Introduction

Lung cancer (LC) is the most common malignant tumor in the world, which causes over a million deaths per year (1,2). In recent years, the morbidity and mortality of LC have increased sharply (3). At present, surgery and chemotherapy are the primary treatment of LC, but there are still problems with drug resistance and easy recurrence (4-6). Over the past 10 years, the 5-year survival rate for patients with advanced LC has not exceeded 20%. With the development of targeted molecular biology, targeted therapy has become a promising method for the treatment of LC. Therefore, the search for new targeted drugs plays a key role in improving the survival rate of patients with advanced LC.

Ligustrazine (LSZ), an alkaloid, is an active ingredient extracted from the Chinese herbal medicine Ligusticum chuanxiong Hort (7). Numerous studies have suggested that LSZ exhibits a variety of biological activities such as anti-depression (8), anti-inflammatory (9), cardioprotective (10) anti-cancer (11,12), and neuroprotective (13). In recent years, pieces of evidence also indicated that LSZ exhibited potent anticancer effects in melanoma (14) and breast cancer (15). Zou et al. found that LSZ-based analogs...
repressed proliferation, motility, and heteroadhesion of HCT-116 cells by inactivating Wnt/β-catenin pathway via inhibiting the Akt and GSK-3β phosphorylation (16). In lung disease, LSZ can alleviate acute lung injury and induce apoptosis in LC cells (17,18).

Phosphatase and tensin homolog (PTEN) is a newly discovered tumor suppressor gene, which is the most popular tumor suppressor gene after p53 and plays a vital role in various cancer cells (19). Studies have shown that PTEN has little effect on the protein product of the gene, but it has a strong catalytic effect on the phosphorylation of the substrate (20). In human malignancies such as breast cancer (21) and liver cancer (22), PTEN exhibits low expression and dysfunction. Similarly, in LC, the inactivation of PTEN induces the movement of non-small cell LC cells (23). Besides, the level of PTEN can be inhibited by MIR, thereby preventing the progression of non-small cell LC (24).

The Wnt/β-catenin signaling pathway is a canonical Wnt signaling pathway that plays a vital role in the progression of LC (25,26). It affects various physiological processes in cells such as cell cycle, proliferation, invasion, migration, apoptosis, and angiogenesis (27-29). Abnormal expression of molecules in Wnt/β-catenin signaling plays a vital role in the development of LC, such as glycogen synthase kinase-3β and β-catenin (30). Recently, Curcumin, Garcinol, and Qiyusanlong have inhibition of LC by regulating the Wnt pathway (31-33). However, the roles and molecular mechanism of LSZ in LC is still unclear. In the study, we investigated the effects of different concentrations of LSZ on the expression of PTEN protein in LC cells. The results show that LSZ can regulate the malignant biological behavior of LC cells by regulating Wnt/β-catenin signaling pathway.

**Methods**

**Samples collection**

From March 2015 to May 2016, 30 patients with LC and 40 patients with the adjacent normal group were admitted to our hospital. All patients did not receive chemotherapy or radiation before surgery. Two pathologists confirmed them after pathological staging. The tumor tissue is removed and quickly placed in the RNA preservation solution.

**Cell lines and reagents**

Human lung fibroblasts (MRC-5) and human LC cells (H1650, A549, H1299, and PC-9) were purchased from the Cell Research Institute of Wuhan University (Wuhan, China) and cultured in RPMI 1640 medium supplemental with 5% fetal bovine serum at a constant temperature. (Hyclone Corporation) Transwell chamber was bought from Millipore, Inc., and matrigel was bought from Bio-Rad (Bio-Rad, Madrid, Spain). Lentivirus kit and retrovirus LV3-PTEN were bought from Shanghai Gemma Biology Co., Ltd.

**Quantitative real-time reverse transcription PCR (RT-qPCR)**

Total RNAs are extracted from cells using RNAprep pure Cell / Bacteria Kit. (Tiangen Biotech. Beijing, China) RT-qPCR was conducted by FastKing One-Step RT-qPCR Kit (Probe) (Tiangen Biotech. Beijing, China) and ABI 7500 Real-Time PCR System (Applied Biosystems, USA). The primers of GAPDH were 5’-GTACAATGCACGCCTAGCCG-3’ (sense) and 5’-CATTCGACTCGACTAATGCC-3’ (antisense); PTEN, 5’-GCATTTACGCAACTCGACG-3’ (sense) and 5’-CCTGCTACTAGACATAGGCT -3’ (antisense). GAPDH was used as an internal reference. Fold the equation 2^{ΔΔCt} calculated changes.

