

The role of microRNA-21 in predicting brain metastases from non-small cell lung cancer

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Objective: This study aimed at exploring the role of microRNA-21 (miR-21) in predicting brain metastases (BM) from non-small cell lung cancer (NSCLC).

Methods: A total of 132 NSCLC patients, including 68 patients with BM and 64 patients without BM, were included in the study. NSCLC cells were collected and assigned to the inhibitor (IN) group, the mock group, and the negative control (NC) group. The quantitative real-time polymerase chain reaction assay was used to detect the miR-21 expression. Cell proliferation, migration, invasion, and apoptosis were detected by colony-forming assay, MTT assay, transwell assay, and flow cytometry, respectively. Angiogenesis was measured by endothelial cell tube formation assay.

Results: The miR-21 expression was higher in NSCLC patients with BM than in those without BM. The miR-21 expression in the IN group was lower than that in the NC and mock groups. Compared with the NC and mock groups, the values of optical density (OD) and the colony-forming number decreased in the IN group. Compared with the NC and mock groups, cell invasion and migration abilities significantly reduced in the IN group. The IN group had higher apoptosis rate than the NC and mock groups. The tube length was shorter and the number of junction points was less in the IN group in comparison to the NC and mock groups.

Conclusion: miR-21 might be a potential biomarker for the development of BM in NSCLC patients and could promote the proliferation, migration, invasion, and angiogenesis of NSCLC cells.

Keywords: non-small cell lung cancer, microRNA-21, brain metastases, angiogenesis

Introduction

Non-small cell lung cancer (NSCLC) is a type of epithelial lung cancer other than small cell lung carcinoma and accounts for approximately 85%–90% of all lung cancers.^{1,2} The incidence rates of NSCLC vary from 22 to 63 per 100,000 men and from 5 to 33 per 100,000 women per year.³ It has been reported that the 5-year survival rate of NSCLC patients ranges from 25% to 73% on the basis of different pathological stages.⁴ Despite advances in NSCLC treatments, the prognosis for NSCLC patients remains poor, with the majority of NSCLC patients dying of pulmonary infection, respiratory failure, brain metastases (BM), and so on.^{5,6} BM is the most common neurologic complication related to systemic cancer, which is up to 10 times more common than primary malignant brain tumors and is a significant burden in the management of patients with advanced cancer.⁷ In addition, among patients with NSCLC, approximately 20%–40% suffer from BM, a major concern in the NSCLC treatment, during the course of the disease, which may significantly influence the survival and quality of life of patients.^{8,9} The prognosis of BM in NSCLC patients has been reported to be very poor, and the median survival of BM patients from lung cancer was less than 1 year.^{9,10} In this regard, it is

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necessary to explore better prognostic markers to predict 1) the occurrence of BM in NSCLC patients and 2) the outcomes to improve the clinical management of NSCLC patients.

MicroRNA-21 (miR-21) is implicated in multiple malignancy-related processes, and overexpressed miR-21 is frequently found in various malignancies, such as breast cancer, liver cancer, esophageal cancer, gastric cancer, brain cancer, colorectal cancer, and NSCLC.^{11–13} Previous studies have also shown that miR-21 is an oncogenic miR and the inhibition of miR-21 expression reduced proliferation, migration, and invasion of cancer cells, including the cells of pancreatic, colorectal, gastric, lung, and NSCLC cancers.^{14–18} However, whether miR-21 leads to the development of BM in NSCLC patients remains unknown. In the present study, we sought to investigate the expression levels of miR-21 in NSCLC patients with or without BM. We also conducted in vitro experiments with the A549 cell line to explore the role of miR-21 in the development of BM in NSCLC patients.

Patients and methods

Study subjects

Between January 2013 and June 2014, a total of 132 NSCLC patients at the First Hospital of Qinhuangdao City were enrolled in this study. Sixty-eight cases were diagnosed with BM (BM+) and 64 cases were diagnosed without BM (BM–). Among the 68 NSCLC patients with BM, 55 (80.9%) had adenocarcinoma, 10 (14.7%) had squamous carcinoma, 2 (2.94%) had sarcoma, and 1 (1.47%) had large cell carcinoma. Among the 64 NSCLC patients without BM, 43 (67.2%) had adenocarcinoma, 10 (15.6%) had squamous carcinoma, 2 (3.13%) had sarcoma, 1 (1.56%) had large cell carcinoma, and 8 (12.5%) had neuroendocrine carcinoma. There were no differences in the clinicopathological features between NSCLC patients with and without BM (Table 1). The diagnosis of NSCLC was confirmed by pathological examination, and the occurrence of BM in NSCLC patients was diagnosed by clinicians and experienced radiologists based on the imaging examination results (cerebral computerized tomography [CT] or magnetic resonance imaging [MRI]) and clinical symptoms. This study was approved by the Ethics Committee of the First Hospital of Qinhuangdao City, and written informed consent was obtained from all subjects.

Sample collection

Blood samples were collected from NSCLC patients and centrifuged at 3,000 rpm at 4°C for 15 min to separate hemocytes and serum. The supernatants were transferred into

Table 1 Clinicopathological characteristics of NSCLC patients with or without brain metastases

Characteristic	BM+ group (n=68)	BM– group (n=64)	P-value
Age (years)	58.5±10.2	61.8±11.4	0.082
Gender			0.325
Male	48	40	
Female	20	24	
Smoking history			0.939
Yes	44	41	
No	24	23	
Pathological type			0.053
Adenocarcinoma	55	43	
Squamous carcinoma	10	10	
Sarcoma	2	2	
Large cell carcinoma	1	1	
Neuroendocrine carcinoma	0	8	
Lymph node metastasis			0.510
Yes	40	34	
No	28	30	

Abbreviations: NSCLC, non-small cell lung cancer; BM+, NSCLC patients with brain metastases; BM–, NSCLC patients without brain metastases.

sterile 2.0 mL Eppendorf (EP) tubes and numbered, and then stored in an ultra-low temperature freezer at –80°C.

