Tumor-associated macrophages secret exosomal miR-155 and miR-196a-5p to promote metastasis of non-small-cell lung cancer

Xiang Li#, Zhipeng Chen#, Yaojun Ni1,2, Chengyu Bian1, Jingjing Huang1, Liang Chen1, Xueying Xie3, Jun Wang1

1Department of Thoracic Surgery, Jiangsu Province People’s Hospital and the First Affiliated Hospital of Nanjing Medical University, Nanjing, China; 2Department of Cardiothoracic Surgery, Huai’an First People’s Hospital Affiliated to Nanjing Medical University, Huai’an, China; 3State Key Laboratory of Bioelectronics, School of Biological Sciences and Medical Engineering, Southeast University, Nanjing, China

Contributions: (I) Conception and design: X Xie, J Wang; (II) Administrative support: L Chen; (III) Provision of study materials or patients: X Xie, Z Chen, C Bian; (IV) Collection and assembly of data: J Wang, X Li, X Xie, Y Ni; (V) Data analysis and interpretation: X Li, Z Chen, J Huang, Y Ni; (VI) Manuscript writing: All authors; (VII) Final approval of manuscript: All authors.

#These authors contributed equally to this work.

Correspondence to: Xueying Xie. State Key Laboratory of Bioelectronics, School of Biological Sciences and Medical Engineering, Southeast University, 2 Sipailou, Nanjing, Jiangsu 210096, China. Email: xxying.rcbs@seu.edu.cn; Jun Wang. Department of Thoracic Surgery, Jiangsu Province People’s Hospital and the First Affiliated Hospital of Nanjing Medical University, Nanjing, Jiangsu 210029, China. Email: drwangjun@njmu.edu.cn.

Background: Understanding the molecular basis underlying metastasis of non-small cell lung cancer (NSCLC) may provide a new therapeutic modality for the treatment of NSCLC. However, the mechanisms by which tumor-associated macrophages (TAMs) affect NSCLC metastasis remain undefined. In this study, we aimed to discover a novel regulatory pathway involved in NSCLC metastasis.

Methods: Cell Counting Kit-8 (CCK-8), Transwell, western blot assays were used to assess cell viability, migration, invasion and epithelial-mesenchymal transition (EMT). Exosomes from macrophages medium were characterized, and in vitro cell coculture was further conducted to investigate M2 derived exosomes mediated crosstalk between TAMs and tumor cells. Besides, miRNA microarray was used to analyze miRNA expression profiles of M0 and M2 derived exosomes. Luciferase reporter assay was used to verify the potential binding between miRNA and mRNA. Moreover, 6-week-old male BALB/c nude mice were performed to establish transplantation tumor model using tail vein injection. Hematoxylin & eosin staining was used to detect the metastasis of tumor tissues.

Results: We found that M2 TAMs were the main TAMs in metastatic tissues of NSCLC patients and exosomes derived from M2 TAMs were able to promote cell viability, cell migration, cell invasion and EMT in NSCLC. We demonstrated that miR-155 and miR-196a-5p were abundant in M2 TAMs and exosomes secreted by M2 TAMs. Functional experiments demonstrated that the deletion of miR-155 and miR-196a-5p in M2 TAMs significantly prevented NSCLC metastasis in vitro and in vivo. To clarify the mechanism governing miR-155 and miR-196a-5p from M2 TAMs, we carried out bioinformatics analysis to predict potential target genes. Mechanistically, miR-155 and miR-196a-5p directly bound to the 3’-UTR of Ras association domain family member 4 (RASSF4), and negatively regulating RASSF4 expression. At last, rescue assays demonstrated that miR-155 and miR-196a-5p exerted its performance by RASSF4.

Conclusions: Overall, we revealed a new regulatory pathway that was M2 TAMs secreted exosomal miR-155 and miR-196a-5p to promote NSCLC metastasis. This dynamic and reciprocal cross-talk between NSCLC and macrophages innovatively provided a potential opportunity for diagnosis and treatment of NSCLC.

Keywords: Tumor-associated macrophages (TAMs); miR-155; miR-196a-5p; exosomes; non-small-cell lung cancer (NSCLC); epithelial-mesenchymal transition (EMT)

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Introduction

According to global cancer statistics in 2018, lung cancer accounts for ~12% of newly diagnosed cancer cases and ~18% of total cancer deaths (1). About 85% of diagnosed lung cancer is non-small cell lung cancer (NSCLC), which have a low 5-year survival rate (<15%) (2,3). Although timely treatments such as surgery, chemotherapy and radiotherapy can prolong the overall survival of patients, NSCLC is still the most fatal disease. The causes of lung cancer death vary and metastasis is a predominant factor (4). Therefore, understanding the molecular basis of NSCLC metastasis is critical for proposing new therapeutic approaches for its treatment (5-7). Several classes of biomolecules, such as microRNAs (miRNAs), matrix metalloproteinases and transcription regulatory protein BACH1, have been well studied in the metastasis of NSCLC (8-11). Although tumor microenvironment is also known to mediate NSCLC metastasis, the role of key components of tumor microenvironment in NSCLC metastasis has not been fully elucidated (12,13).

