JTC-801 inhibited cell growth by promoting apoptosis via the PI3K/AKT signalling pathway (13). Furthermore, JTC-801 demonstrated its anticancer effect in the ovarian cancer SKOV3 cell line, particularly on cell growth and metastasis (16). Song et al (12) reported that JTC-801 may be used for the treatment of pancreatic cancer. Similarly, the present study revealed the inhibitory effect of the drug on Hep G2 hepatoblastoma cells. The anticancer effect of JTC-801 only occurred at a concentration ≥20 µM. The high concentration of JTC-801 (>100 µM) acts on a wide range of receptors, including GPCRs, ion channel receptors and nociception opioid peptide receptors (17). Therefore, the mechanism by
which JTC-801 functions through receptors requires further experimental validation.

The results of the western blot analysis indicated that JTC-801 may affect the expression of apoptosis-related proteins in liver cancer. Apoptosis is an important antitumour pathway and several antitumour drugs serve an important role in cancer by inducing apoptosis (18,19). The activation of apoptosis is regulated by multiple signalling pathways, of which the PI3K/AKT pathway is one of the most important (20). The PI3K/AKT pathway regulates a number of malignant phenotypes, including anti-apoptotic, cell growth and proliferation phenotypes (21,22). In this pathway, AKT activation inhibits cell cycle arrest and angiogenesis, and promotes tumour invasion and metastasis via phosphorylation of the protein kinase mTOR (23,24). Furthermore, P70S6K, which is closely associated with cell proliferation, is located downstream of the PI3K/AKT/mTOR pathway (25). Cyclin D1 has also been reported to be a direct downstream target of the PI3K/AKT signalling pathway (26). The expression of cyclin D1 is increased in different types of tumour tissues. The present western blot results demonstrated that the phosphorylation levels of AKT and mTOR were decreased significantly in JTC-801-treated Hep G2 cells. Similarly, the expression levels of p70S6K and cyclin D1 were reduced following JTC-801 treatment. These results suggest that JTC-801 may have an antitumour effect via the regulation of apoptosis through the PI3K/AKT pathway.

In summary, the present study demonstrated that JTC-801 inhibited cell proliferation, invasion and migration, and promoted cell apoptosis through the PI3K/AKT signalling pathway in hepatoblastoma Hep G2 cells. This study provides a basis for further clinical research on JTC-801 in liver cancer treatment.

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Availability of data and materials

The datasets analysed during the present study are available from the corresponding author on reasonable request.

Authors' contributions

BFZ designed the study and was involved in data collection. TH analysed and interpreted the experimental data. Both authors prepared figures, wrote the initial manuscript and approved the final manuscript.

Ethics approval and consent to participate

Not applicable.

Patient consent for publication

Not applicable.

Competing interests

The authors declare that they have no competing interests.

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