Follow-up

All patients were regularly reviewed with timely follow-up. The follow-up was from the time of the first visit to the death or the last follow-up on September 30, 2015. The patients who lost to follow-up in this study were recorded and analyzed with the clinical information of patients on the last contact date. The occurrence time (month) of newly diagnosed BM for the patients without BM at the first visit was carefully recorded during the follow-up.

Cell culture

Human NSCLC cell lines (A549, NCI-H157, NCI-H460, YTMCL-90, and XWLC-05), purchased from the Cell Culture Center of the Institute of Basic Medical Sciences of Chinese Academy of Medical School (Beijing, People's Republic of China), were cultured in Roswell Park Memorial Institute (RPMI)-1640 containing 10% fetal bovine serum (FBS), at 37°C with 5% CO₂. The cultured cells displayed monolayer growth, and the cell adherent rate of passage cells was up to 90.0%. Culture medium was poured out routinely when passaged. After washing twice with phosphate buffer solution (PBS), the cells were digested with 0.25% trypsin, and when the intercellular gaps were enlarged, the trypsin was poured out. The cells were to be blown and beaten into single-cell suspension with RPMI-1640 culture medium containing 10% FBS and passaged routinely.

Quantitative real-time polymerase chain reaction (qRT-PCR)

Total RNA was extracted by a miRNeasy Mini Kit (Qiagen, NV, Venlo, the Netherlands) from the cytolymph or serum. RNA samples (5 µL) were diluted 20 times in RNase-free ultrapure water. The concentration and purity of RNA were determined by ultraviolet (UV) absorbance at 260 and 280 nm (optical density, OD_{260}/OD_{280} ratio) using an UV spectrophotometer. The OD_{260}/OD_{280} ratio between 1.7 and 2.1 indicated that the RNA was highly pure, which could satisfy the requirements of further research process. The cDNA template was generated by reverse transcription with a PCR amplifier. qRT-PCR was conducted by ABI 7500 quantitative PCR System (Thermo Fisher Scientific, Waltham, MA, USA), and ABI 7300 System (Thermo Fisher Scientific) was used for the quantitative detection of miR-21. The reaction conditions were as follows: initial denaturation at 95°C for 10 min and 40 cycles of denaturation at 95°C for 10 s, annealing at 60°C for 20 s, and extension at 72°C for 34 s. U6 snRNA was used as an internal control. The primer sequences of miR-21 and U6 were illustrated in Table 2. The cycle number at threshold (Ct value) was used to calculate the relative expressions of miR-21. The results were presented as fold change, calculated using the $2^{-\Delta\Delta Ct}$ method,¹⁹ using the formula:

$$\Delta\Delta Ct = [Ct_{\text{target gene}} - Ct_{\text{internal gene}}]_{\text{experiment group}} - [Ct_{\text{target gene}} - Ct_{\text{internal gene}}]_{\text{control group}}$$

where $2^{-\Delta\Delta Ct}$ is the multiple proportions of gene expression between the experiment and control groups and Ct represents the cycle number at which fluorescence reached the fixed threshold. The experiments were repeated for a total of three times.

Cell transfection and grouping

Human NSCLC cell lines in the logarithmic growth phase were collected and assigned to three groups: 1) miR-21 inhibitor group (IN group): cells were transfected with synthesized

specific miR-21 inhibitor; 2) negative control (NC) group: cells were transfected with miR-21 inhibitor negative control; and 3) mock group: cells without transfection. The miR-21 inhibitor and miR-21 inhibitor negative control were purchased from Shanghai GenePharma Company (Shanghai, People's Republic of China). A549 cells were inoculated in 50 mL culture flask and cultured in RPMI-1640 complete medium to 30%–50% density. Lipofectamine™ 2000 (Thermo Fisher Scientific) and miR-21 inhibitor or negative control were configured in sterile EP tubes: 1 µL Lipofectamine™ 2000 transferred into 50 µL serum-free medium at room temperature for 5 min; 20 pM miR-21 inhibitor or negative control transferred into 50 µL serum-free medium at room temperature for 5 min. The diluted miR-21 inhibitor or negative control was mixed with Lipofectamine™ 2000 by gentle swirling, with a total volume of 100 µL, at room temperature for 20 min. Each well was added with 100 µL transfection solutions by gentle swirling, and then placed in a 5% CO₂ incubator at 37°C for 6–8 h. Then the solutions were replaced with complete medium and cultured for 48–72 h, and the cells were collected.

MTT assay

After the density of cultured A549 cells reached up to about 80%, the cells were washed twice by PBS, digested with trypsin, and blown and beaten into single-cell suspension. The cells were counted by a cell counter. Cells (3×10^3 – 6×10^3 /well) were cultured in 96-well plates with a volume of 200 µL in each well and repeated for six wells. The cells were cultured at 37°C in a 5% CO₂ incubator for 24–72 h, then each well was treated with 20 µL of MTT (5 mg/mL; Sigma-Aldrich Co., St Louis, MO, USA) reagent and incubated for an additional 4 h. After the medium was removed, 150 µL of dimethyl sulfoxide (DMSO) was added to the wells and the solutions were shaken for 10 min to solubilize the MTT crystals. The OD at 490 nm for each well was measured by an enzyme-linked immunometric meter at the time points of 12, 24, 48, and 72 h. The OD value was set as ordinate and the time point was set as abscissa to graph the MTT curve. Three independent experiment processes were conducted.