The tumor microenvironment consists of cancer cells and stromal cells that include macrophages, endothelial cells and fibroblasts (14-16). Moreover, products of these cells, such as cytokines, growth factors and enzymes, are also present in the tumor microenvironment. Among these components in the tumor microenvironment, tumor-associated macrophages (TAMs) are the pivotal orchestrators as a class of immune cells (17). TAMs have two populations, M1 and M2 types. It has been found that M1 TAMs have an anti-tumor function, whereas M2 TAMs can promote tumor metastasis (18). For NSCLC, M1 TAMs can suppress the angiogenesis and are positively related with the survival time of NSCLC patients (19,20). In contrast, M2 TAMs are found to promote NSCLC metastasis through activating the epithelial-mesenchymal transition (EMT) of NSCLC cells and the invasion of lung cancer cells (20,21). However, the detailed mechanisms by which M2 TAMs promote NSCLC metastasis remain elusive.

Exosomes are small vesicles secreted by parental cells into extracellular microenvironment to mediate cross-talk between parental cells and target cells (22,23). Almost all cell types have the capacity to release exosomes and the size of exosomes is ~30–150 nm in diameter. To mediate cell-cell communication, exosomes deliver proteins, miRNAs or mRNAs into target cells to change the gene expression (24). For instance, gastric cancer cells secret exosomes carrying miR-21 into peritoneal mesothelial cells to promote peritoneal metastasis (22). For the cross-talk between cancer cells and TAMs, exosomes carrying miRNAs are also key players. Taking an example, ovarian cancer cells secret exosomes containing miR-222 to induce M2 polarization of TAMs (25). Moreover, M2 TAMs also can release exosomes carrying miR-501 to promote the progression of pancreatic cancer (26). Nevertheless, whether M2 TAMs promote NSCLC metastasis through exosomes and miRNAs remains unclear.

Here, we report that M2 TAMs promote EMT of NSCLC cells through secreting exosomes. MiR-155 and miR-196a-5p are the key functional molecule in M2 TAMs-released exosomes that promote EMT of NSCLC cells through targeting RASSF4.

Methods

Patients

NSCLC localized tissues (n=15) and metastatic tissues (n=15) were obtained from patients in Jiangsu Province People’s Hospital between 2015 and 2016. The project has attained informed consent from all patients and was approved by the ethics committee of Jiangsu Province People’s Hospital and the First Affiliated Hospital of Nanjing Medical University (No. 2019-SR-266). The study was conducted in accordance with the Declaration of Helsinki (as revised in 2013).

Cell lines and cell culture

The A549 and THP-1 cells were obtained from Chinese Academy of Sciences, Shanghai Institute of Biochemistry and Cell Biology (Shanghai, China). A549/Luc cells (A549 cells stably express luciferase) were constructed by
Synthgene (Nanjing, China). THP-1 cells were cultured in Roswell Park Memorial Institute (RPMI) 1640 medium and A549 cells were cultured in DMEM (Dulbecco’s Modified Eagle’s Medium) (Gibco, Rockville, USA) supplemented with 10% FBS (fetal bovine serum) (Gibco, Rockville, USA) and 100 µg/mL streptomycin and penicillin (Gibco, Rockville, USA) in a humidified atmosphere at 37 °C. After 7 days in culture, THP-1 cells were treated with 100 ng/mL phorbol 12-myristate-13-acetate (PMA) to differentiate into macrophages for 48 h. Following treatment with 20 ng/mL interleukin-4 (IL-4) for 72 h, the cells were polarized into M2 macrophages.

**Cell transfection**

MiR-155 inhibitor, miR-155 mimic, miR-196a-5p inhibitor, miR-196a-5p mimic, short hairpin (sh-RASSF4) and negative control (NC) were synthesized in Gene Pharma Company (Shanghai, China). Lipofectamine 2000 (Invitrogen, USA) was performed for cell transfection based on the specification.

**Immunohistochemistry assay**

The immunohistochemistry assay was performed according to Yin et al. (26). The sections were incubated with the primary antibody of CD68 (Ab955, 1: 100, Abcam, Cambridge, UK) at 4 °C overnight and horseradish peroxidase labeled goat anti-mouse IgG antibody (A205719, 1:200, Abcam, Cambridge, UK) at room temperature for 1 h. The color reaction was performed with diaminobenzidine chromogen solution (Dako, Carpinteria, USA). Brown-yellow particles represented the positive expression of CD68 protein and the blue particles represented the nucleus stained by hematoxylin (Sigma, USA).

