**Abstract.** The aim of the present study was to investigate the function of long non-coding RNA taurine-upregulated gene 1 (lncRNA TUG1) in chronic obstructive pulmonary disease and further assess the underlying molecular mechanisms. Flow cytometry analysis was performed to detect cell apoptosis of human pulmonary microvascular endothelial cells (HPMECs) treated with 1% cigarette smoke extract (CSE). The activity of caspase-3 was measured using a Caspase-3 Activity assay kit and the protein expression of cleaved caspase-3, caspase-3 and Bcl-2 like 11 (BCL2L11) were measured using western blotting. Reverse transcription-quantitative PCR (RT-qPCR) was performed to measure the expression of TUG1 mRNA levels in the treated cells. The association between TUG1 and miR-9a-5p/BCL2L11 axis was predicted and verified using a dual luciferase reporter assay system. The mRNA expression of miR-9a-5p and BCL2L11, and the transfection efficiency were measured by RT-qPCR. The results showed that CSE induced cell apoptosis and increased lncRNA TUG1 expression in HPMECs. CSE significantly reduced the expression of miR-9a-5p in HPMECs compared with the control group. TUG1-short hairpin RNA relieved cell apoptosis induced by CSE by downregulating miR-9a-5p in HPMECs. The present study predicted and verified that BCL2L11 is a direct target of miR-9a-5p. The mRNA expression of BCL2L11 was increased in HPMECs following CSE treatment compared with the control group. miR-9a-5p mimic and BCL2L11-plasmid markedly increased the expression of miR-9a-5p and BCL2L11, respectively. miR-9a-5p mimic reversed the increase in cell apoptosis induced by CSE by inhibiting BCL2L11 expression in HPMECs. To conclude, the present study demonstrated that lncRNA TUG1 exerted roles in cell apoptosis induced by CSE through modulating the miR-9a-5p/BCL2L11 axis.

**Introduction**

Chronic obstructive pulmonary disease (COPD), which is a progressive degenerative lung disease, is a preventable and treatable disease characterized by incomplete reversibility and airflow limitation, including emphysema and chronic bronchitis (1-3). It is currently the fourth leading cause of death worldwide, just behind ischemic heart disease and cerebrovascular disease, and an increasing number of individuals are diagnosed with the disease (4). Therefore, new and effective treatment strategies targeting COPD are urgently needed. Patients with COPD have difficulty breathing and present with airflow obstruction, which is often accompanied by discomforts, such as coughing and sputum production (5,6). However, the pathogenesis is not yet fully understood. The occurrence of COPD is the result of the combined effects of genetic and environmental pathogenic factors. Cigarette smoke is the most common risk factor for COPD. The rates of respiratory disease, lung function impairment and mortality of smokers are significantly higher compared with non-smokers (7,8). Smoking cessation has been shown to effectively delay the progressive decline of lung function.

Endothelial cells (ECs), as the basic unit of blood vessels, and play key roles in physiological metabolism. Under harmful conditions, the regulatory mechanisms become disrupted, which may be one of the main initiating factors for various diseases (9,10). The pathogenesis of acute lung injury includes alveolitis and tissue recovery, and the vascular response is the central link in the occurrence of inflammation (11). Previous data suggested that abnormal apoptosis serves as a significant factor contributing to the destruction of pulmonary tissue in COPD (12). Both animal and human studies confirmed that endothelial apoptosis plays a key role in the pathogenesis of COPD (13,14). It was reported that human pulmonary microvascular endothelial cells (HPMECs) play important roles in...
the progression of COPD, and the present study further investigated COPD using HPMECs as a model (15).

Long non-coding RNAs (lncRNAs), a class of non-coding RNAs that are >200 nucleotides in length, participate in the structure and function of chromatin, and promote a series of biological functions (16,17). An increasing number of studies have suggested that lncRNAs are associated with a number of human diseases, such as cancer, cardiovascular diseases, gynecological diseases and inflammatory diseases (18,19). Studies have suggested that lncRNAs participate in regulating the process of COPD (20). Previously, taurine-upregulated gene 1 (TUG1), one of the most commonly examined lncRNAs, was shown to be upregulated in patients with COPD, and TUG1 knockdown reduced inflammation and airway remodeling in an animal model and reversed inflammation and collagen deposition induced by cigarette smoke extract (CSE) in vitro (21,22). However, the mechanisms and roles of TUG1 in COPD remains unclear.

