Effects of tanshinones mediated by forkhead box O3a transcription factor on the proliferation and apoptosis of lung cancer cells

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Abstract. According to global cancer statistics in 2012, lung cancer (LC) was the most frequently diagnosed cancer and the leading cause of cancer-associated mortality among males worldwide. Owing to the limited therapeutic approaches available, novel methods for treating LC are required. Tanshinones (Ts) have previously been proved to be effective in treating cardiovascular disease, inflammatory disease and cancer, and have been reported to regulate cell proliferation and apoptosis of LC. The underlying molecular mechanism of action of Ts remains unclear. Furthermore, forkhead box O3a (FoxO3a) has been reported to be a critical gene in cell apoptosis. Therefore, the A549 lung cancer cell line was transfected with FoxO3a small interfering RNA (siRNA) or scrambled siRNA, and the cells which exhibited the most successful transfection efficacy were selected for further investigation into the underlying molecular mechanism of the influence of Ts on FoxO3a mRNA and protein were restored following treatment with Ts in a dose-dependent manner, alongside caspase-3 activation. On the basis of these results, we hypothesize that Ts regulate LC cell proliferation and apoptosis by triggering an apoptotic cascade through the FoxO3a/caspase-3 signaling pathway.

Introduction

According to global cancer statistics in 2012, lung cancer (LC) was the leading cause of cancer-related mortality in developed countries (1). Furthermore, in less-developed countries including China and India, LC rates are predicted to continue increasing as a result of increasing endemic use of tobacco (1). It is well-known that the progression of LC is the result of sequential genetic and epigenetic changes including the dysregulation of oncogenes, tumor suppressor genes and growth factors (2). Currently, the general approach to the treatment of LC (non-small cell LC and small cell LC) is based on surgical resection, chemotherapy and radiation therapy according to the stage (3). However, it is well-known that the intrinsic and potent cytotoxicity to normal cells of a number of anticancer agents imposes a restriction on their prolonged use and their therapeutic effectiveness (4). As a result, exploring novel therapeutic methods is of urgent priority.

Chinese herbs have been considered as novel ways to treat diseases (5). With regard to cancer, a number of previous studies have demonstrated that extractions of specific Chinese herbs exhibit therapeutic effects on certain types of cancer (6-8). The exact underlying molecular mechanism of Chinese medicine-based treatments of cancer remain elusive.

Tanshinones (Ts), including tanshinone I, tanshinone IIA, cryptotanshinone and dihydrotanshinone I, are a class of lipophilic diterpene compounds extracted from Salvia miltiorrhiza Bunge, a plant root used in traditional Chinese medicine (9). Numerous studies have demonstrated that Ts exhibit potential properties of anti-inflammation (10), anticancer (11-13) and cardio-cerebrovascular protection (14). Regarding LC, previous studies have suggested that Ts are involved in the process of inhibiting cell proliferation and inducing apoptosis (13,15,16). However, the underlying molecular mechanism of the effect of Ts on LC remains unknown.

Previous studies have demonstrated that FoxO3a, a member of the forkhead box (Fox) gene family of transcription factors, serves a role in the apoptotic cascade (17,18), which suggests that targeting the AMP-activated protein kinase-FoxO3a axis...
is a potential therapeutic approach for cancer treatment (19). Previous study has demonstrated that silencing FoxO3a prevents control of the apoptotic cascade due to the inhibition of mitochondrial membrane depolarization. In contrast, the presence of FoxO3a is the prerequisite for cleaved caspase-3 expression (17). On the basis of these previous studies, we hypothesize that FoxO3a mediates the activity of Ts, regulating LC cell proliferation and apoptosis through caspase-3 activation.

Materials and methods

Cell lines and cell culture. The human lung cancer cell line A549 was obtained from PeproTech, Inc. (Rocky Hill, NJ, USA). The cells were routinely cultured in Dulbecco’s modified Eagle’s medium (DMEM; Gibco; Thermo Fisher Scientific, Inc., Waltham, MA, USA) supplemented with 10% fetal bovine serum (FBS; Thermo Fisher Scientific, Inc.) at 37°C in a humidified cell incubator containing 5% CO₂.