**Flow cytometry assay**

The macrophages were extracted by the tumor macrophage isolation fluid (Sangon Biotch, China). Then, macrophages were stained with CD163 (Abcam, USA) and CD206 (Abcam, USA) for 30 min at 4 °C. The labeled cells were analyzed by FACScan flow cytometry (BD Biosciences, USA).

**RNA extraction and quantitative real-time PCR analysis**

The extraction and reverse transcription of total RNA were performed according to the previous report (29). The expression levels of TNF-α, IRF5, IRF4, Arg-1 and miR-155 were analyzed by quantitative real-time PCR with the glyceraldehyde-3-phosphate dehydrogenase (GAPDH) gene or U6 as a standard control. Primers of TNF-α, IRF5, IRF4, Arg-1, miR-155, miR-196a-5p, miR-3091-3p, miR-12206-5p, miR-12180-3p, miR-12206-5p, and miR-12180-3p were as follows: TNF-α (Forward: 5’-CCTCTCTCTCTCAGCCTCTCT-3’; Reverse: 5’-GAGGACCTGGGAGTAGATGAG-3’); IRF5 (Forward: 5’-GGCGTATGTTGCTTCTGTTGC-3’; Reverse: 5’-GGCGTATGTTGCTTCTGTTGC-3’); miR-155 (Forward: 5’-GGAGGTTAATGCTAATCGTGATAG-3; Reverse: 5’-GTGCAGGGTCCGAGGT-3’); miR-196a-5p (Forward: 5’-CAGGACTGGGAGCTCGAGATGAG-3’; Reverse: 5’-CGGACTGGGAGCTCGAGATGAG-3’); miR3091-3p (Forward: 5’-GAGGTTAATGCTAATCGTGATAG-3; Reverse: 5’-GTGCAGGGTCCGAGGT-3’); miR-12206-5p (Forward: 5’-CAGGACTGGGAGCTCGAGATGAG-3’; Reverse: 5’-CGGACTGGGAGCTCGAGATGAG-3’); miR-12180-3p (Forward: 5’-GAGGTTAATGCTAATCGTGATAG-3; Reverse: 5’-GTGCAGGGTCCGAGGT-3’). These primers were synthesized and purified by RiboBio (Guangzhou, China).

**CCK-8 assay**

The cell viability was measured by CCK-8 assay (Beyotime, China). A549 cells (1×10⁴ cells/well) were cultured in 96-well plates and cultured for 24, 48 and 72 h. Subsequently, 10 µL of CCK-8 reagent was incubated for additional 4 h at 37 °C. At last, the cell optical density was detected by a Microplate Reader (Bio-Rad, USA) with absorbance at 450 nm.

**Cell migration and invasion assays**

The cell invasion and migration assays were performed by 24-well Transwell cell culture chambers with 8-µm sized pores with or without precoated Matrigel (BD Biosciences, San Jose, CA, USA). Specifically, A549 cells, A549 cells co-cultured with M2 macrophages treated with or without GW4869 and A549 cells co-cultured with exosomes from
M0/M2 macrophages or M2 macrophages transfected with different plasmids, at a density of $5 \times 10^4$ cells/mL, were re-suspended with 200 µL DMEM medium (serum-free) and seeded into the upper chamber, while the lower chamber was placed with 600 µL DMEM medium (10% FBS). After incubation for 24 h, the cells remaining in the upper chamber were removed, the invaded or migrated A549 cells were fixed with the methanol (100%), stained with crystal violet (0.1 mg/mL) and counted under a microscope.

**Isolation, identification and labeling of exosomes**

The exosomes from M0 or M2 macrophages were isolated by density gradient ultracentrifugation according to previously reported protocol (30). Briefly, cell culture medium was collected and centrifuged at 1,000 g for 10 min, 2,000 g for 20 min, 4,000 g for 30 min and 10,000 g for 1 h to obtain the supernatant. The exosomes were collected by centrifuging the supernatant at 100,000 g for 2 h at 4 °C. The size distribution and concentration of exosomes were analyzed at a flow rate of 0.03 Ml per min using a Zetasizer Nano ZS (Malvern Instrument, UK) and NanoSight NS300 (Westborough, MA, USA), respectively. Purified exosomes were labeled with the PKH-67 green fluorescent linker mini kit (Sigma, USA) according to the manufacturer’s instructions.

**miRNA profiling**

The aberrant miRNAs expressions of M0-exosome and M2-exosome were analyzed by miRNAs sequencing. Briefly, total RNA was isolated from M0-exosome and M2-exosome using TRIzol reagent (Invitrogen, MA, USA). 250 ng total RNA of M0-exosome and M2-exosome were extracted to prepare the small RNA sequencing library by using the NEBNext Multiplex Small RNA Library Prep Set for Illumina (NEB, USA). The libraries were finally sequenced and the Solexa CHASTITY quantity filtered reads were harvested as Clean Reads. For data analysis, differentially expressed miRNA profiles between M0-exosome and M2-exosome groups were compared, fold changes were calculated to identify significant differentially expressed miRNAs and hierarchical clustering was performed. The selected miRNAs were verified by qRT-PCR.

