Long non-coding RNA TP73-AS1 accelerates the progression and cisplatin resistance of non-small cell lung cancer by upregulating the expression of TRIM29 via competitively targeting microRNA-34a-5p

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Abstract. Non-small cell lung cancer (NSCLC) is a leading subtype of lung cancer, with high mortality rates. Recently, long non-coding RNAs (lncRNAs) have been associated with NSCLC. The present study aimed to examine the role of the TP73 antisense RNA 1 (TP73-AS1) lncRNA in NSCLC. TP73-AS1 and microRNA(miR)-34a-5p expression levels were measured using reverse transcription-quantitative PCR (RT-qPCR) and chromogenic in situ hybridization (CISH). Cell proliferation, apoptosis, migration and invasion was determined using Cell Counting Kit-8 (CCK-8), flow cytometry, Transwell and Matrigel assays, respectively. The median inhibitory concentration (IC\textsubscript{50}) value of cisplatin (cis-diamminedichloroplatinum; DDP) was assessed using a CCK-8 assay. The interaction between miR-34a-5p and TP73-AS1 or tripartite motif-containing 29 (TRIM29) was predicted using microRNA.org and Starbase, then verified using a dual-luciferase reporter assay. The expression of TRIM29 was quantified at the mRNA and protein level using RT-qPCR and western blot analysis, respectively. TP73-AS1 was significantly upregulated, while miR-34a-5p was downregulated in NSCLC tissues and cells. Functionally, TP73-AS1 knockdown inhibited proliferation, migration, invasion and DDP resistance, whilst inducing apoptosis in NSCLC cells. miR-34a-5p was identified as a target for TP73-AS1, and its inhibition reversed the effects of TP73-AS1 knockdown on NSCLC cells. In addition, TRIM29 was targeted by miR-34a-5p, and its overexpression reversed the effects of miR-34a-5p. Moreover, TP73-AS1 acted as a molecular sponge for miR-34a-5p, increasing the expression of TRIM29. In conclusion, TP73-AS1 contributed to proliferation, migration and DDP resistance but inhibited apoptosis of NSCLC cells by upregulating TRIM29 and sponging miR-34a-5p.

Introduction

Lung cancer remains the leading cause of cancer-associated death worldwide, and the mortality rate in 2019 was higher in males (24%) compared with females (23%) in 2019 in USA (1). Non-small cell lung cancer (NSCLC) is a primary form of lung cancer that accounts for >80% of all lung cancer cases worldwide (2). NSCLC has two major subtypes based on clinical pathology, squamous cell carcinoma and adenocarcinoma (3). The pathogenesis of NSCLC is complex and poorly understood, and a previous study recorded from Brazil that the rates of recurrence (over 50%) and mortality (69.7%) following surgery and different lines of chemotherapy remain extremely high (4). A previous study suggested that novel targets (ROS1 and RET fusions) would benefit NSCLC treatment (5). Therefore, further research on potential therapeutic targets associated with NSCLC remains a necessity.

Long non-coding RNAs (lncRNAs) are RNA molecules >200 nucleotides in length with no or limited protein-coding capacity (6). lncRNAs are involved in multiple biological processes, including cell cycle, cell differentiation and myogenesis, by acting as microRNAs sponges or antisense modulators of mRNA transcripts (7). Recently, lncRNAs have been implicated in several types of cancer, including NSCLC. For example, IncRNA prostate cancer associated transcript 6 acted as an oncogene in NSCLC, contributing to tumor cell growth, migration and invasion (8). Another previous study suggested that small nucleolar RNA host gene 1 was upregulated in NSCLC and promoted tumor cell viability, proliferation and metastasis in NSCLC (9). In addition, IncRNA VPS9D1-antisense RNA 1 (AS1) was overexpressed in NSCLC, which was associated with high rates of metastasis and short survival time within ~10 years (10). Altogether, these previous studies suggested that lncRNAs played essential roles in the progression of NSCLC. Tumor protein 73 AS1 (TP73-AS1) has been found to be involved in several types of cancer (11,12). However, the role of TP73-AS1 in the development of NSCLC remains unclear.
MicroRNAs (miRNAs) are non-coding RNA molecules, ~18-22 nucleotides in length (13). MiRNAs interact with downstream mRNAs, leading to mRNA degradation or translation repression. Therefore, miRNAs modulate diverse cellular processes, including cell differentiation, proliferation and development (14,15). Aberrant increased or decreased expression of miRNAs has been implicated in different human diseases, particularly cancer (16). Previous studies demonstrated that miR-34a-5p acted as a tumor suppressor in various types of cancer, such as colorectal cancer (17), cervical cancer (18) and pancreatic carcinoma (19). However, the role of miR-34a-5p in lung cancer is unknown.