It was hypothesized that TUG1 may be involved in COPD pathogenesis through regulating EC apoptosis. The present study investigated whether TUG1 played a role in cell apoptosis of HPMECs induced by CSE, and further studied its molecular mechanism in order to provide a more theoretical basis and novel strategies for the treatment of COPD.

Materials and methods

Cell culture. HPMECs were obtained from the American Type Culture Collection and cultured in DMEM (Gibco; Thermo Fisher Scientific, Inc.) containing 10% FBS (Gibco; Thermo Fisher Scientific, Inc.) and 1% penicillin/streptomycin at 5% CO₂ and 37°C.

CSE preparation. CSE preparation was performed as described previously (23). Smoke from 10 cigarettes (Chengdu Cigarette Factory) was bubbled through DMEM, and this solution was considered 100% CSE. Then the 100% CSE solution was frozen and stored at -80°C until required. CSE was diluted with DMEM medium to a final concentration of 1% for the present study.

Apoptosis assay. HPMECs were treated with 1% CSE for 24 h, and then collected by trypsinization, washed in PBS and resuspended in 1X binding buffer at a density of 1x10⁶ cells/ml. A total of 100 µl cell suspension was then transferred to a 5 ml tube, and stained with 5 µl Annexin V/FITC and PI solution, according to the manufacturer's protocol (BD Biosciences). Stained cells were analyzed by flow cytometry (FCM) using a BD FACSCalibur flow cytometer (BD Biosciences) within 1 h. All the experiments were performed at least three times. Data were analyzed using FlowJo software (version 7.2.4; FlowJo LLC).

Western blot analysis. HPMECs were treated with 1% CSE for 24 h and dissolved in RIPA buffer (Beyotime Institute of Biotechnology) after washing in cold PBS. The supernatant was centrifuged at 10,000 x g at 4°C for 15 min to obtain total protein. Protein concentration was measured using a BCA protein assay kit (Beyotime Institute of Biotechnology). Equal amounts of protein (40 µg/lane) were separated by 10% SDS-PAGE. Separated proteins were transferred to PVDF membranes (Thermo Fisher Scientific, Inc.), which were then blocked at room temperature for 1 h in PBS-Tween-20 (PBS-T) solution containing 5% non-fat milk. Subsequently, membranes were incubated with cleaved caspase-3 (cat. no. ab32042; 1:1,000; Abcam), caspase-3 (cat. no. ab32351; 1:1,000; Abcam), Bcl-2 like 11 axis (BCL2L11; cat. no. ab32158; 1:1,000; Abcam) or GAPDH (cat. no. sc-47724; 1:500; Santa Cruz Biotechnology, Inc.) antibodies at 4°C overnight. Membranes were then washed three times with PBS-T and incubated with the Goat Anti-Mouse IgG Antibody (cat. no. bc-0296G; 1:1,000; BIOSS) or Goat Anti-Rabbit IgG Antibody (cat. no. bc-0295G; 1:1,000; BIOSS) for 1 h at room temperature. ECL substrate (Thermo Fisher Scientific, Inc.) was used to visualize the protein bands according to the manufacturer's protocol. Band densities were quantified using Gel-Pro Analyzer Densitometry software (version 6.3; Media Cybernetics, Inc.). All the experiments were performed at least three times.