**Western blot analysis**

The radio-immunoprecipitation assay (RIPA) lysis buffer (Solarbio, Shanghai, China) with 0.5% phenylmethanesulfonyl fluoride (PMSF) (Solarbio, Shanghai, China) was used to extract the total protein of exosomes, cells or tissues. The protein concentration was quantified by the BCA protein quantification Kit (Sigma, USA). The primary antibodies in this study were purchased from Abcam: rabbit anti-CD63 (ab68418, 1:500), CD81 (ab109201, 1:200), RASSF4 (ab243709, 1:1,000) and TSG101 (ab30871, 1:500); mouse anti-E-cadherin (ab11512, 1:1,000), vimentin (ab8978, 1:1,000) and glyceraldehyde-3-phosphate dehydrogenase (GAPDH) (ab8245, 1:5,000). The horseradish peroxidase labeled goat anti-rabbit IgG antibody (ab205718, 1:10,000) and goat anti-mouse antibody (ab6789, 1:10,000) were available as the secondary antibodies. Image J software was used to quantify each protein band.

**Luciferase reporter assay**

The 3’-UTR segments of RASSF4 in wild type and mutant were synthesized and inserted into a firefly luciferase reporter construct. Cells were co-transfected with RASSF4 construct and miR-155 mimic or miR-196a-5p mimic. Luciferase activity in this study was measured by the Dual-Luciferase Reporter Assay System (Promega, USA) according to the protocol.

**Animal studies**

Six-week-old male athymic BALB/c nude mice (15–20 g) were purchased from the Model Animal Research Center of Nanjing University (Nanjing, China) and housed in a temperature-controlled room with a 12-hour light/dark cycle. Diet and water were available ad libitum. For the in vivo lung metastases model, exosomes purified from M2 macrophages or M2 macropahes transfected with 1×10^9 ifu of miR-155 inhibitor lentivirus or M2 macrophages transfected with 1×10^9 ifu of miR-196a-5p inhibitor lentivirus were respectively injected into the peritoneum. Four days post-injection, A549/Luc cells were injected into the tail vein of representative mice (n=5 per group, total 25). All mice were grouped randomly. The luciferase signal intensity from days 0 to 28 is on equivalent scales in the models. Bioluminescent flux (photons/s/cm^2/steradian) was determined for the lung metastases. Metastatic progression was monitored and imaged using an IVIS-100 system (Caliper Life Sciences, MA, USA) 10 min after intraperitoneal injection of luciferin (300 mg/kg i.v.) in
80 µL of saline. After 28 days, mice were euthanized using large doses of intraperitoneal pentobarbital (200 mg/kg) in the laboratory and tissues were separated for further experiments. Animal care and euthanasia were carried out with the approval of the Institutional Animal Care and Use Committee (IACUC) of Nanjing Medical University, in compliance with the national or institutional guidelines for the care and use of animals.

**Hematoxylin & eosin (HE) staining**

The dewaxed sections were firstly incubated with hematoxylin to stain the nucleus for 5 min, then 1% ethanol-hydrochloric acid for 30 s and eosin solution for 3 min. Finally, the sections were dehydrated in graded alcohol following by clearing in xylene.

**Statistical analysis**

All statistical analyses were performed by GraphPad Prism 6.0 software. The Student’s *t*-test was used to analyze significant differences in this study. The error bars indicate the standard deviation from the mean of triplicate measurements. Asterisks indicate significant differences (*P<0.05; **P<0.01; ***P<0.001) compared with the corresponding control.

**Results**

**M2 TAMs are abundant in metastatic tissues of NSCLC patients**

We firstly analyzed the enrichment of macrophage in different cancer types through the online database TCIA (https://tcia.at/). The result suggested that macrophages were most abundant in lung cancer (Figure 1A). NSCLC accounts for more than 80% of lung cancer and metastasis causes 70% of deaths. Then, to evaluate the role of macrophages in the metastasis of NSCLC, we utilized gene set enrichment analysis via the Cancer Genome Atlas database and observed that CD68, a macrophage marker, was significantly increased in 122 patients with the metastasis of NSCLC (Figure 1B). To further validate the distribution of macrophages in NSCLC, we analyzed the expression of CD68 by immunohistochemical staining and found that the higher density of TAMs in NSCLC metastasis tissues than that in localized tissues (Figure 1C,D), suggesting TAMs recruitment related to the metastasis of NSCLC. Next, we estimated the two subpopulations of macrophages including macrophage M0 and M2 in NSCLC patients. The results of qRT-PCR showed that the transcription expressions of M1 macrophages marker such as tumor necrosis factor alpha (TNF-α) and interferon regulatory factor 5 (IRF5) had a significant decrease in all localized samples in this study; however, the signatures of M2 macrophages like interferon regulatory factor 4 (IRF4) and arginase-1 (Arg-1) were up-regulated in metastatic tissues compared with localized tissues (Figure 1E). Moreover, flow cytometry assay suggested that M2 macrophages were more abundant in metastatic tissues (Figure 1F,G). These data indicated that M2 TAMs are probably involved in the metastasis of NSCLC.