Plasmid construction. The plasmid (pMSCV-puro) containing FoxO3a small interfering RNA (siRNA) was constructed by Guangzhou Ribobio Co., Ltd. (Guangzhou, China). The sequences were as follows: siRNA1, 5’-UUCUCCGAAGCUGACUACGUUGAGAATT-3’; siRNA2, 5’-AUCCUGGGUCCACUCUCACTT-3’ and 5’-GGUGGACUGUGGACCCGAGUGGUTT-3’; siRNA3, 5’-GGAGACUGUAGCUUCGCAATT-3’ and 5’-UUGCGAGACUCACGUUCCCGG-3’. Scrambled siRNA (ssiRNA) was purchased from Invitrogen; Thermo Fisher Scientific, Inc., and with sequences of 5’-CGUUCACCGCAAUUCAUATT-3’ and 5’-UAGAAUUGUGUGCGUAGACCGA-3’.

Transfection. For transfection, 5x10⁶ A549 cells were seeded in 6-well culture plates until reaching 70% confluence. Subsequently, transfection of cells with recombinant FoxO3a-siRNA/ssiRNA plasmids was performed using Lipofectamine 2000 (Thermo Fisher Scientific, Inc.), according to the manufacturer’s protocol. At 8-12 h after transfection, the morphology of transfected cells was observed using a fluorescence microscope (Olympus Corporation, Tokyo, Japan) at x200 magnification, and the most successfully transfected experimental and control (CON) cells were selected to for treatment with Ts in the subsequent experiments.

Preparation of Ts and experimental groups. Ts (Xian Yuesun Biological Technology, Xi’an, Shaanxi, China) was dissolved at 25 mmol/l in absolute ethyl alcohol and stored at 4°C until use. The transfected A549 cells were divided into four groups and treated with Ts as follows: 5 µmol/l Ts (group I), 10 µmol/l Ts (group II), 20 µmol/l Ts (group III), 30 µmol/l Ts (group IV) and the negative control of ssiRNA-transfected cells without Ts administration (CON group).

Measurement of cell proliferation. Cell proliferation was determined using an MTT assay (Shanghai Yeasen Biotechnology Co., Ltd., Shanghai, China), according to the manufacturer's protocol. Transfected A549 cells were seeded onto 96-well plates (5x10⁴ cells/well). The cell proliferation was measured every 24 h for 3 days. The absorbance of cells was screened at 570 nm using a microplate reader (SM600; Shanghai Utrao Medical Instrument Co., Ltd., Shanghai, China).

Analysis of cell apoptosis. For analysis of apoptosis, the proportions of apoptotic cells were determined using an Annexin V-Fluorescein Isothiocyanate (FITC)/Propidium Iodide (PI) Apoptosis Detection kit (Beijing Solarbio Science and Technology Co., Ltd., Beijing, China), according to the manufacturer’s protocol. Briefly, the transfected A549 cells of each group were cultured for 72 h, collected by trypsinization (Beijing Dingguo Changseng Biotechnology Co., Ltd., Beijing, China) and washed with PBS. The cells were subsequently resuspended in 1X binding buffer at a concentration of 1x10⁶ cells/ml. Subsequently, 5 µl Annexin V-FITC was added to the cell suspension, prior to incubation at 4°C for 15 min in darkness. Subsequently, 10 µl PI was added to the samples, mixed gently and incubated at 4°C for 5 min in darkness. Finally, the cells were screened using flow cytometry (Beckman Coulter, Inc., Brea, CA, USA).