Colony formation assay

The A549 cells were inoculated in a 24-well culture plate, with 500 cells in a well, and with three wells for each group. The plate was shaken to make the cells distributed uniformly and then cells were placed in a 5% CO₂ incubator at 37°C. The cell clones were washed twice with PBS, and then fixed with 4% paraformaldehyde (5 mL) at room temperature for 15 min. After the medium was removed, the cells were

Table 2 The primer sequences of real-time qRT-PCR products

Target	Primer	Sequence
miR-21	RT	5'-GTCGTATCCAGTGCAGGGTCCGAGG TATTCGCACTGGATACGACTCAACA-3'
	Forward:	5'-GTGCAGGGTCCGAGGT-3'
	Reverse:	5'-GCCGCTAGCTTATCAGACTGATGT-3'
U6	RT	5'-AACGCTTCACGAATTTGCGT-3'
	Forward:	5'-CTCGCTTCGGCAGCAC-3'
	Reverse:	5'-AACGCTTCACGAATTTGCGT-3'

Abbreviations: miR-21, microRNA-21; RT, reverse transcription; qRT-PCR, quantitative real-time polymerase chain reaction.

stained with crystal violet solution for 10 min, and then rinsed slowly with running water and dried at room temperature. The number of cells was then counted under an inverted microscope. Individual clones with more than 50 cells were counted and the colony-forming efficiency was calculated with the formula: colony-forming efficiency = the number of clones/the number of inoculated cells $\times 100\%$. This experiment was repeated three times.

Flow cytometry (FCM)

The cells were inoculated in a six-well culture plate with 1×10^6 cells in each well and cultured with serum-free medium for 12 h to synchronize the cells. Then the cells were centrifuged at 1,000 rpm for 5 min and the original medium was removed. The cells were digested and collected after centrifugation. The cells were washed twice with PBS and were resuspended in precooled 75% ethanol and fixed at -20°C overnight. After centrifugation, the supernatant was discarded and was then washed twice with PBS. The cells (with a final concentration of $1 \times 10^6/\text{mL}$) were mixed with 50 μL propidium iodide (PI, 0.5 mg/mL), stained away from light at 4°C for 30 min, and filtered with a screen mesh. A FACSCalibur flow cytometer (BD Biosciences, San Jose, CA, USA) was used to determine and analyze the cell cycle distributions. Argon ion laser was applied to observe the fluorescence of PI at 450 nm.

The A549 cells were collected at 48 h after transfection, washed with PBS for three times, and were then added into a precooled $1 \times$ binding buffer (500 μL). Annexin-V-FITC (fluorescein isothiocyanate) (5 μL) and PI (2.5 μL) were mixed into the cell suspension and analyzed by a BD FACS Aria I cell sorter (BD Biosciences). The cell populations were detected in a dot-matrix plot: viable cells (FITC⁻/PI⁻) were presented in the left lower quadrant (Q4), early apoptotic cells (FITC⁺/PI⁻) were presented in the right lower quadrant (Q3), and necrotic and late apoptotic cells (FITC⁺/PI⁺) were presented in the right upper quadrant (Q2). The apoptosis rate = the percentage of FITC⁺/PI⁻ (Q3) + the percentage of FITC⁺/PI⁺ (Q2).

Transwell migration and invasion assays

At 48 h after transfection, the cells were digested with 2.5% trypsin and suspended in serum-free Dulbecco's Modified Eagle's Medium (DMEM). Two-hundred μL of a single-cell suspension of cells (with a final concentration of $1 \times 10^6/\text{mL}$) was placed in the upper chamber, whereas the lower chamber of the 24-well culture plate was loaded with 600 μL RPMI-1640 containing 10% FBS, and cultured at 37°C for 24 h. The experiment was repeated three times with three parallel wells

in each group. After incubation, the matrix and noninvading cells in the upper chamber were carefully removed, stained with 0.1% crystal violet solution for 5 min, and the redundant stain was removed with clean water. The cells that penetrated through the Matrigel to the lower surface of the filter were calculated and photographed under a light microscope at $\times 200$. Five optional visual fields of the cell invasion, including up, down, middle, left, and right, were chosen for counting, and the average value was calculated. Transwell migration assay was performed using transwell without a layer of Matrigel on the upper surface. The rest of the experimental procedure was the same as the invasion assay.

Endothelial cell tube formation assay

After the extracellular matrix gels were fully solubilized, the cells were placed into a 96-well culture plate and incubated at 37°C with 5% CO_2 for 60 min. The cells were collected by centrifugation at 1,500 rpm for 5 min after removal of dead cells in the culture supernatants. A549 cells were suspended in the supernatants with a final concentration of $2 \times 10^5/\text{mL}$. A volume of 150 μL of supernatant was added into each well of the 96-well culture plate and incubated at 37°C with 5% CO_2 for 18 h. The cell tube formation in each group was observed and imaged under a light microscope at $\times 20$. Five optional visual fields were chosen randomly for counting.