**Cytotoxic effect**

Cell viability was measured by CCK-8 assay according to the manufacturer’s protocol. MRC-5 cells were treated with LSZ (0, 10, 20, 40, 60, 80, 100 and 120 mg/kg) and then incubated in 10% CCK-8 at 37 °C for 3 h. Cell viability was measured, and the absorbance was determined at 570 nm with multifunctional microplate reader SpectraMax M5 (Molecular Devices, Sunnyvale, CA, USA).

**Cell infection**

H1299 cells in the logarithmic growth phase were inoculated on the 6-well culture plate until the fusion degree was 40%. Then, cells were infected with lentivirus holding LV3-NC or LV3-PTEN, respectively. The experiment was randomly divided into two groups: the LV3-NC group and the LV3-PTEN group. The cells were evenly laid into a 6-well plate and incubated in an incubator at 37 °C.

**Immunobistochemistry**

The tissue sections were deparaffinized with xylene and
rehydrated in gradient ethanol. After repairing with 1x citrate, the parts were placed in a 3% aqueous hydrogen peroxide solution and incubated for 10 minutes to remove endogenous peroxidase. Next, the sections were blocked with a blocking solution for 1 hour at room temperature and then incubated with the primary antibody (#9188, 1:125, Cell Signaling Technology, USA) at 4 °C overnight. After washing with TBST, the sections were incubated with the secondary antibody (#8114, Cell Signaling Technology, USA) for 30 minutes at room temperature, and stained with SignalStain® DAB. The images were seen under a light microscope.

Western blotting

The total protein in cells and tissues is extracted with cell lysate and cryopreserved for use. After separating by SDS-PAGE, the proteins (25 μg) were transferred to the PVDF membrane. Then, the membrane was incubated with primary antibodies PTEN (#9188, 1:1000, CST, USA), GAPDH (#5174, 1:1,000, CST, USA), β-catenin (#8480, 1:1,000, CST, USA), Wnt (#2391, 1:1,000, CST, USA) and GSK-3β (#12456, 1:1,000, CST, USA) overnight at 4 °C. The next day, after washing with TBST, the membrane was incubated with a second antibody (#7074, CST, USA) for 1 hour. The ECL developer was developed and photographed.

Transwell assay

H1299 cells were diluted to 1×10⁶ cells/mL. Two hundred and fifty μL medium (fetal bovine serum) and 25 μL cell suspension were added to the upper chamber covered with Matrigel and cultured in 37 °C incubators for 24 hours. The upper room was wiped with aseptic cotton swabs, stained with crystal violet for 10 minutes, washed with PBS for 5 minutes, then observed under a microscope and photographed.

Cloning formation assays

Cloning formation assays were used to detect the ability of clone formation of H1299 cells. Briefly, LSZ-treated cells were resuspended in a single cell suspension and cultured in RPMI 1640 medium supplemental with 5% FBS at a constant temperature. The colonies were fixed with paraformaldehyde and stained with crystal violet.

LC xenografts

Male BALB/c nude mice (4–6 weeks old) were housed under specific pathogen-free conditions. All animal experiments were performed following the NIH Guide for the Care and Use of Laboratory Animals and were approved by The First Affiliated Hospital of Hebei North University. H1299 cells (2×10⁶/mL) were injected subcutaneously into the right-back flanks of the mice. After successful modeling, the mice were randomly divided into 2 groups (n=10): LSZ group; Mice were injected intraperitoneally (i.p.) with 70 mg/kg LSZ; Control group, mice were subjected to the same volume of physiological saline daily. After treatment for 6 weeks, the mice were sacrificed. Tumors were taken out, and volume and weight were measured.

Statistical analysis

SPSS 13.0 software was used for statistical analysis. Data were expressed as mean ± standard deviation, and one-way ANOVA analyzed the data. The variance inequality was treated with the Kruskal-Wallis method. The difference was considered statistically significant at P<0.05.