Tripartite motif-containing 29 (TRIM29) is a member of the TRIM protein family, which contains multiple zinc finger motifs and a leucine zipper motif (20). The TRIM29 gene is located on chromosome 11q23 and plays a role in numerous biological processes, including DNA assembly, protein repair and cancer progression (21,22). Notably, TRIM29 not only serves as a tumor inhibitor, such as in invasive breast cancer (23), but also acts as an oncogene in gastric cancer (24), thyroid carcinoma (25) and bladder cancer (26). Nevertheless, the potential role played by TRIM29 in NSCLC is poorly understood.

The present study examined the expression pattern of TP73-AS1 in NSCLC and investigated the role of TP73-AS1 in the progression and drug chemoresistance of NSCLC. It also provided a potential pathway to explain the mechanism of TP73-AS1 action in NSCLC to further understand the pathogenic processes of NSCLC.

Materials and methods

Lung cancer specimens. A total of 50 lung cancer specimens and paired normal tissues within 5 cm of the tumor were collected from patients at The First People's Hospital of Tianmen. Patients' sample collection and 5-year overall survival investigation were conducted between January 2013 and November 2018. All specimens were promptly placed in liquid nitrogen following surgery, then stored at -80˚C. All patients provided written informed consent prior to surgery. The present study was approved by the Ethics Committee of The First People's Hospital of Tianmen. Patients' sample collection and 5-year overall survival investigation were conducted between January 2013 and November 2018. All specimens were promptly placed in liquid nitrogen following surgery, then stored at -80˚C. All patients provided written informed consent prior to surgery.

Reverse transcription-quantitative PCR (RT-qPCR). Total RNA was isolated using TRIzol® reagent (Invitrogen; Thermo Fisher Scientific, Inc.) and cDNA was synthesized using the PrimeScript RT reagent kit (Takara Biotechnology Co., Ltd.) for miR-34a-5p. RT-qPCR was then performed using the SYBR Premix Ex Taq II kit (Takara Biotechnology Co., Ltd.) on an ABI 7900 system (Applied Biosystems; Thermo Fisher Scientific, Inc.) and cDNA was synthesized using the PrimeScript RT reagent kit (Takara Biotechnology Co., Ltd.) for miR-34a-5p. RT-qPCR was then performed using the SYBR Premix Ex Taq II kit (Takara Biotechnology Co., Ltd.) on an ABI 7900 system (Applied Biosystems; Thermo Fisher Scientific, Inc.), according to the manufacturer's instructions. Expression levels were calculated using the 2^-ΔΔCq method (28) and normalized to GAPDH or small nuclear RNA U6. Primers sequences were as follows: TP73-AS1 forward, 5'-TAAAGGTATCGAGATACGGTGATCTGT-3'; #2: 5'-CTCTGCTCTCTCAAGAGAGGTTATTAA-3' and #3: 5'-GCTCTCGCTCCTGCACTCGTTA-3' and #3: 5'-GCTCTCGCTCCTGCACTCGTTA-3'. miR-34a-5p mimic (miR-34a-5p: 5'-UGCCAGUGUCU AGCGUGUUGU-3'), miR-34a-5p inhibitor (anti-miR-34a-5p: 5'-ACAAACGCATGACACATTCAC-3') and their negative controls (si-NC 5'-CTTCCAACGTCAAACGTAAAGATGACAT T-3'), miR-NC (5'-UCUCCGAACUGUCUGAC-3') and anti-NC (5'-UCUCCGAACUGUCUGAC-3').