Statistical analysis

SPSS 21.0 statistical software (SPSS Inc., Chicago, IL, USA) was applied for statistical analysis. The measurement data were expressed by mean and standard deviation. One-way analysis of variance (ANOVA) was used for multiple group comparisons after the variance homogeneity test. Least significant difference (LSD) unpaired *t*-test was used in pairwise comparison of multiple groups or averages. Receiver operating characteristic (ROC) curve was used to measure the value of apparent diffusion coefficient (ADC), with BM+ or BM- as state variables and miR-21 expression of 64 NSCLC patients without BM as test variables. The sensitivity, specificity, accuracy rate, and the area under the curve (AUC) were calculated. Kaplan-Meier survival curve was used to analyze the influence of miR-21 expression on the occurrence time of BM. $P < 0.05$ was considered statistically significant.

Results

The expression of miR-21 in NSCLC patients

The qRT-PCR assay was used to detect the expression of miR-21 in the BM+ and BM- groups. As presented in

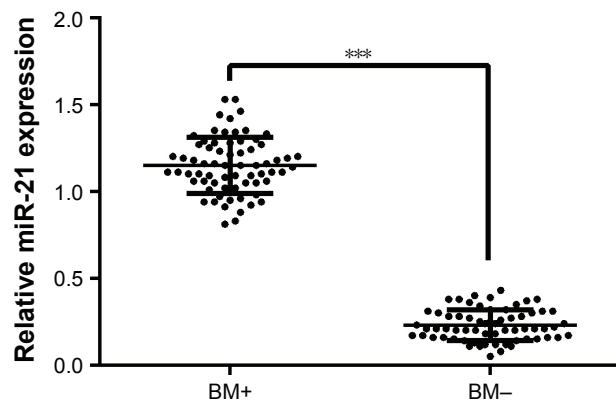


Figure 1 Expression of miR-21 in NSCLC patients with or without BM detected by qRT-PCR.

Note: *** $P < 0.001$.

Abbreviations: miR-21, microRNA-21; NSCLC, non-small cell lung cancer; BM, brain metastases; BM+, NSCLC patients with brain metastases; BM-, NSCLC patients without brain metastases; qRT-PCR, quantitative real-time polymerase chain reaction.

Figure 1, the results showed that the expression of miR-21 in the BM+ group was significantly higher than that in the BM- group ($P < 0.001$).

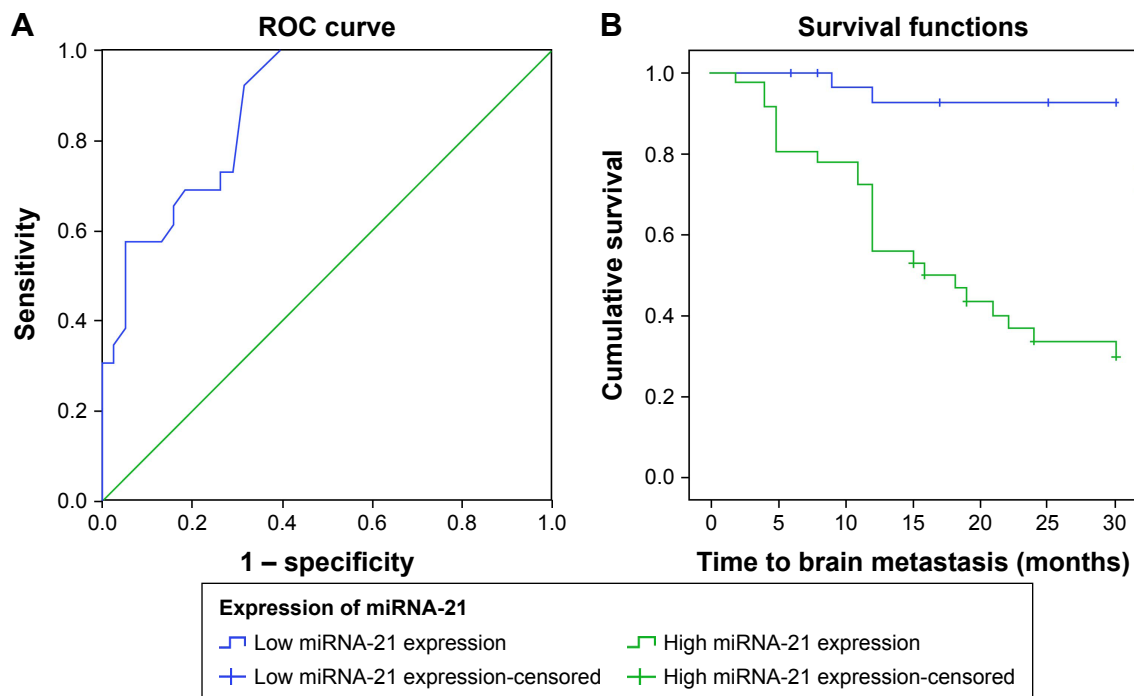
Correlation between miR-21 expression and the risk of BM in NSCLC patients

Among 64 NSCLC patients without BM, 26 cases were newly diagnosed with BM during the substantial follow-ups.