Reverse transcription-polymerase chain reaction (RT-PCR). Total RNA from transfected A549 cells of each experimental group was extracted using TRIzol reagent (Invitrogen; Thermo Fisher Scientific, Inc.). The RT and PCR primers for FoxO3a and GAPDH (internal control) were obtained from KareBay Biochem, Inc. (Monmouth Junction, NJ, USA). The PCR primers for FoxO3a were 5’-CGTGGTAAAGGTGTCC-3’ (forward) and 5’-CAAGGCTCTACACTCGGAC-3’ (reverse); the primers for GAPDH were 5’-CTGTGGAAAGGACTCATGCACC-3’ (forward) and 5’-AGGGATGATGGTTCGAGG-3’ (reverse). The PrimeScript RT Reagent kit (Takara Biotechnology Co., Ltd., Dalian, China) was used to synthesize the first-strand cDNA. The cycling program was as follows: Initiation at 37°C for 10 min, followed by 85°C for 5 sec, then 4°C for 10 min. PCR was performed using SYBR Premix Ex Taq (Guangzhou Ribobio Co., Ltd.). Quantitative RT-PCR was performed using the PCR 7900HT Real-Time PCR System (Applied Biosystems; Thermo Fisher Scientific, Inc.). The cycling program was as follows: Initiation at 95°C for 30 sec, followed by 40 cycles at 95°C for 5 sec and at 60°C for 34 sec. The relative expression level of each circRNA was analyzed using the ΔΔCq method (20).

Western blot analysis. The cells of each group were collected, centrifuged (500 x g at 4°C for 15 min) and transferred into clean test tubes. A 400 µl volume of lysis buffer (Beyotime Institute of Biotechnology, Haimen, China) was added to each tube to lyse the cells at -10°C for 30 min, then centrifuged at 500 x g at 4°C for 10 min. The formula of the lysis buffer was: Collagenase type IV (1 mg/ml), DNase type I (100 µg/ml), CaCl₂ (2 mmol/l) and MgCl₂ (2 mmol/l). The protein concentration of the lysates was examined using the Bradford method (21). Lysates (200 µl) were then boiled, electrophoresed (4% SDS) and transferred onto nitrocellulose membranes (CST Biological Reagents Co., Ltd., Shanghai, China). Non-fat milk solution (10%) in Tris-buffered saline solution with Tween-20 (0.5 ml/l) was used to block the membranes, which were incubated at 25°C for 2 h. Subsequently, 10 µl mouse anti-human FoxO3a monoclonal antibody (cat. no. 05-1075-25UG; 1:250; EMD Millipore, Billerica, MA, USA) and mouse anti-human
caspase-3 (cat. no. ab1271; 1:1,000; Abcam, Cambridge, UK) were added and incubated at room temperature for 2 h, followed by 5 µl alkaline phosphatase-conjugated goat anti-mouse IgG secondary antibody (cat. no. 115-055-006; 1:50; Jackson ImmunoResearch Laboratories, Inc., West Grove, PA, USA). Signals were screened using an automated chemiluminescence immunoassay analyzer (Beckman Coulter, Inc.).

Statistical analysis. All statistical analysis was carried out using the SPSS software package (version 12.0; SPSS, Inc., Chicago, IL, USA) and GraphPad Prism 5.00 software (GraphPad Software, Inc., La Jolla, CA). For comparisons between two groups, Student's t-test was performed. For comparisons between multiple groups, a Student-Newman-Keuls test was performed. P<0.05 was considered to indicate a statistically significant difference.

Results

Recombinant FoxO3a siRNA and ssiRNA were successfully transfected. The transfected A549 cells expressed green fluorescence as determined using fluorescence microscopy. As presented in Fig. 1, the transfection efficiency of siRNA2 was higher than that of siRNA1 and siRNA3 (P<0.05), with similar results in comparison with those for a negative control. As a result, siRNA2-transfected cells were selected as the culture containing the most successfully transfected cells to use for further experiments.

Ts inhibit the proliferation of LC cells. To detect cell viability, the absorbance of cells in each group was screened at 24, 48 and 72 h using an MTT assay. As presented in Fig. 2, Ts inhibited the cell viability in a dose- and time-dependent manner with a maximal dose of 20 µmol/l at 72 h treatment.

Ts induce LC cell apoptosis. Following treatment of the cells of each group with Ts for 72 h, the effect of Ts was examined using the Annexin V-FITC/PI Apoptosis Detection kit. The proportion of apoptotic cells in each experimental group was increased compared with that of CON (P<0.05), and dose-dependence was exhibited by experimental groups I-III (P<0.05; Fig. 3). Similarly, no statistically significant difference was identified in experimental groups III and IV, suggesting that 20 µmol/l may be the most appropriate concentration to treat LC cells.