**M2 TAMs promote migration, invasion and EMT of A549 cells**

The results in Figure 1 showed that M2 macrophages were more enriched in metastatic NSCLC patients than those without metastasis. To evaluate whether TAMs were involved in the metastasis of NSCLC, we co-cultured A549 cells with M0 or M2 macrophages in a Transwell insert. The macrophages have been extensively reported to regulate a series of biological processes of NSCLC through exosomes. To evaluate whether the effect of macrophages on A549 cells was through exosomes, we treated M0 or M2 macrophages with GW4869 (an inhibitor of exosome release). For inhibition of exosome generation, macrophages were cultured in media containing 10 µM GW4869. CCK-8 assay displayed that M2 macrophages significantly promoted cell viability compared to the control, and GW4869 treatment reversed the accumulation of cell viability caused by M2 macrophages (Figure 2A). The results of Transwell assay unveiled that the migration and invasion capacity of A549 cells was enhanced in A549 cells co-cultured with M2 macrophages (Figure 2B,C,D,E). Following GW4869 treatment, M2 macrophages failed to promote the migration and invasiveness of A549 cells, suggesting that exosomes derived from M2 macrophages play an essential role in the migration and invasiveness of A549 cells (Figure 2B,C,D,E). Besides, M2 macrophages decreased the protein expression of epithelial cell marker (E-cadherin) and promoted the expression of mesenchymal cell marker (vimentin), and GW4869 treatment reversed the effects (Figure 2F,G). Above all, these results clarified that M2 macrophages markedly promoted
migration, invasion, and EMT of A549 cells.

**Exosomes secreted by M2 TAMs promote migration, invasion and EMT of A549 cells**

Subsequently, a large number of exosomes were found in the supernatant of M2 macrophages culture by transmission electron microscope. And the shape and size of exosomes were solid with typical structure of two-layer membrane and exosomes had an average diameter of 100 nm as shown in Figure 3A,B. Western blot analysis in Figure 3C showed that the protein levels of exosomal markers TSG101, CD63 and CD81 had a significant increase in exosomes compared with cell lysis, confirming the successful extraction of exosomes. To examine whether M2 macrophages derived exosomes (M2 exosomes) can be taken up by A549 cells, we pre-labeled M2 exosomes with PKH67. In Figure 3D, A549 cells co-cultured with PKH67-labeled M2 exosomes for 48 h exhibited green fluorescence, confirming the exosomes derived from M2 macrophages can be taken up by A549 cells. Furthermore, after 48 h of co-culture of A549 cells with M0 exosomes or M2 exosomes in vitro. CCK-8 assay, Transwell assay and western blot analysis were applied to examine whether exosomes generated from M2 macrophages were sufficient to induce the cell migration and invasion. CCK-8 assay suggested that exosomes secreted from M2 macrophages notably raised the A549 cell viability, while exosomes from M0 macrophages had no effect on cell viability (Figure 3E). The results in Figure 3F,G,H,I revealed that the invasiveness and migration ability of A549 cells co-
Figure 2 M2 TAMs promote migration, invasion and EMT of A549 cells. (A) The viability of A549 cells co-cultured with M0 macrophages, M2 macrophages, M0 macrophages + GW4869, or M2 macrophages + GW4869 was examined by CCK-8 assay. (B,C,D,E) The migration and invasion abilities of A549 cells were measured by Transwell assay (200x magnification). Scale bar =100 µm. (F,G) The protein expression levels of epithelial cell marker (E-cadherin) and mesenchymal cell marker (vimentin) in A549 cells were analyzed by western blot. Error bars represent standard deviations and asterisks show significant differences from corresponding control according to Student’s t-test (*P<0.05, **P<0.01, ***P<0.001).