Results

Low expression of PTEN and the promoting effect of LSZ on PTEN in LC

The results of the immunohistochemical analysis showed that the expression of PTEN in LC tissue was significantly lower than that in normal tissues, indicating that PTEN was low expressed in LC tissues (Figure 1A). RT-qPCR and Western blotting showed that PTEN in LC cell lines (H1650, A549, H1299, and PC-9) was down-regulated compared with lung fibroblasts (MRC-5). Among them, the change of PTEN in H1299 cells was the most significant. Therefore, H1299 cells were selected as the research object in the follow-up experiment (Figure 1B,C). Besides, cytotoxicity experiments showed that when the dose was lower than 60 mg/kg, LSZ had no apparent toxic effect on MRC-5. However, LSZ inhibited the growth of MRC-5 cells in varying degrees when it was higher than 80 mg/kg. RT-PCR and Western blotting were implemented again to investigate the effect of LSZ on PTEN in H1299 cells (Figure 1D). The results showed that LSZ promoted the expression of PTEN in a dose-dependent manner in a certain dose range. When the dose exceeded 60 mg/kg, the level of PTEN began to
decrease (Figure 1E,F). These results show that PTEN was low expressed in LC, while LSZ promoted the expression of PTEN in LC.

**LSZ inhibits invasion and proliferation of LC cells**

Transwell and clone formation assay was implemented to investigate the effect of LSZ on the invasion and proliferation of H1299 cells. As shown in Figure 2, LSZ treatment significantly inhibited the invasion and proliferation of H1299 cells compared with the control group. These results suggest that LSZ inhibits the growth and motility of LC cells to a certain extent.

**Knockdown PTEN counteracts the inhibitory effect of LSZ on the invasion and proliferation of LC cells**

Cells, knockdown experiments, Transwell, and clone formation assays were performed to investigate the effects of PTEN and LSZ on invasion and proliferation of H1299. As shown in Figure 3A,B, the level of PTEN was significantly reduced in the LV3-PTEN group compared to the LV3-NC group. After infecting H1299 cells with a lentivirus holding LV3-PTEN, a PTEN-deficient H1299 cell line was selected. Normal H1299 cells and PTEN-deficient H1299 cells were treated with LSZ, and invasive ability and proliferative ability were measured by Transwell and clone formation assays, respectively. The results showed
that compared with normal H1299 cells, cell invasion, and proliferation of PTEN-deficient H1299 cells were significantly inhibited (Figure 3C, D). These results show that knockdown PTEN counteracts the inhibitory effect of LSZ on the invasion and proliferation of LC cells.

**LSZ inhibits tumor formation by increasing PTEN levels and blocking the Wnt/β-catenin pathway in vivo**

Tumor growth in nude mice after 6 weeks is shown in Figure 4A. The results showed that the tumors in the LSZ group were significantly smaller than the control group. Tumor weight and volume were shown in Figures 4B, C; the weight and volume of the tumor were significantly reduced after LSZ treatment. In terms of mechanism, Western blotting showed that LSZ treatment significantly reduced the levels of Wnt and β-catenin, while increased the level of GSK-3β, indicating that LSZ inhibited the activation of Wnt/β-catenin pathway by increasing the level of GSK-3β. Also, western blotting and immunohistochemical analysis showed that the expression of PTEN in tumor tissues treated with LSZ was significantly higher than that in the control group without LSZ treatment. These results show that LSZ inhibits tumor formation by increasing PTEN levels and blocking the Wnt/β-catenin pathway in vivo.

**Discussion**

LC is one of the malignant tumors with the highest morbidity and mortality. The high metastasis of LC cells leads to a meager early diagnosis rate and easy to deteriorate. The recurrence rate of advanced LC is high, while the survival rate is low with a poor prognosis (34). At present, the early diagnosis, prognosis, and treatment of patients with LC are a hot clinical study. Abnormal PTEN mutations can lead to some tumor gene mutations in the occurrence and development of tumors, resulting in the inactivation of tumor suppressor genes (35). Therefore, we hope to study the relationship between PTEN diagnosis and treatment of LC.

PTEN is a recently discovered tumor suppressor gene, found on chromosome 10q23.3. The transcript is about 515 kb (36). The phosphatase domain of PTEN exists in tyrosine and bispecific phosphatase (37). PTEN
may play an essential role in tumor development and progression through dephosphorylation (38). Li et al. found that microRNA-374b promotes LC cell motility and proliferation by targeting PTEN and downregulating PTEN expression via modulating the PI3K/AKT pathway (39). Liu et al. found that PTEN was significantly decreased in NSCLC cell lines. Knockdown PTEN by shRNA significantly increased cell viability, while overexpression of PTEN significantly inhibited the growth and apoptosis of NSCLC cells (40). In this study, it was found that the levels of PTEN in tumor tissues and LC cells of patients with LC were decreased, while LSZ treatment could significantly increase the level of PTEN, but also inhibit the invasion, proliferation and tumor growth of LC cells. These results suggest that PTEN plays a crucial role in the pathogenesis of LC.