Reagents and antibodies. Cell lines were cultured in DMEM (Gibco; Thermo Fisher Scientific, Inc.) and 16HBE cells were cultured in DMEM (Gibco; Thermo Fisher Scientific, Inc.) and 10% FBS at 37˚C containing 5% CO2. H1299, H460 and H1650 were cultured in RPMI-1640 supplemented with 10% FBS (both from Gibco; Thermo Fisher Scientific, Inc.) and cDNA was synthesized using the PrimeScript RT reagent kit (Takara Biotechnology Co., Ltd.) for miR-34a-5p. RT-qPCR was then performed using the SYBR Premix Ex Taq II kit (Takara Biotechnology Co., Ltd.) on an ABI 7900 system (Applied Biosystems; Thermo Fisher Scientific, Inc.), according to the manufacturer's instructions. Expression levels were calculated using the 2^-ΔΔCq method (28) and normalized to GAPDH or small nuclear RNA U6. Primers sequences were as follows: TP73-AS1 forward, 5'-CCGGTTPCTTCCAGGTCTGGCA-3' and reverse, 5'-GCCCTACAGGGAAACTTCCAGT-3'; TRIM29 forward, 5'-TTGCTAGGTCCAGGAGCACAGAAATGC-3' and reverse, 5'-CAAATGACCCAAATTCCTTGCAAAACA-3'; GAPDH forward, 5'-ACACAGTCCATGGCCTAC-3' and reverse, 5'-TCCACCACCTGGTTGCTGTA-3'. The primers for miR-34a-5p (forward, 5'-ACATCTCAGGCTGGTGCCAGCGTCTTGGAC-3' and reverse, 5'-CTCAGTCTGTGGTGGAGTGCAATCCGTTGAGAACACCACA-3') and U6 (forward, 5'-CTCCGCTCCGCGCAGAGAC-3' and reverse, 5'-AACAGCTTCAAGATTTTGC GTT-3') were purchased from Guangzhou RiboBio Co., Ltd.
**Cell proliferation and drug resistance analysis.** Cell proliferation and viability was performed using a Cell Counting Kit-8 (CCK-8; Beyotime Institute of Biotechnology). Transfected cells were seeded into 96-well plates (2,000 cells/well) and 10 µl CCK-8 solution was added into each well at different time points post-transfection (0, 24, 48 and 72 h) for another 2 h. The absorbance at 450 nm was measured using a micro-plate reader (Bio-Rad Laboratories, Inc.). Cells were planted into 96-well plates (10,000 cells/well) and then treated with varying concentrations of cisplatin (DDP) (0, 1, 2, 4, 8, 16, 32, 64 and 128 µg/ml) for 48 h, and the concentration of DDP resulting in 50% inhibition of growth (IC\textsubscript{50}) was obtained from the dose-response curve.

**Cell apoptosis analysis.** Apoptosis was assessed using flow cytometry with a Cell Apoptosis kit (Invitrogen; Thermo Fisher Scientific, Inc.), according to the manufacturer’s protocol. A total of 1x10^6 transfected HCC827 and H522 cells was measured using a micro-plate reader (Bio-Rad Laboratories, Inc.). Cells were planted into 96-well plates (10,000 cells/well) and then treated with varying concentrations of cisplatin (DDP) (0, 1, 2, 4, 8, 16, 32, 64 and 128 µg/ml) for 48 h, and the concentration of DDP resulting in 50% inhibition of growth (IC\textsubscript{50}) was obtained from the dose-response curve.

**Cell migration and invasion analysis.** Cell migration and invasion was investigated using the Transwell assay, and 24-well chambers (Corning Inc.). For migration, transfected HCC827 and H522 cells (10,000 cells/well) in serum-free RPMI medium were placed into the top of chambers without Matrigel™ (Corning Inc.). For invasion assays, chambers were pre-coated with Matrigel™ at 4˚C overnight, and then the transfected cells (40,000 cell/well) were seeded into the upper chambers coated with Matrigel™. In either assay, RPMI medium containing 10% FBS was added to the bottom chambers. After a 24-h incubation, the migrated or invaded cells on the bottom surface were fixed with 90% methanol for 20 min and stained with crystal violet (Beyotime Institute of Biotechnology) for 10 min at room temperature. Finally, three randomly selected fields were chosen to evaluate the number of cells under a light microscope (magnification, x100; Olympus Corporation).

**Bioinformatics analysis.** The microRNA.org (version: 2010; http://www.microrna.org/microrna/getDownloads.do) and Starbase (version: 3.0; http://starbase.sysu.edu.cn/agoClipRNA.php?source=circRNA) online tools were used to predict the targets and binding sites between lncRNA and miRNA, or miRNA and mRNA, respectively. Then, the expression of these predicted miRNAs was detected in cells with TP73-AS1

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**Table I. Association between TP73-AS1 expression level and clinicopathological characteristics of patients with non-small cell lung cancer.**

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*P<0.05.
knockdown to monitor which miRNAs were upregulated. Similarly, the expression of predicted mRNAs was detected in cells with miR-34a-5p enrichment to monitor which mRNAs were downregulated, using RT-qPCR assay as detailed above.