M2 macrophages-derived exosomes promote migration, invasion and EMT of A549 cells through transporting miR-155 and miR-196a-5p

Emerging evidence indicated that miRNAs involved in cell-cell communications are frequently encapsulated in exosomes, and implement their biological functions in the recipient cells. To explore the mechanisms by which M2-exosome produced a marked effect in NSCLC, we generated miRNA profiles of M0-exosome and M2-exosome by miRNA microarray analysis. The comparison of the expression levels of the miRNAs in the M0-exosome and M2-exosome groups is depicted in Figure 4A,B. Among the miRNAs, miR-155 and miR-196a-5p were the most abundant in M2-exosome group in Figure 4A,B. In further analysis, in Figure 4C, miR-155 and miR-196a-5p were markedly upregulated in M2-exosome group. The results showed that miR-155 and miR-196a-5p were enriched in M2 macrophages-derived exosomes. Exosomes deliver proteins, miRNAs or mRNAs into target cells to mediate a series of physiological processes. Therefore, we hypothesized that M2-exosomes might induce the increased migration of recipient A549 cells through the transfer of functional miR-155 and miR-196a-5p. According to CCK-8 assay, we found that M2-exosomes promoted the cell viability, then the viability was restrained by transfecting miR-155 inhibitor and miR-196a-5p inhibitor (Figure
Figure 3 Exosomes secreted by M2 TAMs promote EMT and migration of A549 cells. (A) The structure of exosome was identified by transmission electron microscope (40.0k× magnification). Scale bar =200 nm. (B) The size of exosome was detected by nanoparticle tracking analysis. (C) The protein levels of exosomal markers TSG101, CD63 and CD81 in exosomes and cell lysis were analyzed by western blot. (D) The fluorescence signal from A549 cells co-cultured with PKH67-labeled M2 exosomes or control was detected (400× magnification). Scale bar =50 μm. (E) The viability of A549 cells co-cultured with M0 exosomes or M2 exosomes were examined by CCK-8 assay. (F, G, H, I) The migration and invasive abilities of A549 cells after M0 exosome or M2 exosome treatment in vitro was examined by Transwell assay (200× magnification). Scale bar =100 μm. (J, K) The protein expression levels of epithelial cell marker (E-cadherin) and mesenchymal cell marker (vimentin) in A549 cells after M0 exosome or M2 exosome treatment were analyzed by western blot. Error bars represent standard deviations and asterisks show significant differences from corresponding control according to Student’s t-test (*P<0.05, **P<0.01, ***P<0.001).

5A). In Figure 5B, C, D, E, we found that the M2-exosomes promoted the migration of A549 cells, increased the protein expression level of vimentin and inhibited the E-cadherin expression, while the opposite pattern could be observed in M2/miR-155-inhibitor exosomes group and M2/miR-196a-5p-inhibitor exosomes group. Taken together, these data suggested that miR-155 and miR-196a-5p carried by exosomes derived from M2 macrophages mediated a series of biological processes of NSCLC.

The direct target of exosomal miR-155 and miR-196a-5p in NSCLC is RASSF4

To explore the common targets between miR-155 and miR-196a-5p participated in the development of NSCLC, online bioinformatics database (TargetScan Release 7.0) was executed to predict the target genes. Based on these, David’s bioinformatics resources and literature review were performed to evaluate analyze genes function in cancer, which showed 8 genes were associated with cancer progression (USP48, TRMT1L, HOOK1, ZMYND11, LCOR, RASSF4, BCAT1, ZBTB39) (Figure 6A). Moreover, RT-qPCR analysis displayed that RASSF4 had the largest fold difference in A549 cells treating with M2 exosomes (Figure 6B). The prediction of TargetScan Release 7.0 database showed that RASSF4 in 3′-untranslated regions (UTRs) possesses putative binding sites for miR-155 and miR-196a-5p (Figure 6C). Subsequently, the luciferase reporter plasmids containing the wild type (WT) or mutation-type of 3′-UTR of RASSF4 were successfully transfected into A549 cells. As shown in Figure 6D, miR-155 or miR-196a-5p significantly decreased the luciferase
Figure 4 MiR-155 and miR-196a-5p are abundant in M2 macrophages-derived exosomes. (A) Volcano plot of the differentially expressed miRNAs as assessed by microarray analysis in M0 and M2 exosomes. (B) Microarray analysis of exosomal miRNAs in M0 and M2 TAMs were presented in a heatmap. (C) The transcription expression of miR-155, miR-196a-5p, miR-3091-3p, miR-12206-5p and miR-12180-3p in M0-exosome and M2-exosome groups. Error bars represent standard deviations and asterisks show significant differences from corresponding control according to Student's $t$-test (**P<0.001).
activity of WT RASSF4 compared with NC. Western blot analysis revealed that miR-155 or miR-196a-5p overexpression significantly down-regulated the expression of RASSF4, while miR-155 or miR-196a-5p inhibitor up-regulated RASSF4 expression (Figure 6E,F). Collectively, these results testified that RASSF4 was the target of miR-155 and miR-196a-5p.