True positive rate (sensitivity) was defined as the percentage of NSCLC patients with BM whose miR-21 expression was higher than the threshold of ROC curve, while true negative rate (specificity) was defined as the percentage of NSCLC patients without BM whose miR-21 expression was lower than the threshold. The ROC curve showed that the AUC was 0.873 (95% confidence interval [CI]: 0.791–0.956), and the sensitivity and specificity were 92.3% and 60.7%, respectively, with a best cutoff point of 0.205 (Figure 2A). Kaplan–Meier survival curve was used to analyze the correlation between the miR-21 expression and the occurrence time (month) of BM. The results indicated that the expression more than 0.205 indicated enhanced expression of miR-21 and the expression less than or equal to 0.205 indicated decreased expression of miR-21. Furthermore, the average time of BM occurrence with high expression of miR-21 was 17.4 months, and that with low expression of miR-21 was 26.1 months (Figure 2B).

Expression of miR-21 in five NSCLC cell lines

As shown in Figure 3, qRT-PCR was used to detect the expression of miR-21 in human NSCLC cell lines A549, NCI-H157, NCI-H460, YTMCLC-90, and XWLC-05. The results suggested that miR-21 expression was the highest



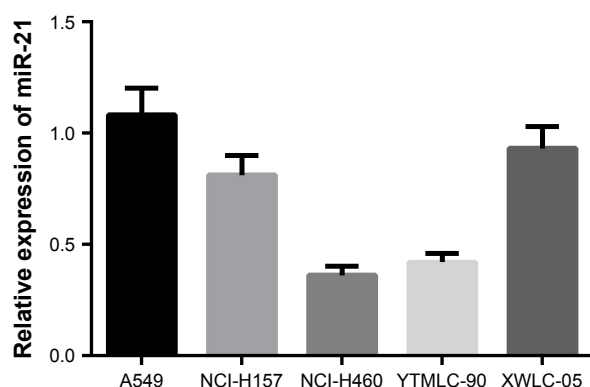


Figure 3 Expression of miR-21 in five human NSCLC cell lines (A549, NCI-H157, NCI-H460, YTMLC-90, and XWLC-05).

Abbreviations: miR-21, microRNA-21; NSCLC, non-small cell lung cancer.

in A549 cells. Therefore, A549 cell line was chosen for further studies.

Expression of miR-21 in A549 cells after transfection

At 48 h after transfection, the expression of miR-21 in the IN group was significantly lower than that in the NC and mock groups (both $P < 0.05$; Figure 4). However, no significant difference in miR-21 expression was observed between the NC and mock groups ($P > 0.05$).

Effects of miR-21 inhibitors on the proliferation and clone formation of A549 cells

Compared with the NC and mock groups, the OD values at 24, 48, and 72 h significantly decreased in the IN group after transfection (all $P < 0.05$), whereas no significant difference was observed between the NC and mock groups ($P > 0.05$;

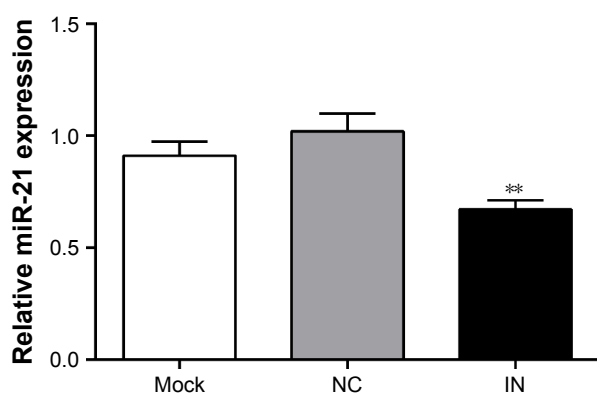


Figure 4 Expression of miR-21 in A549 cells among three groups after transfection detected by qRT-PCR.

Note: **Compared with the NC group or mock group, $P < 0.01$.

Abbreviations: miR-21, microRNA-21; qRT-PCR, real-time quantitative polymerase chain reaction; IN, inhibitor group; NC, negative control group; mock, mock group.

Figure 5A). The results showed that the down-regulated expression of miR-21 might inhibit the proliferation of A549 cells. The number of clones in the IN group significantly decreased at 48 h after transfection ($P < 0.05$), whereas no significant differences were observed between the NC and mock groups ($P > 0.05$; Figure 5B).

Effects of miR-21 inhibitors on cell cycle and apoptosis of A549 cells

As shown in Figure 6A, the proportions of cells at G0/G1 phase in the NC and mock groups were $42.87\% \pm 3.48\%$ and $40.39\% \pm 4.23\%$, respectively, at 48 h after transfection. There were no significant differences between the NC and mock groups ($P > 0.05$). As compared to the NC and mock groups, the proportions of cells at G0/G1 phase in the IN group significantly increased ($62.25\% \pm 3.00\%$; $P < 0.05$). Furthermore, the proportions of cells at S phase in the NC and mock groups were $50.79\% \pm 7.17\%$ and $54.48\% \pm 3.53\%$, respectively, with no significant differences between the groups ($P > 0.05$). As compared to the NC and mock groups, the proportions of cells at S phase in the IN group significantly decreased ($34.35\% \pm 3.14\%$; $P < 0.05$). The proportions of cells at G2/M phase in the NC, mock, and IN groups were $6.33\% \pm 3.88\%$, $5.13\% \pm 1.54\%$, and $3.30\% \pm 0.25\%$, respectively. No significant differences were observed among the three groups ($P > 0.05$). These results indicated that miR-21 inhibitors might reduce the proliferation of A549 cells.

The apoptosis rate of A549 cells in the IN group was $17.58\% \pm 1.42\%$ at 48 h after transfection, while the apoptosis rates of A549 cells in the NC and mock groups were $5.69\% \pm 1.03\%$ and $6.02\% \pm 1.15\%$, respectively. The apoptosis rate in the IN group was significantly higher than in the NC and mock groups ($P < 0.05$), while there was no significant difference in the rate between the NC and mock groups ($P > 0.05$; Figure 6B). These results indicated that miR-21 inhibitors might promote the apoptosis of A549 cells.