RNA extraction and reverse transcription-quantitative PCR (RT-qPCR). Total cellular RNA was extracted using TRIzol® reagent (Invitrogen; Thermo Fisher Scientific, Inc.) and reverse transcribed into cDNA using a cDNA Synthesis kit according to the manufacturer’s protocol (Invitrogen; Thermo Fisher Scientific, Inc.). The relative expression of genes was quantified using a Prism 7000 Real-Time PCR system with SYBR qPCR MasterMix (Vazyme Biotech Co., Ltd.) in accordance with the manufacturer's protocol. Primers were purchased from Sangon Biotech Co., Ltd., and primer sequences were as follows: GAPDH forward, 5'-CTTTTGATCATCGGAAACGTCT-3' and reverse, 5'-GATAGAGCGAGGATGTGTCT-3'; U6 forward, 5'-GCTTCCGCGACAGCATAATCTAAAT-3' and reverse, 5'-CTCCCTGCAATTTGCTGTGCT-3'; lncRNA TUG1 forward, 5'-GACGGCTCAATGACCTTCTCTC-3' and reverse, 5'-TGGCTGAATGCTTTCTTGGGT-3'; miR-9a-5p forward, 5'-GCCTCGGCTCTTGGTCTTATGAG-3' and reverse, 5'-ATCCAGTGCCAGAGGT-3'; and BCL2L11 forward, 5'-CAAACGCACCATAGAAAGG-3' and reverse, 5'-ATAAGGACAGGCACACA-3'. The following thermocycling conditions were used for qPCR: Initial denaturation for 5 min at 95°C; followed by 35 cycles of 95°C for 10 sec and 60°C for 30 sec. U6 or GAPDH were used as the internal controls. The relative mRNA expression levels of lncRNA TUG1, miR-9a-5p and BCL2L11 were calculated using the 2-ΔΔCq method (24). All samples were analyzed at least in triplicate.

miRNA target analysis and dual luciferase reporter assay. The association between miR-9a-5p and BCL2L11 was investigated using TargetScan version 7.1 (targetscan.org/vert_71). The 3'-untranslated region products of BCL2L11 containing the target sequence of miR-9a-5p were obtained by RT-qPCR and fused into a pmirGLO vector (Promega Corporation) to construct the BCL2L11-wild-type (BCL2L11-WT) and BCL2L11-mutated-type (BCL2L11-MUT) reporter vectors. The results showed that BCL2L11 was a potential target of miR-9a-5p. 293 cells (ATCC® CRL-1573; ATCC), that were cultured >24 h, were transfected with BCL2L11-WT or
BCL2L11-MUT and miR-9a-5p mimic or mimic control using Lipofectamine® 2000 reagent (Invitrogen; Thermo Fisher Scientific, Inc.) for 48 h. Luciferase activity was measured using a dual luciferase reporter assay system (Promega Corporation) according to the manufacturer’s instructions. Renilla luciferase activity was used as a control. All the experiments were performed at least three times.

Cell transfection. Normal HPMECs were inoculated in 6-well plates overnight and transfected with control-small interfering RNA (siRNA), TUG1-siRNA, inhibitor control, miR-9a-5p inhibitor, TUG1-siRNA + inhibitor control, miR-9a-5p inhibitor, mimic control, miR-9a-5p mimic, control-plasmid, BCL2L11-plasmid, miR-9a-5p mimic + control-plasmid or miR-9a-5p mimic + BCL2L11-plasmid (Shanghai GenePharma Co., Ltd.) using Lipofectamine® 2000 reagent (Invitrogen; Thermo Fisher Scientific, Inc.). A total of 48 h after transfection, cells were collected, and the transfection efficiency was assessed using RT-qPCR. As shown in Fig. 2D and E, TUG1-siRNA and miR-9a-5p inhibitor significantly reduced the expression of lncRNA TUG1 in HPMECs.

Caspase-3 activity assay. The activities of caspase-3 were measured using a colorimetric assay kit (cat. no. C1115; Beyotime Institute of Biotechnology) in accordance with the manufacturer’s instructions.

Statistical analysis. Data are presented as the mean ± standard deviation of at least three independent tests. Statistical comparisons between groups were assessed using a Student’s t-test or a one-way ANOVA followed by a Tukey’s post hoc test. *P<0.05 was considered to indicate a statistically significant difference.

Results

CSE influences HPMEC growth. Cell apoptosis of HPMECs treated with 1% CSE for 24 h was detected by FCM. The results showed that CSE significantly induced cell apoptosis in HPMECs (Fig. 1A and B). Compared with the control group, 1% CSE markedly increased caspase-3 activity, cleaved caspase-3 protein expression and the cleaved-caspase-3/caspase-3 ratio (Fig. 1C-E). Moreover, lncRNA TUG1 expression in HPMECs was significantly increased with 1% CSE treatment compared with the control group (Fig. 1F). The results confirmed that lncRNA TUG1 was significantly upregulated in 1% CSE induced HPMECs.