**Dual-luciferase reporter assay.** The wild-type (WT) and mutant (MUT) 3’untranslated region (3’UTR) sequences of TP73-AS1 and TRIM29 containing putative binding sites with miR-34a-5p (400 bp, containing 200 bp sequences before or after binding sites) were amplified by PCR and introduced into the luciferase reporter vector PGL3 (Promega Corporation) to generate fusion plasmids, referred to as TP73-AS1-WT, TP73-AS1-MUT, TRIM29-WT and TRIM29-MUT. These fusion plasmids were transfected into HCC827 and H522 cells together with miR-34a-5p mimic or miR-NC using Lipofectamine® 2000 reagent (Invitrogen; Thermo Fisher Scientific, Inc.). After a 48-h transfection, cells were collected and luciferase activity was detected using the dual Glo™ Luciferase Assay System (Promega Corporation), according to the manufacturer's protocol. Relative luciferase activity was expressed as normalization of Renilla luciferase activity to firefly luciferase activity.

**Western blot analysis.** Total protein was isolated using RIPA lysis buffer (Beyotime Institute of Biotechnology) and quantified using BCA Protein Assay kit (Beyotime Institute of Biotechnology). Equal amounts (40 µg) of protein were electrophoresed using 12% SDS-PAGE, then transferred onto PVDF membranes on ice (Bio-Rad Laboratories, Inc.). Next, the membranes were blocked with 5% skimmed milk for 1 h at room temperature and incubated with primary antibodies overnight at 4°C. The following day, the membranes were washed with Tris-buffered saline + Tween-20 buffer, and incubated with a secondary antibody for 2 h. The protein blots were then visualized using an enhanced chemiluminescence kit (Beyotime Institute of Biotechnology) on a ChemiDoc MP imaging system (Bio-Rad Laboratories, Inc.). All antibodies were purchased from Abcam, including primary antibodies against TRIM29 (cat. no. ab244380; 1:1,000) and GAPDH (cat. no. ab181602; 1:10,000) and the HRP-conjugated secondary antibody (cat no. ab205718; 1:5,000). The protein bands were quantified using the ImageJ software (version 1.46; National Institutes of Health).

**Statistical analysis.** All experiments were performed at least three times. Differences between two groups were assessed using paired and unpaired Student's t-test. ANOVA was used for multi-group comparisons, followed by Tukey's post hoc test. Overall survival was evaluated using the Kaplan-Meier method and analyzed by log-rank test. The linear association between variables was analyzed using Pearson correlation coefficient. Statistical analysis was conducted using SPSS v17.0 software (SPSS, Inc.). Data are presented as the mean ± SD. P<0.05 was considered to indicate a statistically significant difference.

**Results**

TP73-AS1 is upregulated in NSCLC tissue and cell lines. Statistical analysis of clinicopathologic features demonstrated that high TP73-AS1 expression was significantly associated with large tumor size, positive lymph node metastasis and advanced TNM stage (Table I). To determine whether the expression of TP73-AS1 was altered in NSCLC, RT-qPCR was performed in tumor samples and normal tissue (n=50 in each group). Compared with that in normal tissue, TP73-AS1 was expressed in tumor tissues at significantly higher levels (Fig. 1A). CISH was also used to detect the expression of...
TP73-AS1 in NSCLC tumor tissues relative to adjacent normal tissues, which revealed a high level of TP73-AS1 in NSCLC tissues (Fig. 1B). Moreover, Kaplan-Meier curves and log-rank analysis indicated that the overall survival rate of patients with NSCLC presenting with high TP73-AS1 levels was lower, compared with patient with low TP73-AS1 expression levels (Fig. 1C).

TP73-AS1 was also highly expressed in NSCLC cell lines, including HCC827, H522, and H23, compared with that in the 16HBE bronchial epithelial cell line (Fig. 1D). In particular, expression of TP73-AS1 was significantly higher in the HCC827 and H522 cell lines relative to the H23 cell line. Therefore, the HCC827 and H522 cell lines were selected for subsequent experiments. Altogether, these findings indicated that increased TP73-AS1 expression levels could play a role in NSCLC progression.