Effects of miR-21 inhibitors on the migration and invasion of A549 cells

The results of transwell migration and invasion assays showed that the number of invading cells in the IN group significantly decreased (25.67 ± 6.21 [mean $\pm$ SD]) compared to the NC group (58.55 ± 7.48) and the mock group (58.74 ± 8.57) ($P < 0.05$; Figure 7A). The results demonstrated that miR-21 inhibitors might reduce the invasion of cells significantly. Moreover, the number of migrating cells in the IN group significantly decreased (52.34 ± 8.74) compared to the

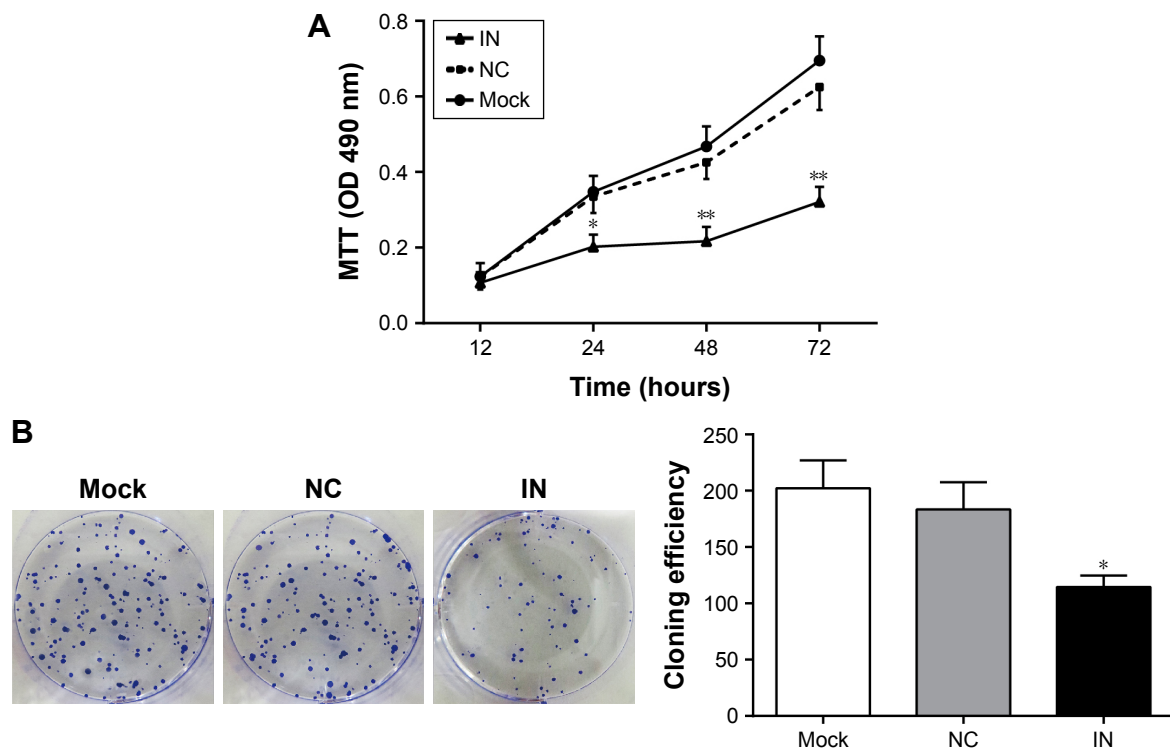


Figure 5 Effects of miR-21 inhibitors on the proliferation and clone formation of A549 cells. **(A)** The effect of miR-21 inhibitors on the proliferation of A549 cells detected by MTT assay; **(B)** the effect of miR-21 inhibitors on the clone formation of A549 cells measured by colony-forming assay (mean and standard deviation).

Notes: *Compared with the NC group or mock group, $P < 0.05$; **compared with the NC group or mock group, $P < 0.01$.

Abbreviations: miR-21, microRNA-21; IN, inhibitor group; NC, negative control group; mock, mock group; OD, optical density.

NC group (98.74 ± 11.24) and the mock group (95.67 ± 10.25) ($P < 0.05$; Figure 7B).

Effect of miR-21 inhibitors on angiogenesis of A549 cells

The A549 cells in the IN group rarely showed tube formation, whereas the A549 cells in the NC and mock groups had dense and homogeneous lumens (Figure 8). The differences

in tube length were analyzed by ImageJ software, and the results showed that the tube length of A549 cells in the IN group was significantly shorter and the number of junction points was less compared to those in the NC group ($P < 0.05$).