TUG1 negatively regulates miR-9a-5p expression in HPMECs. A previous study showed the presence of a binding site between TUG1 and miR-9a-5p (25) (Fig. 2A). In the present study, the targeted relationship between TUG1 and miR-9a-5p was verified using a dual luciferase reporter assay (Fig. 2B). Moreover, miR-9a-5p expression in HPMECs was significantly decreased following the 1% CSE treatment compared with the control group (Fig. 2C). To further assess the regulatory relationship between TUG1 and miR-9a-5p, HPMECs were transfected with different plasmids, and the transfection efficiency was determined using RT-qPCR. As shown in Fig. 2D and E, TUG1-siRNA and miR-9a-5p inhibitor significantly reduced the expression levels of miR-9a-5p (Fig. 2F).
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Figure 2. TUG1 negatively regulates miR-9a-5p in HPMECs. (A) The association between TUG1 and miR-9a-5p was predicted using bioinformatics software. (B) The association between TUG1 and miR-9a-5p was verified using a dual luciferase reporter assay. (C) Levels of miR-9a-5p in HPMECs following CSE treatment were detected using RT-qPCR. (D-F) RT-qPCR was used to quantify the mRNA expression levels of TUG1 and miR-9a-5p in transfected HPMECs. Experiments were repeated three times. A one-way ANOVA with a post-hoc Tukey's test or an unpaired Student's t-test were used for analysis. *P<0.01 vs. mimic control; **P<0.01 vs. Control; ***P<0.01 vs. Control-siRNA; ****P<0.01 vs. TUG1-siRNA; ####P<0.01 vs. TUG1-siRNA+inhibitor control. TUG1, taurine-upregulated gene 1; miR, microRNA; CSE, cigarette smoke extract; HPMEC, human pulmonary microvascular endothelial cells; RT-qPCR, reverse transcription-quantitative PCR.

of TUG1 and miR-9a-5p, respectively. Compared with the control-siRNA group, TUG1-siRNA significantly increased miR-9a-5p expression in HPMECs, which was reversed by co-transfection with miR-9a-5p inhibitor (Fig. 2F). The data indicated that TUG1 negatively regulated miR-9a-5p expression in HPMECs. TUG1 inhibits cell apoptosis induced by CSE in HPMECs by regulating miR-9a-5p. As shown in Fig. 3A-F, 1% CSE significantly induced cell apoptosis, increased caspase-3 activity, enhanced cleaved caspase-3 protein expression, increased the cleaved caspase-3/caspase-3 ratio and downregulated miR-9a-5p expression in HPMECs. Moreover, compared with
the CSE + control-siRNA group, TUG1-siRNA significantly reduced cell apoptosis, caspase-3 activity, cleaved caspase-3 protein expression and the cleaved caspase-3/caspase-3 ratio, and upregulated miR-9a-5p expression in HPMECs. All these changes were reversed by miR-9a-5p inhibitor co-transfection. The above results suggested that TUG1 inhibited cell apoptosis induced by CSE in HPMECs by regulating miR-9a-5p.

**BCL2L11 is a direct target gene of miR-9a-5p.** TargetScan analysis showed the presence of a binding site between BCL2L11 and miR-9a-5p (Fig. 4A), which was verified by a dual luciferase reporter assay (Fig. 4B). Moreover, the mRNA expression of BCL2L11 in HPMECs was significantly increased with 1% CSE treatment compared with the control group (Fig. 4C). Further results showed that miR-9a-5p expression in the miR-9a-5p mimic group was higher compared with the mimic control group (Fig. 4D). BCL2L11-plasmid significantly increased BCL2L11 mRNA expression in HPMECs (Fig. 4E). Compared with the mimic control group, miR-9a-5p mimic significantly decreased BCL2L11 mRNA expression in HPMECs, which was reversed by the co-transfection with the BCL2L11-plasmid (Fig. 4F). Thus, it was shown that BCL2L11 was the direct target gene of miR-9a-5p, and it was negatively regulated by miR-9a-5p.