TP73-AS1 knockdown inhibits proliferation, migration and invasion but induced apoptosis of NSCLC cells, and enhances sensitivity to DDP.

To determine the potential role of TP73-AS1 in NSCLC cells, si-TP73-AS1 was transfected into HCC827 and H522 cells for functional analysis. The mRNA expression level of TP73-AS1 was reduced following transfection with si-TP73-AS1#1, si-TP73-AS1#2 and si-TP73-AS1#3 transfection, and the largest decrease was in the si-TP73-AS1#1 group, in both cell lines (Fig. 2A and B). Therefore, si-TP73-AS1#1 was used in subsequent experiments.

Cell proliferation was significantly decreased following si-TP73-AS1 transfection (Fig. 2C and D). By contrast, apoptosis was significantly increased in HCC827 and H522 cells transfected with si-TP73-AS1 (Fig. 2E). In addition, the numbers of migrating and invading cells were significantly reduced in the NSCLC cell lines following transfection with si-TP73-AS1 (Fig. 2F-G). In addition, the viability of HCC827 and H525 cells transfected with si-TP73-AS1 was reduced with increasing DDP concentrations, compared with that in the si-NC-transfected cells (Fig. 2H and I). Furthermore, TP73-AS1 knockdown significantly reduced IC_{50} of DPP in NSCLC cells (Fig. 2J). Taken together, these results demonstrated that TP73-AS1 knockdown inhibited the growth of NSCLC cells, and increased their sensitivity to DDP.

TP73-AS1 targets miR-34a-5p and negatively regulates the expression of miR-34a-5p in NSCLC cells. To explore the mechanism underlying the effect of TP73-AS1 on NSCLC cell lines, the target miRNAs of TP73-AS1 were predicted using microRNA.org and were subsequently screened according to their expression levels. The sectional predicted target miRNAs of TP73-AS1 are shown in Table S1. A significant increase in
miR-34a-5p expression was observed following TP73-AS1 knockdown compared with that in cells transfected with si-NC, while there was no significant difference in the other predicted target miRNAs (Fig. S1). The binding sites between TP73-AS1 and miR-34a-5p were identified using microRNA.org (Fig. 3A). The expression of miR-34a-5p was subsequently found to be increased in HCC827 and H522 cells transfected with TP73-AS1, compared with that in cells transfected with si-NC (Fig. 3B and C). However, there was no difference in luciferase activity in cells co-transfected with TP73-AS1-WT and miR-34a-5p (Fig. 3C and D).

The expression of miR-34a-5p in NSCLC cells was measured, and was found to be significantly downregulated in HCC827 and H522 cells, compared with that in H16HBE cells (Fig. 3E). The efficiency of TP73-AS1 overexpression was subsequently verified. The expression of TP73-AS1 was significantly increased in HCC827 and H522 cells transfected with TP73-AS1, compared with that in cells transfected with the empty vector (Fig. 3F). Further analysis suggested that TP73-AS1 knockdown in HCC827 and H522 cells increased miR-34a-5p expression, while TP73-AS1 overexpression reduced miR-34a-5p expression (Fig. 3G and H). Furthermore, the expression of miR-34a-5p was significantly lower in NSCLC tumor tissues compared with that in normal tissues (Fig. 3I), which was also confirmed using CISH (Fig. 3J). In addition, Pearson correlation coefficient analysis demonstrated that miR-34a-5p expression was negatively associated with TP73-AS1 expression in NSCLC tissues (Fig. 3K). These results demonstrated that miR-34a-5p was targeted by TP73-AS1.

miR-34a-5p inhibition reverses the effects of TP73-AS1 knockdown on proliferation, apoptosis, migration, invasion and drug resistance in NSCLC cells. To determine whether TP73-AS1 exerted its effects by targeting miR-34a-5p, the HCC827 and H522 cell lines were transfected with si-TP73-AS1 and anti-miR-34a-5p. Transfection with anti-miR-34a-5p significantly reduced the expression of miR-34a-5p in HCC827 and H522 cells, compared with that in cells transfected with anti-NC (Fig. 4A). In a CCK-8 assay, cell proliferation was inhibited
following si-TP73-AS1 transfection, compared with that in cells transfected with si-NC. However, this effect was reversed following transfection with si-TP73-AS1+anti-miR-34a-5p, in both HCC827 and H522 cells (Fig. 4B and C). Apoptosis was significantly increased by TP73-AS1 knockdown but was significantly decreased by anti-miR-34a-5p (Fig. 4D). The number of migrating and invading cells was inhibited by TP73-AS1 knockdown, but rescued by miR-34a-5p inhibition (Fig. 4E and F). Furthermore, the IC_{50} of DDP was reduced following TP73-AS1 knockdown. By contrast, anti-miR-34a-5p increased the IC_{50} value in NSCLC cells (Fig. 4G and H). Taken together, these findings suggest that TP73-AS1 knockdown blocked proliferation, migration and invasion, accelerated apoptosis, and reduced resistance to DDP, by upregulating miR-34a-5p in NSCLC cells.