LSZ is an effective alkaloid isolated and purified from *Chuanxiong*. It has a wide range of anti-platelet and depolymerized platelet coagulation function and has a good alleviation effect on tissue and organ ischemia and hypoxia (41). Feng et al. found that LSZ can inhibit autophagy and apoptosis of liver cancer cells, thereby affecting the metastasis and spread of liver cancer cells and inhibiting the development of liver cancer (42). Li and

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**Figure 3** Knockdown PTEN counteracts the inhibitory effect of LSZ on the invasion and proliferation of LC cells. (A) The protein level of PTEN was detected by western blotting in H1299 cells infected with lentivirus holding LV3-NC or LV3-PTEN. (**, P<0.01 vs. LV3-NC group). (B) The mRNA level of PTEN was detected by RT-qPCR in H1299 cells infected with lentivirus holding LV3-NC or LV3-PTEN. (**, P<0.01 vs. LV3-NC group). (C,D) Normal H1299 cells and PTEN-deficient H1299 cells were treated with LSZ. (B) Invasive ability was measured by the Transwell assay. (C) Proliferative ability was measured by the clone formation assay. Magnification ×400 (**, P<0.01 vs. control group). LSZ, ligustrazine; LC, lung cancer.
other studies have shown that LSZ can effectively alleviate the proliferation of glioma in glioma (43). It is reported that LSZ exhibited therapeutic effects on sepsis-induced acute lung injury in rats (17). Besides, LSZ induces cycle arrest and apoptosis of LC cells (18). Jia et al. also found that LSZ suppressed proliferation, migration, and capillary tube formation of HMEC-1 cells by blocking the BMP/Smad/Id-1 pathway \textit{in vitro} and \textit{in vivo} (44). Consistent with these results, this study found that LSZ has a certain promoting effect on the expression of PTEN in LC cells. Increased PTEN further inhibits invasion and proliferation of LC cells. Also, animal experiments have shown that LSZ inhibits tumor formation by down-regulating Wnt/β-catenin pathway in \textit{vivo}.

Wnt signaling pathway has irreplaceable physiological functions \textit{in vivo}. Wnt signaling pathway can not only regulate the formation of the brain and nervous system but also closely related to the self-renewal and differentiation regulation of various stem cells (45). Wnt signaling can modulate the biological behavior of malignant cells. Studies have shown that the Wnt signaling pathway in malignant tumor cells such as rectal cancer, colon cancer, and ovarian cancer is activated, which has certain reference value for the diagnosis and prognosis of cancer (46-48). Wang et al. found that curcumin inhibited the proliferation of A549 cells by inhibiting the Wnt pathway and reduced oxidative stress damage (31). Huang et al. found that Garcinol inhibited the phenotype of human non-small cell LC stem cells by inhibiting the Wnt/β-catenin/STAT3 axis signaling pathway (32). Similarly, this study found that LSZ significantly inhibited the activation of the Wnt/β-catenin pathway in tumor tissues by up-regulating the level of GSK-3β, thereby inhibiting tumor formation in mice.

In conclusion, current studies have explored the effects of LSZ and PTEN on the formation of LC. The results showed that the level of PTEN was significantly down-regulated in LC tissues or cells, while LSZ treatment promoted the expression of PTEN in LC cells in a dose-dependent manner. The increase of PTEN further inhibited the proliferation and invasion of LC cells. Animal experiments show that LSZ treatment can inhibit the formation of tumors in mice by up-regulating the
levels of PTEN and GSK-3β and inactivating Wnt/β-catenin pathway. These results suggest that LSZ and PTEN are of great significance in the treatment and prognosis of LC.

Acknowledgments

Funding: None.

Footnote

Conflicts of Interest: The authors have no conflicts of interest to declare.

Ethical Statement: The authors are accountable for all aspects of the work in ensuring that questions related to the accuracy or integrity of any part of the work are appropriately investigated and resolved. All animal experiments were performed following the NIH Guide for the Care and Use of Laboratory Animals and were approved by The First Affiliated Hospital of Hebei North University.

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