miR-9a-5p inhibits cell apoptosis induced by CSE in HPMECs by regulating BCL2L11. Cells were treated...
with 1% CSE following transfection. Cell apoptosis, caspase-3 activity, cleaved caspase-3 expression, the cleaved caspase-3/caspase-3 ratio and BCL2L11 expression were detected. The results showed that CSE significantly induced cell apoptosis, increased caspase-3 activity, cleaved caspase-3 expression and the cleaved caspase-3/caspase-3 ratio, and enhanced BCL2L11 protein and mRNA expression in HPMECs. Similarly, compared with the CSE + mimic control group, miR-9a-5p mimic inhibited cell apoptosis, decreased caspase-3 activity, cleaved caspase-3 expression and the
cleaved caspase-3/caspase-3 ratio, and reduced BCL2L11 protein and mRNA expression in HPMECs. These changes were significantly reversed by BCL2L11-plasmid co-transfection (Fig. 5A-F). In conclusion, miR-9a-5p inhibited cell apoptosis induced by CSE in HPMECs by regulating BCL2L11.

Discussion

COPD is a chronic lung disease characterized by incomplete reversible airflow limitation. Common complications include chronic respiratory failure, spontaneous pneumothorax and chronic pulmonary heart disease (4,26). The global incidence of COPD amongst individuals >40 years of age is 9-10%. COPD not only causes respiratory impairment, but also affects circulatory function and can be life-threatening in severe cases (27). HPMEC growth is associated with the process of COPD. HPMECs were used in the present study, and it was found that 1% CSE induced cell apoptosis and increased lncRNA TUG1 expression in HPMECs, consistent with previous reports. The results showed that increased lncRNA TUG1 levels could be used as a diagnostic marker for COPD.

Furthermore, the present study investigated the mechanisms of lncRNA TUG1 in COPD. It was previously reported that there is a binding site between TUG1 and miR-9a-5p (25),
and this association was verified in the present study. In addition, a negative regulatory mechanism was found between lncRNA TUG1 and miR-9a-5p in HPMECs. Endothelial apoptosis serves a key role in the pathogenesis of COPD (13,14). The lncRNA TUG1/miR-9a-5p axis has also been reported to serve a significant role in the regulation of cell apoptosis (25). Thus, it was hypothesized that TUG1 may be involved in COPD pathogenesis through regulating EC apoptosis. Therefore, whether TUG1 affected cell apoptosis in HPMECs via regulation of miR-9a-5p was determined, and several experiments in HPMECs were performed following knockdown of TUG1 or miR-9a-5p expression. FCM analysis indicated that knocking down TUG1 reduced cell apoptosis induced by CSE in HPMECs, and the effects were reversed by downregulating miR-9a-5p. Similarly, the present study predicted and verified that BCL2L11 is the direct target gene of miR-9a-5p. miR-9a-5p was found to negatively regulate BCL2L11 expression in HPMECs. BCL2L11, a member of the BCL-2 family, is located in the outer membrane of mitochondria and it serves as an important regulator of apoptosis (28,29). Thus, miR-9a-5p may regulate HPMECs apoptosis via negatively regulating BCL2L11 expression. The FCM results showed that miR-9a-5p overexpression relieved cell apoptosis induced by CSE in HPMECs, and the effect was reversed by upregulating BCL2L11. The caspase family participates in mediating cell apoptosis, and caspase-3 is a key regulatory molecule involved in numerous pathways in apoptotic signal transduction (30,31). The present study detected caspase-3 activity and expression. The results indicated that TUG1, miR-9a-5p and BCL2L11 affected HPMEC apoptosis by regulating caspase-3.

In conclusion, the present study found that lncRNA TUG1 serves a role in cell apoptosis induced by CSE in HPMECs via modulating a miR-9a-5p/BCL2L11 axis; the mechanism identified in the present study is shown in Fig. S1. These findings suggested that TUG1 was a potential effective target for COPD treatment.

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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

XC and MM conceived and designed the current study, as well as acquired, analyzed and interpreted the data, and prepared the manuscript. YS and XJ contributed to the acquisition and analysis of the data, and prepared the manuscript. All authors read and approved the final manuscript, and confirm the authenticity of all the raw data.

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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