Ts upregulate the expression of FoxO3a mRNA. As presented in Fig. 4A, the expression of FoxO3a mRNA in experimental groups III and IV was increased compared with that of CON (P<0.05). Similarly, a dose-dependent association was identified in experimental groups I-III, but not in experimental groups III and IV.

Ts modulate FoxO3a and caspase 3. As presented in Fig. 4B and C, compared with the CON group, the expression of FoxO3a increased as the protein concentration was increased, indicating that FOXO3a was associated with the dosage of Ts in experimental groups I-III (P<0.05), which was consistent with the results of cell proliferation and apoptosis. Simultaneously, as FoxO3a protein was restored, caspase-3 was activated, suggesting that Ts were able to restore the function of FOXO3a, which governs apoptotic transduction pathways by inducing caspase-3 activity.

Discussion

FoxO3a has been recognized as a potential tumor suppressor of lung adenocarcinoma, the expression of which is associated with the aggressiveness of the cancer (22-24). Certain Chinese medicines, including berberine, have been demonstrated to contribute to the inhibition of LC cell proliferation and the induction of apoptosis by regulating the expression of FoxO3a, which serves a role in the activation of the p38a mitogen-activated protein kinase signaling pathway (25). Although the results of previous studies suggest that Ts are implicated in anticancer activities including dysregulation of cell proliferation, apoptosis, cell cycle arrest, metastasis, differentiation and angiogenesis (15,16,26-28), the underlying molecular
The mechanism remains unknown. As a result, the aim of the present study on LC cell treatment with Ts was to investigate the potential signaling pathway mediated by FoxO3a.

To address the aforementioned questions, effective transfected LC cells were established by silencing FoxO3a, whereas the CON cells transfected with FoxO3a ssiRNA retained intact FoxO3a in LC cells. In addition, construction of the siRNA nucleotide sequence was optimized with processes including the presence of a phosphate group at the 5'-terminus and the strand with a less stable 5'-end (29). The successful knockdown of FoxO3a provided a good foundation for the subsequent experiments.

Dysregulation of cell proliferation is the critical characteristic of LC, therefore four distinct concentrations of Ts (5, 10, 20 and 30 µmol/l) were used to treat the experimental groups. The absorbance of experimental groups with FoxO3a knockdown decreased in a dose- and time-dependent manner when compared with that of CON, excluding the concentration of 30 µmol/l, which indicates that Ts inhibit LC cell proliferation. Additionally, the results demonstrated that the optimal cell inhibition ratio was with 20 µmol/l at 72 h, so subsequent measurements were made at the 72 h time point.

Circumventing of cell apoptosis is another feature of LC. In accordance with a previous study of the effect of Ts on LC (15), the results of the present study illustrated that Ts induce cell apoptosis in a dose-dependent manner; however, experimental group IV did not continue the trend of dose-dependence. Cells transfected with non-specific scrambled siRNA without Ts did not exhibit markedly early and late programs of apoptosis. In contrast, experimental groups demonstrated increased proportions of apoptotic cells, identifying the unique ability of Ts to control the early and late apoptotic programs.
Furthermore, the results of the present study demonstrated that the expression levels of FoxO3a protein in cells treated with Ts were increased compared with those in CON cells, consistent with the expression of FoxO3a mRNA with RT-PCR. Previous studies have demonstrated that silencing FoxO3a depletes the induction of caspase-3 activity (30). The results of the present study support this hypothesis by restoring the expression of FoxO3a with Ts treatment, inducing apoptosis by activating a downstream molecule, caspase-3.

The results of the present study reveal that Ts may serve a direct role in regulating LC cell proliferation and apoptosis, mediated by FoxO3a activation; however, further studies are required to determine the exact signaling pathways mediated by Ts. Addressing this deficit in our knowledge may lead to novel therapeutic approaches to treat LC clinically.

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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

DJL, FY and DHZ made substantial contributions to conception and design, RBY analysis and interpretation of data. DJL agreed to be 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.

Ethics approval and consent to participate

Not applicable.

Consent for publication

Not applicable.
Competing interests

The authors declare that they have no competing interests.

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