TRIM29 is a miR-34a-5p target in NSCLC cells. To determine whether miR-34a-5p functioned by regulating downstream genes, potential target mRNAs were obtained using Starbase. The sectional predicted target mRNAs of miR-34a-5p are shown in Table SII. A significant decrease was found in TRIM29 expression in cells following miR-34a-5p transfection compared with that in cells transfected with miR-NC (Fig. S2). A binding site was predicted between miR-34a-5p and TRIM29 3’UTR (Fig. 5A). In a dual-luciferase reporter assay, luciferase activity was significantly reduced in HCC827 and H522 cells co-transfected with TRIM29-WT and miR-34a-5p. By contrast, TRIM29-MUT and miR-34a-5p transfection did not affect luciferase activity compared with that in the control group (Fig. 5B and C). The expression of TRIM29 at the protein level was suppressed following transfection with miR-34a-5p mimic, but enhanced by anti-miR-34a-5p transfection, in both HCC827 and H522 cells (Fig. 5D and E). These findings suggested that TRIM29 was a target for miR-34a-5p.

TP73-AS1 knockdown inhibits TRIM29 expression by increasing the expression of miR-34a-5p in NSCLC cells. To determine whether TRIM29 expression was modulated by TP73-AS1 and miR-34a-5p, the expression levels of TRIM29 in HCC827 and H522 cells were investigated under...
different transfection conditions. The protein expression level of TRIM29 decreased following si-TP73-AS1 transfection but was reversed following si-TP73-AS1+anti-miR-34a-5p co-transfection (Fig. 6A). In addition, the expression of TRIM29 was significantly higher in NSCLC tumor tissues compared with that in normal tissues (Fig. 6B). Pearson’s correlation coefficient analysis demonstrated that TRIM29 expression was negatively associated with miR-34a-5p expression (Fig. 6C), but positively associated with TP73-AS1 expression (Fig. 6D). These results suggested
that the expression of TRIM29 was regulated by TP73-AS1 through miR-34a-5p.

**TRIM29 overexpression reverses the effects of miR-34a-5p enrichment on proliferation, apoptosis, migration, invasion and drug resistance in NSCLC cells.** To further investigate the mechanism underlying the effects of TP73-AS1 on NSCLC cells, miR-34a-5p, miR-NC, miR-34a-5p+pcDNA-TRIM29 or miR-34a-5p+pcDNA were introduced into HCC827 and H522 cells. Transfection with pcDNA-TRIM29 led to a significant increase in TRIM29 protein expression compared with that in cells transfected with pcDNA (Fig. 7A). A CCK-8 assay demonstrated that NSCLC cell proliferation was reduced by miR-34a-5p mimic, but promoted following miR-34a-5p+pcDNA-TRIM29 co-transfection (Fig. 7B and C). Apoptosis was significantly increased following miR-34a-5p mimic transfection. However, the rates of apoptosis were reduced following miR-34a-5p+pcDNA-TRIM29 co-transfection, in both HCC827 and H522 cells (Fig. 7D). The numbers of migrating and invading cells were significantly reduced by miR-34a-5p mimic transfection and significantly increased by TRIM29 overexpression (Fig. 7E and F). Furthermore, the IC_{50} of DDP was significantly decreased with miR-34a-5p mimic. By contrast, IC_{50} values were significantly increased by TRIM29 overexpression (Fig. 7G and H). Taken together, this indicated that miR-34a-5p overexpression inhibited NSCLC cell growth and drug resistance by downregulating TRIM29.