**MiR-155 and miR-196a-5p promote migration, invasion and EMT of A549 cells through regulating RASSF4 expression**

We carried out the rescue assays to further explore whether miR-155 and miR-196a-5p executed its performance by RASSF4. The result of CCK-8 assay indicated that overexpression of RASSF4 suppressed the cell viability induced by miR-155 mimic and miR-196a-5p mimic, while the repression of RASSF4 abrogated the anti-proliferation effects of miR-155 inhibitor and miR-196a-5p inhibitor (Figure 7A). Similar results were observed in cell migration and invasion (Figure 7B,C,D,E). Western blot analysis revealed that the significant down-regulation in the expression of RASSF4 and E-cadherin, and up-expression of vimentin resulting from miR-155 and miR-196a-5p overexpression, these results were reversed when RASSF4 was overexpressed. Correspondingly, the
Figure 6 The direct target of exosomal miR-155 and miR-196a-5p in NSCLC is RASSF4. (A) The underlying targets of miR-155 and miR-196a-5p were predicted by online bioinformatics databases (TargetScan Release 7.0, David's bioinformatics resources). (B) RT-qPCR assay was carried out to detect the mRNA expression of predicted targets in A549 cells treating M2-exosome. (C) The putative binding sites of RASSF4 for miR-155 and miR-196a-5p by TargetScan Release 7.0 database. (D) The luciferase activities in A549 transfected with wild-type or mutated 3’UTR of RASSF4 were measured. (E,F) The protein expression levels of RASSF4 in A549 cells transfected with negative control (NC), miR-155 mimic (miR-155 OE) or miR-155 inhibitor (miR-155 KD) or miR-196a-5p OE or miR-196a-5p KD were analyzed by western blot. Error bars represent standard deviations and asterisks show significant differences from corresponding control according to Student’s t-test (**P<0.01, ***P<0.001).

promoted expressions of RASSF4 and E-cadherin by miR-155 and miR-196a-5p knockdown were alleviated by RASSF4 inhibition (Figure 7F,G). Taken together, the effects of miR-155 and miR-196a-5p on A549 cells, including cell viability, cell migration, cell invasion, and the protein levels of RASSF4, vimentin and E-cadherin, could be reversed by RASSF4.

M2 TAMs secret exosomal miR-155 and miR-196a-5p to promote NSCLC metastasis in vivo

6-week-old healthy male BALB/c nude mice were performed to establish transplantation tumor model. To assess the effect of miR-155 and miR-196a-5p carried by exosomes from M2 macrophages on lung metastasis in vivo, M2 exosome, M2/miR-155 knockdown exosome, M2/miR-196a-5p knockdown exosome, M2/miR-155 knockdown + miR-196a-5p knockdown exosome were injected respectively after intravenous injection of A549/Luc cells into mice to construct lung metastasis model. In Figure 8A,B, M2-derived exosomes markedly contributed to the lung dissemination. And the high expression of luciferase activity in M2 exosome group could be effectively decreased through the knockdown of miR-155 and miR-196a-5p. Next, HE staining from lung tissues of each group in Figure 8C,D showed that M2-derived exosomes with the knockdown of miR-155 and miR-196a-5p markedly contributed to alleviate the metastasis of NSCLC resulting from M2 exosomes. In Figure 8E,F, western blot analysis from tumor tissues of each group revealed that the protein levels of RASSF4 in M2/miR-155 knockdown and miR-196a-5p knockdown exosome group were significantly up-regulated compared with M2-exosome group. Additionally, M2/miR-155 knockdown and miR-196a-5p knockdown exosome group promoted E-cadherin expression and decreased vimentin expression compared with M2 exosome group. In general, our findings reveal that NSCLC metastasis can be
Figure 7 MiR-155 and miR-196a-5p promote migration, invasion and EMT of A549 cells through regulating RASSF4 expression. (A) The viability of A549 cells with the corresponding transfection was examined by CCK-8 assay. (B,C,D,E) The migration and invasion abilities were measured by Transwell assay in A549 cells (200× magnification). Scale bar =100 µm. (F,G) The protein expression levels of RASSF4, E-cadherin and vimentin were analyzed by western blot in A549 cells. Error bars represent standard deviations and asterisks show significant differences from corresponding control according to Student’s t-test (*P<0.05, **P<0.01, ***P<0.001).

Discussion

Due to the metastasis to bone, brain and liver, poor prognosis is common in lung cancer patients, resulting in high mortality (31). Although various biomolecules have been found to mediate lung cancer metastasis, knowledge regarding to roles of tumor microenvironment in lung cancer metastasis has largely lagged behind. Stromal cells including TAMs in the tumor microenvironment are either induced or directly interacting with cancer cells to promote the progression and metastasis of lung cancer (32,33). The modes of action of stromal cells include the secretion of extracellular components, such as growth factors, chemokines and exosomes (34). However, due to the extreme complexity of these processes, the molecular mechanisms underlying lung metastasis in the tumor microenvironment have not been fully understood.