Discussion

Nowadays, NSCLC patients show a higher incidence of BM, which may cause central nervous system dysfunction

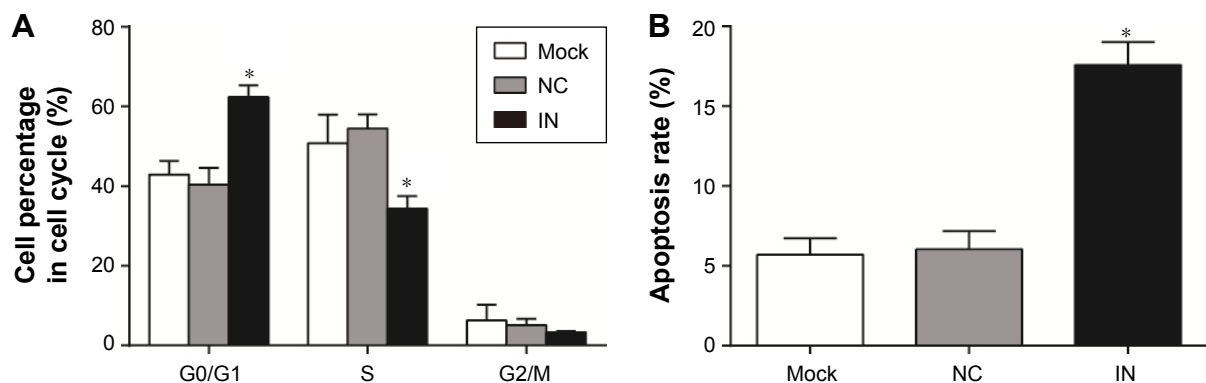


Figure 6 Effects of miR-21 inhibitors on cell cycle and apoptosis of A549 cells. **(A)** The effect of miR-21 inhibitors on cell cycle of A549 cells detected by FCM; **(B)** the effect of miR-21 inhibitors on apoptosis of A549 cells detected by FCM.

Note: *Compared with the NC group or mock group, $P < 0.05$.

Abbreviations: miR-21, microRNA-21; FCM, flow cytometry; IN, inhibitor group; NC, negative control group; mock, mock group.

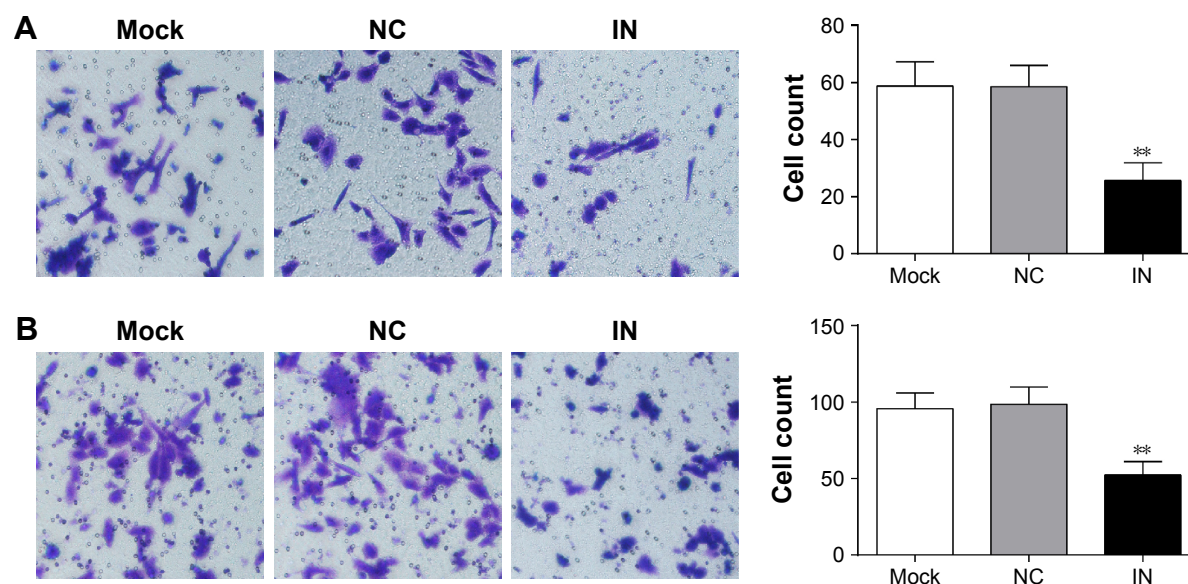


Figure 7 Effects of miR-21 inhibitors on the migration and invasion of A549 cells. **(A)** The effect of miR-21 inhibitors on the invasion of A549 cells detected by transwell assay; **(B)** the effect of miR-21 inhibitors on the migration of A549 cells detected by transwell assay.

Notes: **Compared with the NC group or mock group, $P < 0.01$. Magnification was 200 $\times$.

Abbreviations: miR-21, microRNA-21; IN, inhibitor group; NC, negative control group; mock, mock group.

and may seriously affect the quality of life of patients.⁶ BM causes significant neurologic, cognitive, and emotional difficulties and negatively impacts survival,²⁰ which is one of the leading causes of death in NSCLC patients. Therefore, early diagnosis and timely treatment of BM are essential to the prognosis of NSCLC patients with BM.^{5,21} In this study, we investigated the expression of miR-21 in NSCLC patients with or without BM to explore the correlation between the miR-21 expression and the development of BM in NSCLC patients. The results showed that the NSCLC patients with BM showed higher expression of miR-21 compared to those without BM, indicating that enhanced expression of miR-21

might be correlated with the development of BM. Similarly, Yang et al have demonstrated that overexpression of miR-21 was found in NSCLC tissues, and overexpression of miR-21 was significantly correlated with tumor size and tumor-node-metastasis (TNM) stage of NSCLC patients.²² Furthermore, Oh et al have revealed that NSCLC patients with distal metastasis had higher miR-21 expression than those without distal metastasis.²⁰

Furthermore, the ROC curve showed that the sensitivity and specificity for the miR-21 in predicting the occurrence of BM were 92.3% and 60.7%, respectively, and the Kaplan–Meier curve showed that miR-21 was a potential