**Discussion**

Substantial progress has been made into the management and treatment of lung cancer; however, it remains the leading cause of cancer-related mortality worldwide (29). The identification of additional biomarkers are still required for the treatment of NSCLC. In the present study, TP73-AS1 was highly expressed in NSCLC cells. Functional analysis demonstrated...
that TP73-AS1 downregulation inhibited NSCLC cell growth and resistance to DDP. An interaction between miR-34a-5p, TP73-AS1 and TRIM29 was also identified. Thus, TP73-AS1 promoted tumor cell growth, invasion and drug resistance to DDP through the modulation of TRIM29 and miR-34a-5p in NSCLC cells.

Numerous molecules and genes that are abnormally expressed in tumor tissues significantly affect the occurrence and development of tumors. Previous studies have found that TP73-AS1 was associated with the development of NSCLC. For example, Zhu et al (30) found that TP73-AS1 was expressed at high levels in NSCLC tumor tissues and was associated with poor survival. In addition, increased TP73-AS1 endogenous levels promoted migration and invasion of NSCLC cells (30). Zhang et al (31) found that inhibition of TP73-AS1 reduced NSCLC cell proliferation, inhibited cell cycle progression in vitro, and blocked tumor growth in vivo. Furthermore, a previous study demonstrated that TP73-AS1 promoted temozolomide resistance in glioblastoma multiform cancer stem cells (32). However, whether TP73-ASa could affect DDP resistance in NSCLC remains unclear. In the present study, TP73-AS1 was also upregulated in NSCLC tissues and cells. Moreover, TP73-AS1 knockdown suppressed cell growth, migration and invasion, and inhibited drug resistance to DDP. These findings demonstrated that TP73-AS1 functioned as a tumor promoter in NSCLC. Reactive oxygen species (ROS) play an important role in numerous biological processes, such as cell survival, differentiation and apoptosis (33). Notably, apoptosis induced by intracellular production of ROS has been recognized as one of the basic antitumor mechanisms of DDP (34,35). Therefore, further investigations should focus on the association between ROS production and TP73-AS1 dysregulation in NSCLC.

miR-34a-5p was identified as a target of TP73-AS1 in the present study. A previous study demonstrated that miR-34a-5p was involved in the inhibition of lung cancer progression and metastasis induced by curcumin (36). Another study, based on miRNA microarray and quantitative PCR, found that miR-34a-5p was upregulated in NSCLC tumor tissues treated with luteolin, resulting in the inhibition of tumorigenesis (37). Moreover, miR-34a was downregulated in lung cancer cells, which enhanced their sensitivity to DDP (38). miR-34a overexpression also enhances DDP sensitivity in gastric cancer (39). In agreement with these previous studies, miR-34a-5p was downregulated in NSCLC cells in the present study. In addition, miR-34a-5p overexpression reduced proliferation, migration, invasion and drug resistance to DDP, whilst inducing apoptosis in NSCLC cells. These results provide insight into the role of miR-34a-5p in the growth and DDP resistance of NSCLC cells.

Further experiments suggested that TRIM29 was targeted by miR-34a-5p. Previous studies have documented that TRIM29 functions as a tumor suppressor in several types of cancer, including breast cancer and hepatocellular carcinoma (23,40). By contrast, TRIM29 appears to function as a tumor promoter in lung-related cancers, and accelerates the progression of lung cancer (41-43). Moreover, Liu et al (43) also found that TRIM29 downregulation could augment DDP chemosensitivity in lung cancer cells. Similarly, TRIM29 was upregulated in NSCLC tissues and cell lines in the present study. TRIM29 overexpression reversed the inhibitory effect of miR-34a-5p on progression and DDP resistance in NSCLC. In conclusion, TRIM29 enhanced the growth and DDP resistance to DDP of NSCLC cells.

In summary, the expression of TP73-AS1 was increased in NSCLC tissues and cells, leading to poor overall survival. TP73-AS1 promoted NSCLC progression and resistance to DDP, which was accomplished partially by modulating the TP73-AS1/miR-34a-5p/TRIM29 axis. Thus, the present study suggested that TP73-AS1 might represent a potential biomarker in the treatment of NSCLC.

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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

MS and HC designed the study and performed the experiments. WL and CC collated and analyzed the data. SL and HC were involved in performing the experiments. All authors read and approved the final manuscript.

Ethics approval and consent to participate

The present study was approved by the Ethical Review Committee of The First People's Hospital of Tianmen. All patients provided written informed consent prior to the start of the study.

Patient consent for publication

Not applicable.

Competing interests

The authors declare that they have no competing interests.

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