Multiple evidence has expounded that M2 TAMs participated in the progression of tumors (35). In the study, we found that macrophages were most abundant in lung cancer, and M2 macrophages were rich in metastatic tissues. Based on these results, we speculated M2 TAMs were involved in the metastasis of NSCLC. The alveolar M2 TAM density was reported to be associated with tumor differentiation, invasive size, pathological stage and poor prognosis (36). The results indicated that M2 TAMs promoted NSCLC metastasis, which were consistent with Yuan and Guo (20,21). Nevertheless, the exact mechanisms by which M2 TAMs promote NSCLC metastasis remained undefined. There is increasing evidence that stromal exosomes play an important role in the tumor microenvironment as a medium for intercellular communication (37). For instance, Zheng et al. suggested...
that M2 TAMs-derived exosomes promoted migration of gastric cancer cells (38). Similarly, our result indicated that exosomes secreted from M2 macrophages can markedly promote migration, invasion, and EMT.

Exosomes from macrophages easily selectively deliver biologically active substances, such as proteins, miRNAs or mRNAs, to recipient cells to achieve efficient interflow between macrophages and tumor cells (39). MiRNAs are important gene regulators, which control various physiological and pathological processes (40). For instance, Yin et al. reported that exosomal miR-501-3p from macrophages induced the progression of pancreatic ductal adenocarcinoma through TGFBR3 (26). Besides, Lan et al. clarified that M2 macrophage-derived exosomes...
transferred miR-155-5p and miR-21-5p to colorectal cancer cells to accelerate tumor progression (41). In this paper, we demonstrated that the levels of miR-155 and miR-196a-5p were up-regulated in M2 TAMs-derived exosomes. MiR-155 is previously identified as an oncogenic miRNA in various cancers, such as breast cancer, ovarian cancer and lung cancer (42-44). Previous studies have also suggested that miR-19a-5p facilitated the cell proliferation in colorectal cancer (45), bladder cancer (46), gastric cancer (47). However, whether miR-196a-5p regulates the progression of NSCLC is not fully understood. In this paper, a series of gain-of-function and loss-of-function experiments illustrated exosomal miR-155 and miR-196a-5p promoted EMT and migration of NSCLC cells and promoted NSCLC metastasis in vivo.

It is confirmed that miRNAs regulate the biological effects of cancer cell by directly binding to the 3'UTR of mRNA (48). Therefore, targeting certain miRNAs involved in NSCLC has been considered as an efficient therapeutic approach (49). Moreover, we found that RASSF4 was a direct downstream target of both miR-155 and miR-196a-5p. RASSF4 is considered as a protein of the RASSF family (27). RASSF proteins directly interact with RAS proteins with effector characteristics to induce RAS-dependent cell death (50). In addition, RASSF protein has been reported to play an important role in tumor inhibition and is involved in many important biological functions, such as proliferation, cell cycle, apoptosis, and DNA repair (51). A number of studies suggested that RASSF4 was widely expressed in normal tissues but down-regulated in tumors, indicating that RASSF4 acted as a tumor suppressor in numerous malignancies (27,52). In our study, we found that exosomal miR-155 and miR-196a-5p secreted by M2 TAMs in NSCLC metastasis modulated RASSF4 to promote NSCLC progression.

Inhibiting miRNA-enriched exosomes from M2 TAMs is useful for treating cancer metastasis (53,54). In our work, we found that inhibiting exosomal miR-155 and miR-196a-5p could alleviate NSCLC metastasis activated by exosomes derived from M2 TAMs. Therefore, targeting exosomal miR-155 and miR-196a-5p from M2 TAMs may be a potential therapeutic strategy for NSCLC metastasis. Moreover, previous researches have also indicated that miRNA can control polarization of TAMs (55). Whether miR-155 and miR-196a-5p are involved in the M2 polarization of TAMs deserve further investigation, which is now underway in our group.

Overall, this study showed that M2 TAMs were the main population of TAMs in metastic tissues of NSCLC and promoted NSCLC metastasis. The cross-talk between M2 TAMs and lung cancer cells was mediated by exosomes that carry miR-155 and miR-196a-5p. MiR-155 and miR-196a-5p, which were abundant in both M2 TAMs and exosomes from M2 TAMs, promoted EMT and migration of NSCLC through targeting RASSF4. Inhibition of miR-155 or miR-196a-5p in M2 TAMs could prevent NSCLC metastasis. These findings suggest an undescribed mechanism of exosomes transfer from M2 macrophages to NSCLC cells, which explains the way immune cells participate in tumor progression.

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**Footnote**

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**Ethical Statement:** The authors are accountable for all aspects of the work in ensuring that questions related to the accuracy or integrity of any part of the work are appropriately investigated and resolved. The study was conducted in accordance with the Declaration of Helsinki (as revised in 2013). The project has attained informed consent from all patients and was approved by the Ethics Committee of Jiangsu Province People’s Hospital and the First Affiliated Hospital of Nanjing Medical University (No. 2019-SR-266). The Laboratory Animal...
Care and Use Committees of the hospital approved all experimental procedures, which are in compliance with the national or institutional guidelines for the care and use of animals.

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