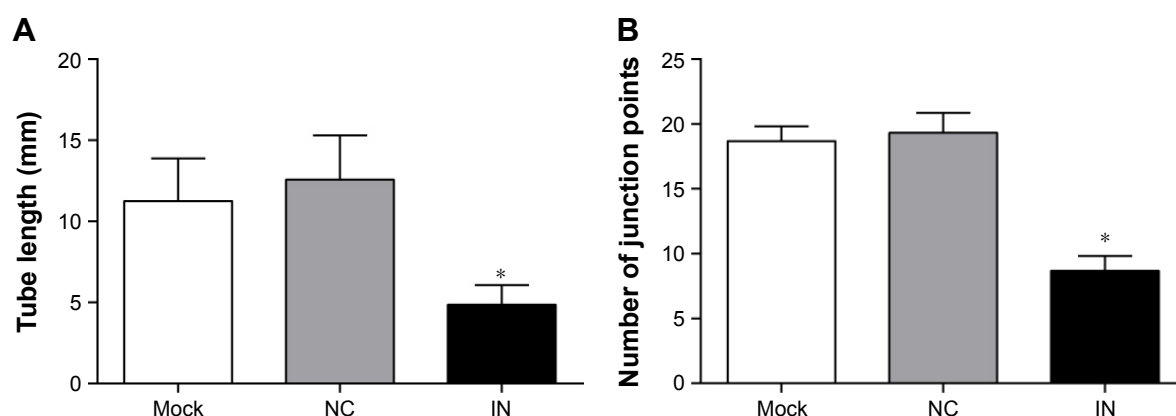


Figure 8 Effects of miR-21 inhibitors on angiogenesis of A549 cells. **(A)** The tube circumference of A549 cells in the three groups analyzed by ImageJ software; **(B)** the tube junctions of A549 cells in the three groups analyzed by ImageJ software.

Note: *Compared with the NC group or mock group, $P < 0.05$.

Abbreviations: miR-21, microRNA-21; IN, inhibitor group; NC, negative control group; mock, mock group.

risk factor for the development of BM. These results demonstrated that the overexpression of miR-21 might be closely implicated in the development and progression of BM in NSCLC patients, which could act as a potential biological marker in predicting the development of BM in NSCLC patients. A variety of miRs were observed in brain-metastatic carcinomas, which can act as potential diagnostic markers and therapeutic targets for various cancers with BM.^{23,24} Nadiya et al have found that the miR-21 expression was significantly increased in the cerebrospinal fluid of patients with glioblastoma and BM of breast and lung cancers, indicating that miR-21 may be a potential biomarker for both glioblastoma and BM.²⁵ Singh et al have demonstrated that miR-21 is overexpressed in lung cancer patients and is associated with a poor prognosis; meanwhile, Singh et al suggested that miR-21 might regulate the metastatic behavior of BM initiating cells by targeting STAT3, and hence blocking the STAT3-miR-21 pathway could be a novel therapeutic treatment in patients with lung-to-brain metastasis.²⁶ A previous study performed by Zhang et al has demonstrated that NSCLC patients with distal metastasis had higher miR-21 expression than those without distal metastasis, and the overexpression of miR-21 may be involved in the metastasis and invasion of tumor cells.¹⁸

For a better understanding of the underlying mechanisms of miR-21 in the development of BM in NSCLC patients, we conducted a cell experiment with the A549 cell line. The results of in vitro experiment showed that miR-21 inhibitors might reduce cell proliferation, clone, migration, invasion, and tumor angiogenesis. Furthermore, miR-21 inhibitors might promote the apoptosis of A549 cells. These results indicated that miR-21 might be implicated in the development of BM in NSCLC patients, and the inhibition of miR-21 expression might be a novel therapeutic approach for the NSCLC patients with BM. It is well known that miR-21 is overexpressed in various human tumors and has been linked to various cellular processes that are altered in cancer cells. Recently, studies have showed that knockdown of miR-21 expression inhibited tumor cell proliferation, migration, and invasion, as well as induced apoptosis in vitro and suppressed tumor cell growth in vivo.^{27,28} Yang et al have found that knockdown of miR-21 expression can inhibit the cell proliferation and migration in vitro, and that miR-21 may regulate the metastatic behavior by promoting cell proliferation, survival, migration, and invasion.²⁹ In addition, down-regulation of miR-21 expression following transfection of miR-21-inhibitor may restrain the cell proliferation and migration of A549 cells through up-regulation

of PDCD4.³⁰ Bornachea et al have found that the expression of miR-21 increased in human metastatic lung cancer, and the elevated miR-21 expression may be closely correlated with distant metastasis.³¹

Conclusion

In conclusion, our analysis suggested that miR-21 was overexpressed in NSCLC patients with BM, and knockdown of miR-21 expression might restrain cell proliferation, migration, invasion, and angiogenesis, and might induce tumor cell apoptosis, indicating that miR-21 might be a potential biomarker for the development of BM in NSCLC patients and knockdown of miR-21 might be a novel therapeutic approach for NSCLC patients with BM. We found that the specificity was 66.7%, 67.8%, and 61.5% in the ROC curve analysis of other relative references,^{32,33} while the specificity was only 60.7% in our experiment. Because the number of NSCLC patients visiting our hospital every year was small, there was separation in the data of miR-21 expression. Therefore, more samples shall be used in the future investigations. In addition, further studies on the targets of miR-21 will be conducted for better understanding of correlation between the underlying mechanism of miR-21 and BM.

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Disclosure

The authors report no conflicts of interest in this work.

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